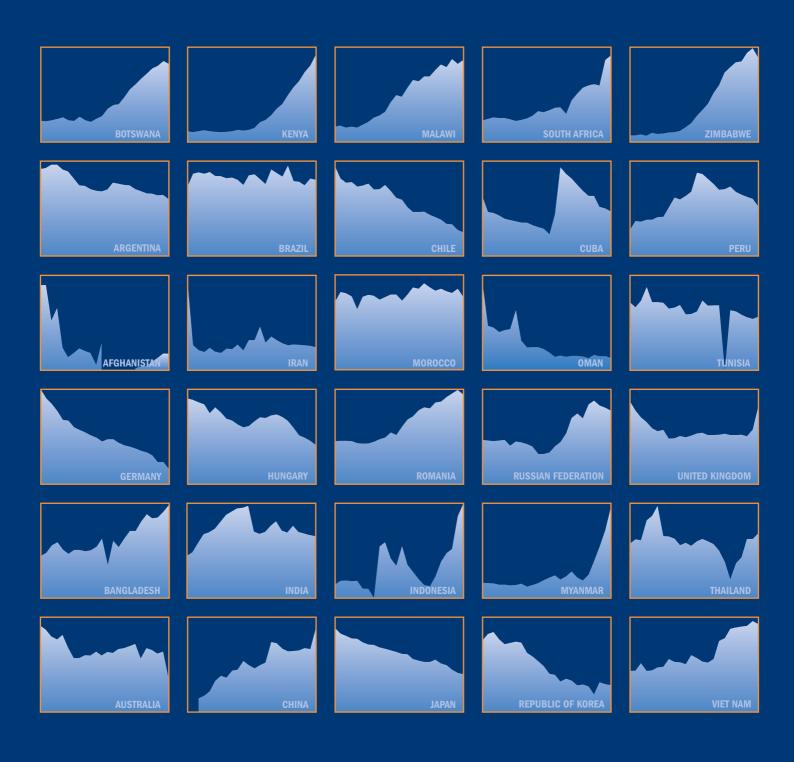
## **WHO REPORT 2005**

# **Global Tuberculosis Control**

Surveillance, Planning, Financing





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For reasons of space, the names of Member States are sometimes shortened in certain figures.

**Cover:** Cover graphics show trends in numbers of notified TB cases, 1980–2003. Each row has an arbitrary selection of countries from one of WHO's six regions. With careful interpretation, these trends can help to assess progress towards Millennium Development Goal 6, Target 8: "to have halted by 2015 and begun to reverse the incidence of malaria and other major diseases".

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As this report went to press, we learnt of the tragic death of our colleague Lisa Véron. During her short period in Africa, Lisa contributed enormously to TB control in the region, and especially to the development of the financial monitoring project. We miss her greatly.

# **Abbreviations**

ADB	Asian Development Bank	GDF	Global TB Drug Facility
AFB	Acid fast bacilli	GFATM	Global Fund to Fight AIDS, Tuberculosis
AFR	WHO African Region		and Malaria
AFRO	WHO Regional Office for Africa	GLC	Green Light Committee
AIDS	Acquired immunodeficiency syndrome	GLRA	German Leprosy and TB Relief Association
AMR	WHO Region of the Americas	GMS	German Medical Service
AMRO	WHO Regional Office for the Americas	GNI	Gross national income
ART	Antiretroviral therapy	GTZ	Deutsche Gesellschaft für Technische
BPHS	Basic package of health-care services		Zusammenarbeit (German development
BRAC	Bangladesh Rural Advancement		agency)
	Committee	HBC	High-burden country of which there are 22
CAREC	Caribbean Epidemiology Centre		that account for approximately 80% of all
CARE Interna	ational International relief and		new TB cases arising each year
	development organization	HIV	Human immunodeficiency virus
CB DOTS	Community-based DOTS	HR	Human resource
CCM	Country Coordinating Mechanism (Global	HRD	Human resource development
	Fund)	HRDP	Human resource development plan
CDC	Centers for Disease Control and	HSD	Health subdistrict
000	Prevention, Atlanta, GA, USA	HSSP	Health Sector Strategic Plan
CENAT	Centre National Anti-Tuberculeux	IEC	Information, education, communication
CI	Confidence interval	IFRC	International Federation of Red Cross and
CIDA	Canadian International Development	11110	Red Crescent Societies
OIDA	Agency	IRR	Incidence rate ratio
COOPI	Cooperazione Internazionale (Italian NGO)	ISAC	Intensified support and action in
COMBI	Communication for behavioural impact	10/10	countries, an emergency initiative to reach
DANIDA	Danish International Development Agency		targets for DOTS implementation by 2005
DCPP	Disease Control Priorities Project	IUATLD	International Union Against Tuberculosis
DDC		IUAILD	
	Department of Disease Control	шол	and Lung Disease
DEWG	DOTS Expansion Working Group of the	JICA	Japan International Cooperation Agency
DED	Stop TB Partnership	KIT	Royal Tropical Institute (Netherlands)
DFB	Damien Foundation Belgium	KNCV	Royal Netherlands Tuberculosis
DFID	UK Department for International	LEDGG	Association
DUT	Development	LEPCO	Tuberculosis and Leprosy Control (German
DHT	District health team	1.04	NGO)
DoH	Department of Health	LGA	Local government area
DOT	Directly observed treatment	MDG	Millennium Development Goal
DOTS	The internationally recommended strategy	MDR	Multidrug resistance
	for TB control	MDR-TB	Multidrug-resistant tuberculosis
DRS	Drug Resistance Surveillance	MEDAIR	An international humanitarian aid
DST	Drug susceptibility testing		organization
DTC	District TB coordinator	МоН	Ministry of Health
DRS	Drug resistance surveillance	MoPH	Ministry of Public Health
EMR	WHO Eastern Mediterranean Region	MSF	Médecins Sans Frontières
EMRO	WHO Regional Office for the Eastern	MSH	Management Sciences for Health
	Mediterranean	NGO	Nongovernmental organization
EQA	External quality assurance	NHLS	National Health Laboratory Services
EU	European Union	NICC	National interagency coordinating
EUR	WHO European Region		committee
EURO	WHO Regional Office for Europe	NLR	Netherlands Leprosy Relief
FCT	Federal Capital Territory	NPO	National programme officer (WHO-
FDC	Fixed-dose combination (or FDC anti-TB		appointed)
	drug)	NRL	National reference laboratory
FIDELIS	Fund for Innovative DOTS Expansion,	NTI	National TB institute
	managed by IUATLD	NTP	National tuberculosis control programme

PAHO	Pan-American Health Organization	TB	Tuberculosis
PAL	Practical Approach to Lung Health	TBCTA	Tuberculosis Coalition for Technical
PATH	International health NGO that focuses on		Assistance
	advancing technologies, strengthening	UNAIDS	Joint United Nations Programme on HIV/
	systems and encouraging healthy		AIDS
	behaviour	UNDP	United Nations Development Programme
PHC	Primary health care	USAID	United States Agency for International
PHIICAT	Philippines Coalition against TB		Development
PLWHA	People living with HIV/AIDS	USTP	Uganda Stop TB Partnership
PPM	Public-private or public-public mix	VCT	Voluntary counselling and testing for HIV
PPP	Purchasing power parity		infection
RIT	Research Institute of Tuberculosis, Japan	WHO	World Health Organization
	Anti-Tuberculosis Association	WHO-CHOICE	Choosing interventions that are cost-
SARS	Severe acute respiratory syndrome		effective, a WHO project
SEAR	WHO South-East Asia Region	WPR	WHO Western Pacific Region
SEARO	WHO Regional Office for South-East Asia	WPRO	WHO Regional Office for the Western
STD	Sexually transmitted disease		Pacific
STI	Sexually transmitted infection	ZTLS	Zonal TB and leprosy supervisor
SVS	Secretary of health surveillance		

# Summary

### **Background and methods**

- 1. The 9th World Health Organization (WHO) annual report on surveillance, planning and financing for TB control includes data on case notifications and treatment outcomes from all national TB control programmes (NTPs) that have reported to WHO, together with an analysis of plans, budgets, expenditures and progress in DOTS expansion for 22 high-burden countries (HBCs).
- 2. Ten consecutive years of data (1994-2003) are now available to assess progress towards the Millennium Development Goals (MDGs) for TB control. The five MDG targets directly relevant to TB control are: by 2005, to detect 70% of new smearpositive cases and successfully treat 85% of these cases; by 2015, to have halted and begun to reverse incidence; between 1990 and 2015, to halve TB prevalence and deaths rates.

## Improving case detection and treatment

- 3. A total of 199 countries reported to WHO on their strategies for TB control, and on TB case notifications and/ or treatment outcomes.
- 4. Using surveillance and survey data to update estimates of incidence, we calculate that there were 8.8 million new cases of TB in 2003 (140/ 100 000 population), of which 3.9 million (62/100 000) were smearpositive and 674 000 (11/100 000) were infected with human immunodeficiency virus (HIV). There were 15.4 million prevalent cases (245/ 100 000), of which 6.9 million were smear-positive (109/100 000). An estimated 1.7 million people (28/ 100 000) died from TB in 2003, including those coinfected with HIV  $(229\ 000).$
- 5. A total of 182 countries were implementing the DOTS strategy during 2003, two more countries than in 2002. By the end of 2003, 77% of the world's population lived in coun-

- tries, or parts of countries, covered by DOTS. DOTS programmes notified 3.7 million new and relapse TB cases, of which 1.8 million were new smearpositive. In total, 17.1 million TB patients, and 8.6 million smearpositive patients, were treated in DOTS programmes between 1995 and 2003.
- 6. The 1.8 million smear-positive cases notified by DOTS programmes in 2003 represent a case detection rate of 45%. The increment in smearpositive cases notified under DOTS between 2002 and 2003 (324 000) was greater than ever before (the average annual increment from 1995-2000 was 134 000). The acceleration in notifications was more pronounced for all TB cases, which increased by 693 000 between 2002 and 2003, compared with the average annual increment of 270 000 in the interval 1995-2000.
- 7. While the number of TB cases reported by DOTS programmes appears to have been accelerating since 2000, the total number of TB cases reported to WHO (all forms from all sources) increased very little over the period 1995-2003 (average detection rate 42%). The number of smear-positive cases reported from all sources has been increasing (50% detection rate in 2003), but much more slowly than the increases reported under DOTS.
- 8. Of the additional smear-positive cases reported under DOTS in 2003, 63% were in just two countries: India and China. Among those individuals who are thought to have developed smear-positive TB in 2003, but were not detected by DOTS programmes, 67% were living in just eight countries: Bangladesh, China, Ethiopia, India, Indonesia, Nigeria, Pakistan and the Russian Federation.
- 9. As DOTS programmes have expanded geographically, the new smearpositive case detection rate within DOTS areas has remained roughly constant since 1995 (average 52%),

- although there are signs of a slow increase in the HBCs, especially in Bangladesh, India, Myanmar and the Philippines.
- **10.** The rate of treatment success in the 2002 DOTS cohort was 82% on average, unchanged since 2000. As in previous years, the treatment success rate was substantially below average in the WHO African Region (73%) and the WHO European Region (76%). Low treatment success rates in these two regions can be attributed, in part, to the complications of TB/HIV coinfection and drug resistance, respectively. Equally important, though, is the failure of DOTS programmes in these two regions to monitor the outcome of treatment for all patients.
- 11. Based on case reports and WHO estimates, 22 countries had reached the targets for case detection and treatment success by the end of 2003. Viet Nam was still the only member of the current group of HBCs1 among them, although Cambodia, Myanmar and the Philippines are within reach.

## **Epidemiological trends and DOTS** impact

- 12. In 2003, the TB incidence rate was falling or stable in five out of six WHO regions, but growing at 1.0% per year globally. The exception is the African Region, where incidence has been rising more quickly in countries with higher HIV prevalence rates. In Eastern Europe the incidence rate increased during the 1990s, but peaked around 2001, and has since fallen. The rise in global incidence is slowing because HIV epidemics are slowing in Africa, but it is unclear when the global incidence rate will begin to decline.
- 13. We calculate that, as a consequence of DOTS expansion between 1990 and 2003, the global TB preva-

<sup>&</sup>lt;sup>1</sup> Peru was excluded from the original group of high-burden countries, having met the targets and successfully reduced incidence.

lence rate fell from 309 to 245 per 100 000 (including HIV-positive TB patients), and by 5% between 2002 and 2003, even though incidence continued to rise. The global mortality rate peaked during the 1990s, and fell at 2.5% (including HIV-positive TB patients) or 3.5% per year (excluding HIV-positive TB patient) between 2002 and 2003. But for the strongly adverse trends in Africa, prevalence and death rates would be falling more quickly worldwide

## Planning and DOTS implementation

- **14.** All HBCs have a strategic plan for DOTS expansion and, during 2005, many will begin a new planning cycle, ideally working towards the MDG target year of 2015. Although the health systems of many countries are still undergoing reform and restructuring, all HBCs except the Russian Federation and Thailand reported that TB control functions are fully integrated with essential national health services.
- 15. Among the obstacles to DOTS expansion, five are of overriding importance: shortages of trained staff; lack of political commitment; weak laboratory services; and inadequate management of multidrug-resistant TB (MDR-TB), and of TB in people infected with HIV. Concerning drug resistance, few countries have national policies for the diagnosis and treatment of MDR-TB: even in those that do. treatment commonly fails to meet acceptable standards. Concerning TB/HIV, NTPs reported that few TB patients are tested for HIV (3% of notified cases), still fewer are assessed for antiretroviral therapy (ART) and a very small fraction begin ART (1349 patients reported in 2003). The report discusses a wide range of remedial actions to overcome these constraints.
- **16.** Intensified support and action in countries (ISAC) is a new initiative designed to catalyse and accelerate DOTS expansion towards 2005 targets. The goal of ISAC is to improve technical capacity so as to facilitate the spending of large grants from the Global Fund to Fight AIDS, Tuberculosis and Malaria (GFATM) and other

- major donors. Participants in 2004 included China, India, Indonesia, Kenya, Pakistan, Romania, the Russian Federation and Uganda.
- 17. The increasing contributions of nongovernmental organizations (NGOs) and community groups are clear expressions of the growing commitment of civil society to TB control. The work of these groups puts patients at the centre of the DOTS strategy, and improves access to TB services in remote areas and among disadvantaged and marginalized populations.
- **18.** Public–private and public–public mix (PPM) projects are showing a measurable impact on case detection in several Asian countries, and may prove to be a mechanism for expanding TB control services in African cities.

### **Financing DOTS expansion**

- **19.** Financial data were received from 134 out of 211 (64%) countries, up from 123 in 2003. Complete budget and expenditure data were provided by 70 and 69 countries, respectively. Data were received from all 22 HBCs, except South Africa.
- 20. There has been a big increase in NTP budgets and a big improvement in the funding available for TB control in the HBCs since 2002, with particularly large increases between 2003 and 2004. The NTP budgets reported for 2005 total US\$ 741 million. The total estimated costs of TB control are projected to be US\$ 1.3 billion in 2005, and available funding is US\$ 1.2 billion. The large increase in available funding is almost entirely due to additional government funding in China, Indonesia and the Russian Federation, and to GFATM grants.
- **21.** The countries with the largest NTP budgets in 2005 are China, India, Indonesia and the Russian Federation. When costs beyond those included in NTP budgets are also considered, China, India, the Russian Federation and South Africa account for US\$ 946 million (73%) of the US\$ 1.3 billion total. Eight HBCs have total costs of US\$ 20–50 million in 2005; the rest are US\$ 18 million or less.

- 22. Per patient treated, there is considerable variation in budgets for firstline drugs, in total NTP budgets and in total costs for each year 2002-2005. Among HBCs, the NTP budget per patient is lowest in India (US\$ 34). Most countries have budgets in the range US\$ 100-200 per patient, and costs in the region of US\$ 150-300. The Russian Federation and South Africa are notable exceptions, with costs per patient treated above US\$ 1000. Budgets per patient treated are generally stable or increasing, and as a consequence costs per patient treated are also generally stable or increasing.
- **23.** In 2005, HBC governments are providing 62% of NTP budgets (including loans), the GFATM 15%, and grants from other sources 7%, leaving a gap equivalent to 16% of the reported budgets. HBC governments contribute more (79%) to total costs than to NTP budgets because they finance the general health services staff and infrastructure used for TB control. High average contributions to the financing of TB control by HBC governments conceal the fact that many HBCs rely extensively on grant funding.
- **24.** Despite progress in securing additional funding, HBCs reported a funding gap of US\$ 119 million in 2005. This is higher than the gaps reported for 2003 and 2004. The largest funding gaps are those reported by China, India, Pakistan, the Russian Federation and Zimbabwe (US\$ 93 million, or 78% of the total gap). Proportional to budgets, the largest gaps are in Kenya, Nigeria, Pakistan, Uganda and Zimbabwe.
- **25.** Planned activities are not in line with meeting the case detection target in 2005 in 12 countries. In addition, the budgets for collaborative TB/HIV activities and for second-line drugs to treat MDR-TB are currently small. This means that the gaps currently reported by NTPs could be regarded as underestimates, and that the total resources required for TB will be higher than US\$ 1.3 billion in future.

- **26.** Absorption capacity is a major issue for HBCs that have secured substantial amounts of additional funding. Expenditures were lower than available funding in 2003; it remains to be seen whether NTPs can effectively spend the extra money available in 2004 and 2005.
- 27. In financing terms, the HBCs fall into four categories: (a) four countries (India, Myanmar, the Philippines and Viet Nam) that have budgets consistent with reaching the 2005 targets, and which are likely to have minimal or no funding shortfall; (b) four countries that have adequate budgets, but which need to make up funding shortfalls (Cambodia, China), or where it is unclear how many more cases will be detected and successfully treated as a result of the substantial additional funds now available (Bangladesh, Indonesia); (c) five countries whose plans are not in line with meeting the 2005 targets, but which report mini-

mal or no funding gaps; (d) nine countries that report large funding gaps and whose plans are less than required to meet the targets for case detection (eight countries) and/or it is not clear if they are sufficient to meet the target for treatment success. These nine countries merit particular attention from donors and other support agencies.

## **Progress towards the Millennium Development Goals**

28. If the improvement in case-finding between 2002 and 2003 can be maintained, the case detection rate will be 60% in 2005. To reach the 70% target, DOTS programmes must recruit TB patients from non-participating clinics and hospitals, especially in the private sector in Asia, and from beyond the present limits of public health systems in Africa. To reach the target of 85% treatment success, a special effort must be made to improve cure rates in Africa and Eastern Europe.

- 29. Our analysis of epidemiological trends suggests that the TB incidence rate is still slowly rising globally, but prevalence and death rates are falling. Whether the burden of TB can be reduced sufficiently to reach the MDGs by 2015 depends on how rapidly DOTS programmes can be implemented by a diversity of health-care providers, and how effectively they can be adapted to meet the challenges presented by HIV coinfection (especially in Africa) and drug resistance (especially in eastern Europe).
- 30. Financing for global TB control has improved since 2002, dramatically in some countries. Some HBCs now have sufficient funds to meet targets, but must show that they can spend them effectively; some have no apparent shortfall, but should verify that their budgets are sufficient; some have an obvious funding gap, and must focus on raising the money needed to improve programme performance.

#### Key epidemiological and financial indicators

EPIDEMIOLOGICAL INDICATORS (WORLD)	MDG TARGET	TARGET YEAR	ESTIMATE 2003	CHANGE, REFERENCE YEAR TO 2003 (%)	REFERENCE YEAR
DOTS case detection (%)	70	2005	45	+7.5	2002
DOTS treatment success (%)	85	2005	82 (2002 cohort)	0.0	2001 cohort
Incidence rate (per 100 000 per year exc HIV)	falling	2015	129	+0.6	2002
Incidence rate (per 100 000 per year inc HIV)*			140	+1.0	2002
Prevalence rate (per 100 000 exc HIV)	half 1990 level	2015	240	-22	1990
Prevalence rate (per 100 000 inc HIV)			245	-21	1990
Mortality rate (per 100 000 per year exc HIV)	half 1990 level	2015	24	-12	1990
Mortality rate (per 100 000 per year inc HIV)			28	-1.6	1990
FINANCIAL INDICATORS (HIGH-BURDEN COUNTRIES)			ESTIMATE 2005	CHANGE, 2002-2005 (%)	REFERENCE YEAR
Total costs of TB control (US\$ millions)			1321	+49	2002
NTP budgets for TB control (US\$ millions)			741	+79	2002
Total funding available for TB control (US\$ millions	s):		1202	+36	2002
Government (excl. loans)			982	+26	2002
Loans			56	+102	2002
Grants (excl. GFATM)			55	+29	2002
GFATM			109	NA	2002
Funding gap as reported by NTPs (US\$ millions)			119	+34	2002
Costs per patient (US\$) (median values)					
Total cost			213	+22	2002
NTP budget			133	+45	2002
First-line drugs budget			28	-12	2002

<sup>\*</sup> inc HIV: including HIV+ TB patients; MDG indicators for TB exclude HIV+ patients, but these statistics are also useful in TB control. NA: not applicable: funds first distributed in 2003.

## Résumé

#### Contexte et méthodes

- 1. Le neuvième rapport annuel de l'Organisation Mondiale de la Santé (OMS) sur la surveillance, la planification et le financement de la lutte antituberculeuse contient des informations sur le nombre de cas notifiés et les résultats du traitement en provenance de tous les programmes nationaux de lutte antituberculeuse (PNT) qui ont envoyé des rapports à l'OMS, ainsi qu'une analyse des plans, budgets, dépenses et progrès concernant l'extension de la stratégie DOTS dans les 22 pays les plus touchés par la tuberculose.
- 2. On dispose désormais de données s'étendant sur dix années consécutives (1994-2003) pour évaluer les progrès accomplis en vue d'atteindre les objectifs du Millénaire pour le développement (OMD) qui concernent la lutte antituberculeuse. Les cinq objectifs des OMD concernant directement la lutte contre la tuberculose sont: d'ici 2005, dépister 70 % des nouveaux cas à frottis positif et traiter avec succès 85 % d'entre eux ; d'ici 2015, arrêter l'augmentation de l'incidence et commencer à inverser la tendance; entre 1990 et 2015, diminuer de moitié le taux de prévalence de la tuberculose et le taux de mortalité.

## Améliorer le dépistage et le traitement

- **3.** Au total, 199 pays ont présenté à l'OMS un rapport sur leur stratégie pour lutter contre la tuberculose, sur le nombre de cas de tuberculose notifiés et/ou les résultats du traitement.
- **4.** Nous avons calculé, en utilisant les données de surveillance et d'enquête pour établir de nouvelles estimations de l'incidence, qu'il y a eu 8,8 millions de nouveaux cas de tuberculose en 2003 (140 pour 100 000 habitants), dont 3,9 millions (62 pour 100 000) avaient un frottis positif et 674 000 (11 pour 100 000) étaient porteurs

- du virus de l'immunodéficience humaine (VIH). Le nombre total de cas était de 15,4 millions (245 pour 100 000), sur lesquels 6,9 millions avaient un frottis positif (109 pour 100 000). Le nombre de décès dus à la tuberculose en 2003 est estimé à 1,7 millions (28/100 000); ce chiffre englobe les cas de co-infection tuberculose-VIH (229 000).
- **5.** En 2003, 182 pays au total appliquaient la stratégie DOTS, soit 2 de plus qu'en 2002. A la fin de 2003, 77 % de la population mondiale vivaient dans des pays, ou des régions de pays, où la stratégie était appliquée. Les programmes ont rapporté 3,7 millions de cas nouveaux et de rechutes, parmi lesquels on recense 1,8 millions de cas nouveaux à frottis positif. Au total, 17,1 millions de tuberculeux et 8,6 millions de sujets à frottis positif ont suivi un traitement dans le cadre des programmes DOTS entre 1995 et 2003.
- 6. Les 1,8 million de cas à frottis positif signalés par les programmes DOTS en 2003 représentent un taux de détection de 45 %. L'augmentation du nombre de cas à frottis positif notifiés dans le cadre de la stratégie DOTS n'a jamais été aussi forte qu'entre 2002 et 2003 (324 000) (l'augmentation annuelle moyenne entre 1995 et 2000 était de 134 000). L'augmentation des cas notifiés a été plus marquée encore pour tous les cas de tuberculose confondus : elle a été de 693 000 entre 2002 et 2003, alors que l'augmentation annuelle moyenne était de 270 000 pendant la période 1995-2000.
- 7. Alors que l'augmentation du nombre de cas de tuberculose notifiés par les programmes DOTS semble s'accélérer depuis 2000 le nombre total de cas notifiés à l'OMS (toutes formes et toutes sources confondues) n'a que très peu augmenté entre 1995 et 2003 (taux moyen de notification de 42 %). Le nombre de cas à frottis positif raporté par toutes les

- sources a augmenté (taux de notification de 50 % en 2003), mais bien plus lentement que la hausse dont il est fait état dans le cadre de la stratégie DOTS.
- 8. Deux pays, l'Inde et la Chine, concentraient à eux seuls 63 % de tous les cas supplémentaires à frottis positif signalés dans le cadre de la stratégie DOTS en 2003. Parmi les personnes qui ont développé une tuberculose à crachat positif en 2003 (nombre estimé) mais qui n'ont pas été détectées par les programmes DOTS, 67% habitaient dans seulement 8 pays: le Bangladesh, la Chine, l'Ethiopie, la Fédération de Russie, l'Inde, l'Indonésie, le Nigéria et le Pakistan.
- 9. Alors que les programmes DOTS se sont étendus géographiquement, le taux de détection des cas nouveaux à frottis positif dans les zones couvertes par la stratégie DOTS est resté relativement constant depuis 1995 (52 % en moyenne), encore qu'on observe une légère hausse dans les pays les plus touchés, en particulier au Bangladesh, en Inde, au Myanmar et aux Philippines.
- 10. Le taux de succès thérapeutique dans la cohorte DOTS de 2002 était de 82 % en moyenne, inchangé depuis 2000. Comme pour les années précédentes, il était nettement inférieur à la moyenne dans la Région africaine (73 %) et dans la Région européenne (76 %). Cela s'explique en partie par les complications de la co-infection tuberculose-VIH dans la Région africaine et par la pharmacorésistance dans la Région européenne. Un autre facteur tout aussi important dans ces deux régions est l'ncapacité des programmes DOTS à documenter les résultats du traitement pour tous les patients.
- **11.** D'après les cas déclarés et les estimations de l'OMS, 22 pays avaient atteint à la fin de 2003 les objectifs en matière de détection et de succès

thérapeutique. Le Viet Nam est cependant le seul pays du groupe des pays les plus touchés1 parmi eux, mais le Cambodge, le Myanmar et les Philippines ne sont pas loin d'atteindre les objectifs.

## Tendances épidémiologiques et impact de la stratégie DOTS

12. En 2003, le taux d'incidence de la tuberculose fléchissait ou se stabilisait dans cinq des six Régions de I'OMS, mais augmentait de 1,0 % à l'échelle mondiale. La région qui fait exception est la Région africaine, où l'incidence a augmenté plus rapidement dans les pays à plus haut taux de prévalence VIH. En Europe de l'Est, les taux d'incidence ont augmenté pendant les années 90 pour atteindre un pic vers 2001 et ont baissé depuis. La hausse de l'incidence mondiale ralentit parce que les épidémies d'infection au VIH ralentissent en Afrique, mais on ignore encore quand l'incidence mondiale de la tuberculose commencera à décroître.

13. Nous calculons que, suite à l'extension de la stratégie DOTS entre 1990 et 2003, le taux de prévalence mondial est passé de 309 à 245 pour 100 000 (cas de co-infection tuberculose-VIH compris), et a diminué de 5 % entre 2002 et 2003 alors que l'incidence continuait d'augmenter. Le taux de mortalité mondial a atteint un record dans les années 90, puis a diminué à 2,5 % (cas de co-infection tuberculose-VIH compris) ou 3,5 % par an (cas de co-infection tuberculose-VIH non compris) entre 2002 et 2003. Si les tendances n'étaient pas si contraires en Afrique, les taux de prévalence et de mortalité baisseraient beaucoup plus rapidement à l'échelle mondiale

## Planification et mise en oeuvre de la stratégie DOTS

**14.** Tous les pays les plus touchés ont un plan stratégique d'extension de la stratégie DOTS et beaucoup d'entre

eux commenceront en 2005 un nouveau cycle de planification, de préférence axé sur l'année cible des OMD, 2015. Bien que les systèmes de santé de nombreux pays subissent encore des réformes et des restructurations, tous les pays les plus touchés, sauf la Fédération de Russie et la Thaïlande, indiquent que les fonctions de lutte antituberculeuse sont entièrement intégrées aux services de santé nationaux généraux.

15. L'extension de la stratégie DOTS se heurte à cinq obstacles d'une importance capitale : pénurie de personnel qualifié, absence d'engagement politique, insuffisance des services de laboratoire, prise en charge inadéquate de la tuberculose multirésistante et de la tuberculose associée au VIH. En ce qui concerne la pharmacorésistance, les pays sont peu nombreux à avoir une politique nationale en matière de diagnostic et de traitement de la tuberculose résistante, et même dans ceux qui en ont une, le traitement n'est souvent pas conforme aux normes acceptables. En ce qui concerne la co-infection tuberculose-VIH, les programmes nationaux de lutte antituberculeuse indiquent que peu de tuberculeux ont un test de dépistage du VIH (3 % des cas notifiés), qu'ils sont moins nombreux encore à être examinés en vue de bénéficier d'un traitement antirétroviral et qu'une très faible proportion commence un traitement de ce type (1347 patients recensés en 2003). Le rapport discute une série de mesures pour remédier à cette situation.

16. L'intensification du soutien et de l'action dans les pays (ISAC) est une nouvelle initiative destinée à catalyser et à accélérer l'extension de la stratégie DOTS en vue d'atteindre les objectifs de 2005. L'initiative a pour but d'améliorer les capacités techniques afin de faciliter l'utilisation des subventions importantes provenant du Fonds Mondial de lutte contre le SIDA, la Tuberculose et le Paludisme (FMSTP) et autres bailleurs de fonds importants. En 2004, ont participé à l'initiative la Chine, la Fédération de Russie, l'Inde, l'Indonésie, le Kenya, l'Ouganda, le Pakistan et la Roumanie.

- 17. Les contributions croissantes des organisations non gouvernementales (ONG) et de groupes communautaires manifestent clairement l'engagement de plus en plus important de la société civile en faveur de la lutte antituberculeuse. L'action de ces groupes place les patients au centre de la stratégie DOTS et améliore l'accès aux services antituberculeux dans les zones éloignées et au sein des populations défavorisées et marginalisées.
- 18. Les projets public-privé et publicpublic ont des effets mesurables sur le dépistage des cas dans plusieurs pays d'Asie, et pourraient être un moyen d'élargir les services de lutte antituberculeuse dans les villes afri-

## Financement de l'extension des programmes DOTS

- 19. Des données financières ont été reçues de 134 pays sur 211 (64 %), contre 123 en 2003 ; 70 et 69 pays ont fourni des données complètes concernant respectivement le budget et les dépenses. Les 22 pays les plus touchés ont tous fourni des données sauf l'Afrique du Sud.
- 20. On a observé une forte augmentation des budgets des PNT ainsi qu'un net accroissement des fonds disponibles pour la lutte antituberculeuse dans les pays les plus touchés depuis 2002, avec des augmentations particulièrement importantes entre 2003 et 2004. Les budgets PNT prévus pour 2005 atteignent au total US \$741 millions. La projection des coûts totaux estimés de la lutte antituberculeuse s'élève à US \$1,3 milliard en 2005 et les fonds disponibles à US \$1,2 milliard. La forte augmentation des crédits disponibles est pratiquement entièrement due à des fonds publics supplémentaires aloués par la Chine, la Fédération de Russie et l'Indonésie et à des subventions du fond mondial (FMSTP).
- 21. Les pays dont le budget du PNT pour 2005 est le plus important sont la Chine, la Fédération de Russie, l'Inde et l'Indonésie. Si l'on prend également en considération des coûts non inclus dans les budgets PNT, l'Afrique du Sud, la Chine, la Fédération

<sup>&</sup>lt;sup>1</sup> Le Pérou a été exclu du groupe initial des pays les plus touchés car ce pays a atteint les objectifs de détection et de succès thérapeutique et a réussi à réduire l'incidence de la tuberculose.

de Russie et l'Inde représentent à elles seules US \$946 millions (73 %) du montant total de US \$1,3 milliard. Huit pays parmi les plus touchés prévoient des coûts totaux de US \$20–50 millions en 2005 ; le reste représente US \$18 millions ou moins.

- 22. On observe des variations considérables, par patient traité, dans les budgets pour les médicaments de première ligne, dans les budgets PNT totaux et dans les coûts totaux pour chaque année entre 2002 et 2005. Parmi les pays les plus touchés, le budget PNT par patient est le plus faible en Inde (US \$34). Dans la plupart des pays, les budgets se situent entre US \$100 et 200 par patient et les coûts entre US \$150 et 300. L'Afrique du Sud et la Fédération de Russie sont des exceptions notables, avec des coûts par patient traité supérieurs à US \$1000. Les budgets par patient traité sont généralement stables ou en augmentation et, par conséquent, les coûts par patient traité le sont aussi.
- **23.** En 2005, les gouvernements des pays les plus touchés financent 62 % des budgets des PNT (y compris par des prêts), le FMSTP 15 % et 7 % provient d'autres sources, ce qui correspond à un déficit équivalent à 16 % des budgets prévus. Les gouvernements des pays les plus touchés contribuent davantage (79 %) aux coûts totaux qu'aux budgets PNT parce qu'ils financent le personnel des services de santé généraux et les infrastructures utilisées pour la lutte antituberculeuse. Les contributions moyennes au financement de la lutte antituberculeuse des gouvernements des pays les plus touchés sont élevées et masquent le fait que nombre de ces pays sont largement dépendants de subventions.
- 24. Malgré des progrès dans la mobilisation de crédits supplémentaires, les pays les plus touchés ont signalé un déficit de financement de US \$119 millions en 2005. Ce chiffre est plus élevé que ceux qui avaient été enregistrés en 2003 et 2004. Les plus importants déficits de financement sont signalés par la Chine, la Fédération de Russie, l'Inde, le Pakistan et

le Zimbabwe (US \$93 millions, soit 78 % du déficit total). Proportionnellement aux budgets, les déficits les plus importants sont ceux du Kenya, du Nigéria, de l'Ouganda, du Pakistan et du Zimbabwe.

- 25. Les activités planifiées ne sont pas en mesure d'atteindre les objectifs fixés pour le dépistage des cas en 2005 dans 12 pays. En outre, les budgets pour les activités concertées contre la tuberculose et le VIH et pour les médicaments de deuxième ligne pour traiter la tuberculose multirésistante sont actuellement peu élévés. Cela veut dire que l'on peut considérer que les déficits actuellement rapportés par les PNT sont sousestimés et que les ressources nécessaires pour la tuberculose seront dans le futur plus élevées que US \$1,3 milliard.
- **26.** La capacité d'absorption est l'un des grands problèmes pour les pays les plus touchés qui sont parvenus à mobiliser un important financement supplémentaire. En 2003, les dépenses ont été inférieures au financement disponible ; reste à évaluer si les PNT peuvent effectivement dépenser les crédits supplémentaires disponibles en 2004 et 2005.
- 27. En termes financiers, les pays les plus touchés entrent dans quatre catégories : a) quatre pays (l'Inde, le Myanmar, les Philippines et le Viet Nam) dont les budgets devraient permettre d'atteindre les objectifs de 2005, et qui auront sans doute un déficit de financement minime, voire nul; b) quatre pays dont les budgets sont suffisants, mais qui devront combler des déficits de financement (Cambodge, Chine), ou qui ne savent pas très bien combien de cas supplémentaires seront détectés et traités avec succès grâce aux fonds supplémentaires importants désormais disponibles (Bangladesh, Indonésie); c) cinq pays dont les plans ne sont pas de nature à leur permettre d'atteindre les objectifs de 2005, mais qui signalent des déficits de financement minimes ou nuls; d) neuf pays qui signalent d'importants déficits de financement et dont les plans sont loin d'être de nature à leur permettre d'atteindre les

objectifs de détection des cas (huit pays) et/ou dont on ne sait pas s'ils seront suffisants pour atteindre l'objectif de succès thérapeutique. Ces neuf pays méritent une attention particulière de la part des donateurs et d'autres organismes d'aide.

## Progrès vers la réalisation des objectifs du Millénaire pour le développement

- 28. Si l'amélioration de la détection des cas observée entre 2002 et 2003 peut être maintenue, le taux de détection des cas sera de 60 % en 2005. Pour atteindre l'objectif de 70 %, les programmes DOTS doivent recruter des patients tuberculeux dans les centres de santé et les hôpitaux qui ne participent pas encore aux programmes, notamment dans le secteur privé en Asie, et au-delà des limites actuelles des systèmes de santé publique en Afrique. Pour atteindre l'objectif de 85 % de succès thérapeutique, un effort particulier doit être fait afin d'améliorer les taux de guérison en Afrique et en Europe de l'Est.
- **29.** Notre analyse des tendances épidémiologiques laisse supposer que le taux d'incidence de la tuberculose est encore en légère augmentation dans le monde, mais que les taux de prévalence et de mortalité sont en diminution. Quant à savoir si la diminution du poids de la tuberculose sera suffisante pour atteindre les OMD d'ici 2015, dépendra de la rapidité avec laquelle les programmes DOTS seront mis en œuvre par les divers prestataires de soins, et de l'efficacité avec laquelle les programmes seront adaptés pour répondre aux problèmes que présentent la co-infection tuberculose-VIH (notamment en Afrique) et la pharmacorésistance (notamment en Europe de l'Est).
- **30.** Le financement de l'effort mondial de lutte antituberculeuse s'est amélioré depuis 2002, de façon spectaculaire dans certains pays. Certains des pays les plus touchés disposent désormais de fonds suffisants pour atteindre les objectifs, mais doivent encore montrer qu'ils sont capables de les utiliser efficacement; certains n'ont pas de déficit apparent, mais

## Principaux indicateurs epidemiologiques et financiers

INDICATEURS ÉPIDÉMIOLOGIQUES (MONDE)	CIBLE OMD	ANNÉE CIBLE	ESTIMATION 2003	EVOLUTION PAR RAPPORT À 2003 (%)	ANNÉE DE RÉFÉRENCE
DOTS détection des cas (%)	70	2005	45	+7,5	2002
DOTS succès thérapeutique (%)	85	2005	82 (cohorte 2002)	0,0	cohorte 2001
Taux d'incidence (pour 100 000 par an, VIH exclus)	En diminution	2015	129	+0,6	2002
Taux d'incidence (pour 100 000 par an, VIH inclus*)			140	+1,0	2002
Taux de prévalence (pour 100 000, VIH exclus)	moitié du niveau de 1990	2015	240	-22	1990
Taux de prévalence (pour 100 000, VIH inclus)			245	-21	1990
Taux de mortalité (pour 100 000 par an, VIH exclus)	moitié du niveau 1990	2015	24	-12	1990
Taux de mortalité (pour 100 000 par an, VIH inclus)			28	-1,6	1990
INDICATEURS FINANCIERS (PAYS LES PLUS TOUCHÉS)			ESTIMATION 2005	EVOLUTION 2002-2005 (%)	ANNÉE DE RÉFÉRENCE
Dépenses totales pour la lutte antituberculeuse (US \$	millions)		1321	+49	2002
Budget PNT pour la lutte antituberculeuse (US \$ millio	ns)		741	+79	2002
Total des fonds disponibles pour la lutte contre la tube	erculose (US \$ millio	ons)	1202	+36	2002
Etat (à l'exclusion des prêts)			982	+26	2002
Prêts			56	+102	2002
Subventions (à l'exclusion du FMSTP)			55	+29	2002
FMSTP			109	NA	2002
Déficit de financement tel que raporté par les PNT (US Coûts par patient (US \$) (valeurs médianes)	\$\$ millions)		119	+34	2002
Coût total			213	+22	2002

VIH inclus: y compris les patients souffrant à la fois de tuberculose et d'une infection à VIH; les indicateurs OMD pour la tuberculose excluent les patients également atteints d'infection à VIH cependant ces statistiques sont également utiles dans la lutte antituberculeuse.

NA: non applicable car les fonds ont été distribués pour la première fois en 2003

Budget pour les médicaments de première ligne

devraient vérifier que leurs budgets sont suffisants; certains ont un déficit de financement évident et doivent se concentrer sur la mobilisation des fonds nécessaires pour améliorer la performance du programme.

**Budget PNT** 

+45

-12

133 28 2002

2002

## Resumen

### Antecedentes y métodos

- **1.** El noveno informe anual de la Organización Mundial de la Salud (OMS) sobre vigilancia, planificación y financiación de la lucha contra la tuberculosis (TB) incluye datos sobre las notificaciones de casos y los resultados del tratamiento procedentes de todos los programas nacionales de lucha contra la TB (PNT) que han informado a la OMS, así como un análisis de los planes, presupuestos y gastos, y de los progresos de la expansión de la estrategia DOTS en 22 países con alta carga de TB (PACT).
- 2. En la actualidad se dispone de datos reunidos durante diez años consecutivos (1994-2003), que permiten evaluar los progresos realizados para alcanzar los Objetivos de Desarrollo del Milenio (ODM) relativos a la lucha contra la TB. Las cinco metas de los ODM que guardan relación directa con la lucha antituberculosa son: para 2005, detectar el 70% de los nuevos casos bacilíferos y tratar con éxito el 85% de esos casos; para 2015, haber detenido y comenzado a reducir la incidencia; entre 1990 y 2015, reducir a la mitad las tasas de prevalencia y de mortalidad de la TB.

## Mejorar la detección y el tratamiento de los casos

- **3.** Un total de 199 países han informado a la OMS de sus estrategias de lucha contra la TB, así como de las notificaciones de casos y/o de los resultados del tratamiento.
- **4.** Tras actualizar las estimaciones de la incidencia tomando como base los datos de la vigilancia y de las encuestas, hemos calculado que en 2003 hubo 8,8 millones de nuevos casos de TB (140/100 000 habitantes), de los cuales 3,9 millones (62/100 000) eran bacilíferos y 674 000 (11/100 000) estaban infectados por el virus de la inmunodeficiencia humana (VIH). Hubo 15,4 millones de casos prevalentes (245/100 000), de

- los cuales 6,9 millones eran bacilíferos (109/100 000). Se estima que 1,7 millones de personas (28/100 000) murieron de TB en 2003, incluidos los casos de coinfección por el VIH (229 000).
- **5.** Ciento ochenta y dos países aplicaron la estrategia DOTS en 2003, dos más que en 2002. A finales de 2003, el 77% de la población mundial vivía en países (o regiones de países) que disponían de cobertura de DOTS. Los programas DOTS notificaron 3,7 millones de casos de TB nuevos y recidivantes, de los cuales 1,8 millones eran nuevos bacilíferos. Entre 1995 y 2003, 17,1 millones de pacientes con TB y 8,6 millones de pacientes bacilíferos recibieron tratamiento en los programas DOTS.
- **6.** Los 1,8 millones de casos bacilíferos notificados por los programas DOTS en 2003 representan una tasa de detección del 45%. El aumento de los casos bacilíferos notificados en el ámbito de los programas DOTS entre 2002 y 2003 (324 000) fue mayor que nunca (el incremento medio anual entre 1995 y 2000 había sido de 134 000). El aumento de las notificaciones fue todavía mayor si se consideran todos los casos de TB: 693 000 entre 2002 y 2003, en comparación con un incremento medio anual de 270 000 en el periodo 1995–2000.
- 7. Mientras que el número de casos de TB notificados por los programas DOTS parece haber crecido de forma acelerada desde 2000, el número total de casos de TB notificados a la OMS (todas las formas, de todas las fuentes) aumentó muy poco entre 1995 y 2003 (tasa media de detección del 42%). El número de casos bacilíferos notificados por la totalidad de las fuentes ha ido en aumento (tasa de detección del 50% en 2003), pero mucho más lentamente que los notificados en el marco del DOTS.
- **8.** El 63% de los casos bacilíferos adicionales notificados a través de

- DOTS en 2003 provenían de tan sólo dos países: China e India. Dos tercios (67%) de los nuevos casos estimados para 2003 que no fueron detectados por medio de los programas DOTS procedían de ocho países: Bangladesh, China, Etiopía, la Federación de Rusia, la India, Indonesia, Nigeria y Pakistán.
- **9.** A medida que los programas DOTS se han extendido geográficamente, la tasa de detección de nuevos casos bacilíferos en las zonas donde se aplica la estrategia DOTS ha permanecido prácticamente constante desde 1995 (media del 52%), aunque hay signos de un lento aumento en los PACT, sobre todo en Bangladesh, Filipinas, la India y Myanmar.
- 10. La tasa media de éxito del tratamiento en la cohorte de DOTS de 2002 fue del 82%, la misma que se viene observando desde 2000. Como en años anteriores, dicha tasa fue considerablemente inferior a la media en las regiones de África (73%) y Europa (76%). Las bajas tasas de éxito del tratamiento en esas dos regiones pueden atribuirse en parte a la coinfección por el VIH y a la farmacorresistencia, respectivamente. Sin embargo, igualmente importante es el fracaso de los programas DOTS en la vigilancia de los resultados del tratamiento en todos los pacientes en esas dos regiones.
- **11.** Con base en los casos notificados y las estimaciones de la OMS, 22 países habían alcanzado a finales de 2003 las metas fijadas en materia de detección de casos y éxito del tratamiento. Viet Nam era aún el único miembro del actual grupo de PACT¹ entre ellos, aunque Camboya, Filipinas y Myanmar están a punto de lograrlo.

Perú ha sido excluido del grupo original de PACT, ya que ha alcanzado la metas y la incidencia ha disminuido.

## Tendencias epidemiológicas e impacto de la estrategia DOTS

- 12. En 2003, la tasa de incidencia de TB estaba disminuyendo o era estable en cinco de las seis regiones de la OMS, pero aumentando en todo el mundo a razón de 1,0% al año. La excepción fue la región de África, donde la incidencia ha aumentado con mayor rapidez en los países con mayores tasas de prevalencia de infección por VIH. En Europa Oriental, las tasas de incidencia aumentaron en la década de los noventa, pero alcanzaron su valor máximo en 2001, y desde entonces han disminuido. El aumento de la incidencia mundial se está haciendo más lento debido a la desaceleración de la epidemia de VIH en África, pero aún no está claro cuándo comenzará a disminuir la tasa de incidencia mundial.
- 13. Hemos calculado que, debido a la expansión de la estrategia DOTS entre 1990 y 2003, la tasa mundial de prevalencia de TB disminuyó de 309 a 245 por 100 000 (incluidos los pacientes tuberculosos con VIH), y en un 5% entre 2002 y 2003, aun cuando la incidencia siguió aumentando. La tasa mundial de mortalidad alcanzó su valor máximo en la década de los noventa y disminuyó al 2,5% (incluidos los pacientes VIH-positivos con TB) o al 3,5% anual (excluidos los pacientes VIH-positivos) entre 2002 y 2003. De no ser por las tendencias extremadamente adversas que se observan en África, las tasas de prevalencia y de mortalidad estarían disminuyendo más rápidamente en todo el mundo.

## Planificación y aplicación de la estrategia DOTS

14. Todos los PACT disponen de un plan estratégico de expansión de la estrategia DOTS; en 2005, muchos comenzarán un nuevo ciclo de planificación con miras a alcanzar la meta de 2015 fijada por los ODM. Si bien los sistemas de salud de numerosos países todavía son objeto de reformas y de reestructuración, todos los PACT, salvo la Federación de Rusia y Tailandia, informaron que las funciones relacionadas con la lucha antituberculosa están completamente integradas en los servicios de salud esenciales de la nación.

- 15. Entre los obstáculos con que se enfrenta la expansión de la estrategia DOTS, hay cinco de importancia capital: la escasez de personal capacitado, la falta de compromiso político. la debilidad de los servicios de laboratorio y la gestión inadecuada de la tuberculosis multirresistente (MDR-TB) y de la TB asociada al VIH. Con respecto a la farmacorresistencia, pocos países cuentan con políticas nacionales para el diagnóstico y el tratamiento de la MDR-TB, e incluso en aquellos que disponen de ellas, el tratamiento no suele estar a la altura del nivel exigido. Por lo que se refiere a TB-VIH, los PNT informaron que son pocos los pacientes con TB sometidos a pruebas de detección del VIH (el 3% de los casos notificados), aún menos los evaluados con vistas a la administración de tratamiento antirretrovírico y que sólo una fracción pequeña inicia dicho tratamiento (1347 pacientes en 2003). Este informe examina un amplio abanico de medidas correctivas para superar dichos obstáculos.
- 16. ISAC (actuaciones y apoyo intensificados en los países) es una nueva iniciativa destinada a catalizar y acelerar la expansión de DOTS con miras a las metas de 2005. Su objetivo consiste en mejorar la capacidad técnica para facilitar el gasto de grandes subsidios del Fondo Mundial de Lucha contra el SIDA, la Tuberculosis y la Malaria (FMSTM) y de otros donantes importantes. En 2004, los participates fueron China, la Federación de Rusia, la India, Indonesia, Kenya, Pakistán, Rumania y Uganda.
- 17. Las contribuciones cada vez más importantes de las organizaciones no gubernamentales y de los grupos comunitarios constituyen una clara manifestación del compromiso creciente de la sociedad civil en la lucha contra la TB. El trabajo de esos grupos sitúa a los pacientes en el centro de la estrategia DOTS y mejora el acceso a los servicios relacionados con la TB en zonas remotas y entre las poblaciones desfavorecidas y marginadas.

18. Los proyectos mixtos de carácter publicoprivado y público-público están ejerciendo un impacto perceptible en la detección de casos en varios países asiáticos y podrían llegar a constituir un mecanismo de expansión de los servicios de lucha contra la TB en las ciudades africanas.

## Financiación de la expansión de la estrategia DOTS

- 19. Se ha recibido información financiera de 134 países sobre un total de 211 (64%), en comparación con 123 en 2003. Han presentado datos completos en materia de presupuesto y gasto 70 y 69 países, respectivamente. Se recibieron datos de los 22 PACT, con excepción de Sudáfrica.
- 20. Desde 2002 ha habido un gran aumento de los presupuestos de los PNT y de la financiación disponible para la lucha antituberculosa en los PACT, en particular entre 2003 y 2004. Los presupuestos de los PNT previstos para 2005 ascienden a US\$ 741 millones. Se calcula que los costos totales de la lucha contra la TB en 2005 serán de US\$ 1,3 mil millones, y los fondos disponibles son de US\$ 1,2 mil millones. El gran aumento de fondos disponibles se debe casi por completo a nuevos recursos proporcionados por los gobiernos de China, la Federación de Rusia e Indonesia, así como a subsidios del FMSTM.
- 21. Los países que disponen de mayores presupuestos para sus PNT en 2005 son China, la Federación de Rusia, la India e Indonesia. Si también se toman en consideración los costos que no figuran en los presupuestos de los PNT, los costos de China, la Federación de Rusia, la India y Sudáfrica reflejan el 73% del costo total (US\$ 946 millones de US\$ 1,3 mil millones). En otros ocho PACT, los costos totales oscilan entre US\$ 20 y US\$ 50 millones, y en el resto de los PACT ascienden a US\$ 18 millones, o menos.
- 22. Por paciente tratado, hay variaciones considerables en los presupuestos destinados a medicamentos de primera línea, en los presupuestos de

los PNT y en los costos totales en cada uno de los años del período 2002-2005. Entre los PACT, la India es el país con menor presupuesto de PNT por paciente (US\$ 34). La mayoría de los países tienen presupuestos que van de US\$ 100 a US\$ 200 por paciente, y costos que varían entre US\$ 150 y US\$ 300. La Federación de Rusia y Sudáfrica constituyen excepciones notables, con costos por paciente tratado que superan los US\$ 1000. En general, los presupuestos por paciente tratado son estables o tienden a aumentar, de modo que los costos por paciente tratado también son generalmente estables o tienden al alza.

- 23. En 2005, el 62% de los presupuestos de los PNT (incluidos los préstamos) será proporcionado por los gobiernos de los PACT, el 15% por el FMSTM, el 7% por subsidios de otras fuentes, con lo que queda un déficit del 16% con respecto a los presupuestos notificados. Los gobiernos de los PACT contribuven más a los costos totales (79%) que a los presupuestos de los PNT, pues financian el personal y las infraestructuras de los servicios de salud generales utilizados en la lucha contra la tuberculosis. La elevada contribución media de los gobiernos de los PACT a la financiación de la lucha antituberculosa oculta el hecho de que muchos de esos países dependen en gran medida de la financiación bajo la forma de subsidios.
- 24. A pesar de los progresos realizados en la obtención de fondos adicionales, los PACT acusan un déficit financiero de US\$ 119 millones en 2005, cifra que es superior a las registradas en 2003 y 2004. Los mayores déficit corresponden a China, la Federación de Rusia, la India, Pakistán y Zimbabwe (US\$ 93 millones, es decir, el 78% del déficit total). Proporcionalmente a los presupuestos, los mayores déficit corresponden a Kenya, Nigeria, Pakistán, Uganda y Zimbabwe.
- 25. En 12 países las actividades planificadas no son compatibles con el logro de la meta de detección de casos para 2005. Además, los presupuestos actuales para las activida-

des de colaboración TB-VIH y para los medicamentos de segunda línea para el tratamiento MDR-TB son pequeños. Esto significa que los déficit notificados por los PNT pueden estar subestimados, y que el total de recursos necesarios para el control de TB será superior a US\$ 1,3 mil millones en el futuro.

- 26. Una cuestión fundamental para los PACT que han conseguido cuantiosos fondos adicionales es su capacidad de absorberlos. En 2003, los gastos fueron inferiores a los fondos disponibles, y queda por ver si los PNT pueden gastar eficazmente el dinero extra disponible en 2004 y 2005.
- 27. En materia de financiación, los PACT pueden clasificarse en cuatro categorías: a) cuatro países cuyos presupuestos son compatibles con el logro de las metas para 2005 y que probablemente no tendrán déficit de fondos o, en el caso de que los tengan, serán mínimos (Filipinas, la India, Myanmar, y Viet Nam); b) cuatro países cuyos presupuestos son suficientes, pero que tendrán que encontrar la forma de completar los fondos que les faltan (Camboya y China) o en los que no está claro cuántos casos adicionales se detectarán y tratarán con éxito como resultado de los considerables fondos adicionales de que disponen actualmente (Bangladesh e Indonesia); c) cinco países cuyos planes no se ajustan al logro de las metas para 2005, pero que tienen un pequeño o nulo déficit de fondos, y d) nueve países con un gran déficit de fondos y cuyos planes están por debajo de lo necesario para alcanzar las metas de detección de casos (ocho países) y/o en los que no está claro si los fondos son suficientes para alcanzar el objetivo del éxito del tratamiento. Estos nueve países merecen especial atención por parte de los organismos donantes y de otros organismos de apoyo.

## Progresos en la consecución de los Objetivos de Desarrollo del Milenio

28. Si se mantiene la mejora en la detección de casos que se produjo entre 2002 y 2003, la tasa de detección de casos será del 60% en 2005. Para alcanzar la meta del 70%, los programas DOTS deben reclutar pacientes con TB de clínicas y hospitales que no participan en esos programas, especialmente los del sector privado en Asia, y los que están fuera de los límites actuales de los sistemas de salud pública en África. Para lograr la meta del 85% de éxito del tratamiento habrá que hacer un esfuerzo especial por mejorar las tasas de curación en África y Europa Oriental.

- 29. Nuestro análisis de las tendencias epidemiológicas indica que la tasa de incidencia de la TB sigue aumentando lentamente en todo el mundo, pero que las tasas de prevalencia y de mortalidad están descendiendo. Que la carga de TB pueda disminuir lo suficiente como para alcanzar los ODM en 2015 dependerá de la rapidez con que los diversos prestadores de atención de salud puedan poner en marcha los programas DOTS, y de cuán eficazmente se puedan adaptar esos programas para hacer frente a los retos que suponen la coinfección por VIH (especialmente en Africa) y a la farmacorresistencia (especialmente en Europa Oriental).
- 30. La financiación de la lucha mundial contra la TB ha mejorado desde 2002, y en algunos países lo ha hecho de forma espectacular. Algunos PACT disponen ahora de fondos suficientes para alcanzar las metas, pero deben demostrar que son capaces de utilizarlos de forma eficaz, otros no tienen déficit aparente, pero deben comprobar que disponen de suficiente presupuesto, y otros presentan un déficit financiero evidente y deben centrarse en conseguir el dinero necesario para mejorar el rendimiento del programa.

## Principales indicadores epidemiológicos y financieros

INDICADORES EPIDEMIOLÓGICOS (NIVEL MUNDIAL)	META DE LOS ODM	AÑO DE Consecución Previsto	I ESTIMACIÓN 2003	CAMBIO, DEL AÑO DE REFERENCIA A 2003 (%)	AÑO DE REFERENCIA
Detección de casos bajo DOTS (%)	70	2005	45	+7,5	2002
Éxito del tratamiento bajo DOTS (%)	85	2005	82 (cohorte de 2002)	0,0	2001
Tasa de incidencia (por 100 000 por año, excluido VIH)	En descenso	2015	129	+0,6	2002
Tasa de incidencia (por 100 000 por año, incluido VIH)*			140	+1,0	2002
Tasa de prevalencia (por 100 000, excluido VIH)	Mitad del nivel de 1990	2015	240	-22	1990
Tasa de prevalencia (por 100 000, incluido VIH)			245	-21	1990
Tasa de mortalidad (por 100 000 por año, excluido VIH)	Mitad del nivel de 1990	2015	24	-12	1990
Tasa de mortalidad (por 100 000 por año, incluido VIH)			28	-1,6	1990

INDICADORES FINANCIEROS (PACT)	ESTIMACIÓN 2005	CAMBIO, DE 2002 A 2005 (%)	AÑO DE Referencia
Costos totales del control de la TB (en millones de US\$)	1321	+49	2002
Presupuestos de los PNT para el control de la TB (en millones de US\$)	741	+79	2002
Fondos disponibles totales para el control de la TB (en millones de US\$)	1202	+36	2002
Gobierno (excluidos préstamos)	982	+26	2002
Préstamos	56	+102	2002
Subsidios (excluidos los del FMSTM)	55	+29	2002
FMSTM	109	NA	2002
Déficit financiero según los PNT (en millones de US\$)	119	+34	2002
Costos por paciente (US\$) (valores medianos)			
Costo total	213	+22	2002
Presupuesto de los PNT	133	+45	2002
Presupuesto para medicamentos de primera línea	28	-12	2002

<sup>\*</sup> Incluido VIH: incluidos los pacientes con TB VIH-positivos; los indicadores de los ODM para la TB excluyen a los pacientes VIH-positivos, pero estas estadísticas también son útiles en el control de la TB.

NA: no aplicable, puesto que los recursos fueron distribuidos por primera vez en 2003.

## Introduction

The goal of this series of annual reports is to chart progress in global TB control and, in particular, to evaluate progress in implementing the DOTS strategy.1,2 The first targets set for global TB control were ratified in 1991 by WHO's World Health Assembly.3 They are to detect 70% of new smearpositive TB cases, and to successfully treat 85% of these cases. Since these targets were not reached by the end of year 2000 as originally planned, the target year was deferred to 2005.4

In 2000, the United Nations created a new framework for monitoring progress in human development, the MDGs. Among 18 MDG targets, the eighth is to "have halted by 2015 and begun to reverse the incidence of malaria and other major diseases". Although the objective is expressed in terms of incidence, the MDGs also specify that progress should be measured in terms of the reduction in TB prevalence and deaths. The target for these two indicators, based on a resolution passed at the 2000 Okinawa (Japan) summit of G8 industrialized nations, and now adopted by the Stop

TB Partnership, is to halve TB prevalence and death rates (all forms of TB) between 1990 and 2015. All three measures of impact (incidence, prevalence and death rates) have been added to the two traditional measures of DOTS implementation (case detection and treatment success), so that the MDG framework includes five principal indicators of progress in TB control. All five MDG indicators will, from now on, be evaluated by WHO's Global TB Surveillance, Planning and Financing Project. The focus is on the performance of NTPs in 22 HBCs, and in priority countries in WHO's six regions.

Some other MDGs are indirectly relevant to TB control. For example, Goal 1 is to eradicate extreme poverty and hunger, and Goal 3 is to promote gender equality and empower women. Goal 8 is to develop a global partnership for development, in which the Stop TB Partnership will have a role. A discussion of these goals is beyond the scope of this report, but further details can be found at web site: unstats.un.org/unsd.

While the MDGs set out the main objectives for global TB control, numerous specific activities must be carried out to meet these larger goals. This technical report, the ninth in the series, describes many of the details. It presents plans and budgets for DOTS expansion, and costs, expenditures and sources of funding. It also summarizes the progress made on special initiatives such as collaborative TB/HIV activities, improvements in the laboratory network and DOTS-Plus projects for the management of drug-resistant TB.

Since 1980, 81 million TB patients have been reported through WHO's surveillance system, including 17 million notified by DOTS programmes since 1995. The financial monitoring system has accounted for US\$ 4.3 billion spent on TB control in the HBCs since its inception in 2002. Thus, the Global TB Surveillance, Planning and Financing Project has become a formidable instrument for monitoring and evaluating progress in TB control.

Framework for effective tuberculosis control. Geneva, World Health Organization (WHO/TB/94.179).

<sup>&</sup>lt;sup>2</sup> An expanded DOTS framework for effective tuberculosis control. Geneva, World Health Organization (WHO/CDS/TB/ 2002.297).

<sup>&</sup>lt;sup>3</sup> Resolution WHA44.8. Tuberculosis control programme. In: Handbook of resolutions and decisions of the World Health Assembly and the Executive Board. Volume III, 3rd edition (1985-1992). Geneva, World Health Organization, 1993 (WHA44/1991/REC/1).

<sup>&</sup>lt;sup>4</sup> Stop Tuberculosis Initiative. Report by the Director-General. Fifty-third World Health Assembly. Geneva, 15-20 May 2000 (A53/5, 5 May 2000); available at http:/ /www.who.int/gb/ebwha/pdf\_files/ WHA53/ea5.pdf, accessed 11 January 2005).

## **Methods**

## Monitoring progress towards the **Millennium Development Goals**

MDGs for tuberculosis control

The MDG framework consists of a hierarchy of indicators that measure progress towards "targets", which are the specific achievements needed to satisfy higher "goals". Those most directly relevant to TB control are Goal 6 (to combat HIV/AIDS, malaria and other diseases) and Target 8 (to have halted by 2015 and begun to reverse the incidence of malaria and other major diseases, including TB). Among the indicators for Target 8 are two groups that can be used to evaluate the implementation and impact of TB control:

Indicator 23: between 1990 and 2015, to halve the prevalence and death rates associated with tuberculosis; and

Indicator 24: by 2005, to detect 70% of new smear-positive TB cases arising annually, and to successfully treat 85% of these cases.

The MDG indicators exclude HIVpositive TB patients, mainly to avoid double-counting in death statistics (deaths of HIV-positive people are recorded as AIDS deaths by WHO). However, we routinely calculate all TB indicators with and without HIVpositive TB patient, because TB control programmes need to know both.

This report focuses on the five principal indicators: incidence, prevalence, deaths, case detection and treatment success. The objective of reducing incidence is made explicit by Target 8; the targets for case detection and treatment success have been set by WHO's World Health Assembly;3 the targets for prevalence and deaths are based on a resolution of the year 2000 meeting of the Group of Eight (G8) industrialized countries, held in Okinawa, Japan.

#### Data collection and verification

Every year, WHO requests information from TB control programmes (or rel-

#### TABLE 1

#### Technical elements of the WHO TB control strategy (DOTS)<sup>a</sup>

MICROSCOPY Case detection among symptomatic patients self-reporting to health services, using sputum smear microscopy.b

**SCC/DOT** Standardized short-course chemotherapy using regimens of 6-8 months for at least all confirmed smear-positive cases. Good case management includes directly observed treatment (DOT) during the intensive phase for all new smear-positive cases, during the continuation phase of regimens containing rifampicin, and during the entirety of a re-treatment regimen.c

**DRUG SUPPLY** Establishment and maintenance of a system to supply all essential anti-tuberculosis drugs, and to ensure no interruption in their availability.

**RECORDING AND REPORTING** Establishment and maintenance of a standardized recording and reporting system, allowing assessment of treatment results (see Table 2).

- The DOTS strategy comprises five elements in all, including political commitment.
- Sputum culture is also used for diagnosis, but direct sputum smear microscopy should still be performed for all suspected cases.
- In countries that have consistently documented high treatment success rates, direct observation of treatment may be reserved for a subset of patients, as long as cohort analysis of treatment results is provided to document the outcome of all cases.

evant public health authorities) in 211 countries or territories via a standard data collection form. The latest form was distributed in mid 2004. The section dealing with monitoring and surveillance asked for the following data: TB control strategies implemented up to the end of 2003; TB case notifications in 2003; and treatment outcomes for TB patients registered during 2002, following definitions given in Table 1. The most recent form can be downloaded from www.who.int/tb.

The data collection form is a tool for collecting aggregated national data. The process of national and international reporting is quite distinct from WHO's recommendations about procedures for recording and reporting data within NTPs. The information gathered from the form includes a core set of data (questions remain more or less the same each year), plus new or timely information (questions may change from year to year). In the latest form, there are new questions about TB/HIV collaboration, about financing (the second year of collection but somewhat expanded), and about the outcomes of re-treatment, for patients who have received two or more courses of anti-TB drugs.

Completed forms are collected and reviewed at all levels of WHO - in WHO country offices, regional offices and at headquarters - and an acknowledgement form that tabulates all data submitted and shows WHO's calculations of principal indicators, is sent back to the national correspondent in order to complete any missing responses and to resolve any inconsist-

In the WHO European Region only, data collection and verification are performed jointly by the regional office and a WHO collaborating centre, EuroTB (Paris), using a different format. EuroTB subsequently publishes an annual report with additional analyses, using more detailed data for the European Region (see: www.eurotb.org).

## High-burden countries and WHO regions

Much of the data submitted to WHO is shown, country by country, in the annexes of this report. The analysis and interpretation that precedes these annexes focuses on 22 HBCs and the six WHO regions. The 22 HBCs account for approximately 80% of the estimated number of new TB cases (all forms) arising worldwide each year. These countries are the focus of intensified efforts in DOTS expansion (Annex 1). The HBCs are not necessarily those with the highest incidence

#### TABLE 2

#### Definitions of tuberculosis cases and treatment outcomes

#### A. DEFINITIONS OF TUBERCULOSIS CASES

CASE OF TUBERCULOSIS A case of TB which has been bacteriologically confirmed, or has been diagnosed by a clinician.

**DEFINITE CASE** Patient with positive culture for the *Mycobacterium tuberculosis* complex. In countries where culture is not routinely available a patient with two sputum smears positive for acid-fast bacilli (AFB+) is also considered a definite case.

PULMONARY CASE A case of TB disease involving the lung parenchyma.

SMEAR-POSITIVE PULMONARY CASE At least two initial sputum smear examinations (direct smear microscopy) AFB+; or one sputum examination AFB+ and radiographic abnormalities consistent with active pulmonary tuberculosis as determined by a clinician; or one sputum specimen AFB+ and culture positive for M. tuberculosis.

SMEAR-NEGATIVE PULMONARY CASE Pulmonary tuberculosis not meeting the above criteria for smearpositive disease. Diagnostic criteria should include: at least three sputum smear examinations negative for AFB; and radiographic abnormalities consistent with active pulmonary TB; and no response to a course of broad-spectrum antibiotics; and decision by a clinician to treat with a full course of antituberculosis therapy; or positive culture but negative AFB sputum examinations.

EXTRAPULMONARY CASE Patient with tuberculosis of organs other than the lungs e.g. pleura, lymph nodes, abdomen, genitourinary tract, skin, joints and bones, meninges. Diagnosis should be based on one culture-positive specimen, or histological or strong clinical evidence consistent with active extrapulmonary disease, followed by a decision by a clinician to treat with a full course of anti-tuberculosis chemotherapy. Note: a patient diagnosed with both pulmonary and extrapulmonary tuberculosis should be classified as a case of pulmonary tuberculosis.

NEW CASE Patient who has never had treatment for tuberculosis, or who has taken anti-tuberculosis drugs

RELAPSE CASE Patient previously declared cured but with a new episode of bacteriologically positive (sputum smear or culture) tuberculosis.b

RE-TREATMENT CASE Patient previously treated for tuberculosis, undergoing treatment for a new episode of bacteriologically positive tuberculosis.b

## **B. DEFINITIONS OF TREATMENT OUTCOMES**

(expressed as a percentage of the number registered in the cohort)

CURED Initially smear-positive patient who was smear-negative in the last month of treatment, and on at least one previous occasion.b

**COMPLETED TREATMENT** Patient who completed treatment but did not meet the criteria for cure or failure.

**DIED** Patient who died for any reason during treatment.

FAILED Smear-positive patient who remained smear-positive at five months or later during treatment.

**DEFAULTED** Patient whose treatment was interrupted for two consecutive months or more.

TRANSFERRED OUT Patient who transferred to another reporting unit and for whom the treatment outcome is not known.

SUCCESSFULLY TREATED Patients who were cured and those that completed treatment.

COHORT A group of TB cases diagnosed (and in principal notified and started on treatment) during a specified time period, e.g., the cohort of new smear-positive cases for the calendar year 2003. This group forms the denominator for calculating treatment outcomes. The sum of the above treatment outcomes, plus any cases for which no outcome is recorded (e.g. still on treatment) should equal the number registered. Some countries monitor outcomes among cohorts defined by smear and/or culture, and define cure and failure according to the best laboratory evidence available for each patient.

- Cases reported as "history unknown" in the European Region are included as new cases in this report.
- In the EuroTB database, bacteriologically positive re-treatment cases for some countries could not be distinguished from other re-treatment cases. For the purposes of this report, where this occurred, all relapse cases were included in the category "relapse", and the remainder of re-treatment cases (after default and after failure) were included as "retreatment excluding relapse" (applies to countries in the European Region only).

rates per capita; many of the latter are medium-sized African countries with high rates of TB/HIV coinfection.

The WHO regions are the African Region, the Region of the Americas, the Eastern Mediterranean Region, the European Region, the South-East Asia Region and the Western Pacific Region. All essential statistics are summarized for each of these regions and globally. However, to make clear the differences in epidemiological trends within regions, we divide the African Region into countries that have low and high rates of HIV infection (boundary at an estimated infection rate of 4% in adults aged 15-49 years), and include those countries in the Eastern Mediterranean Region which are actually on the African continent (Djibouti, Somalia and Sudan) in the low-HIV Africa group. Furthermore, we distinguish central from eastern Europe (countries of the former Soviet Union plus Bulgaria and Romania), and combine western European countries with the other established market economies. The countries within each of the resulting nine regions are listed in the legend to Figure 6.

### DOTS classification

DOTS is the internationally recommended approach to TB control. It is not simply a clinical approach to patient management, but rather a strategy for TB control primarily within public health systems. Countries reporting to WHO classify themselves as DOTS or non-DOTS, referring to the elements listed in Table 2. DOTS countries must have officially accepted and adopted the strategy, and must have implemented the essential components of DOTS in at least part of the country (Annex 2). Based on NTP responses to standard questions about policy, and usually on further discussion with the NTP, WHO accepts or revises each country's own determination of its DOTS status.

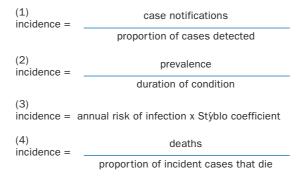
### DOTS coverage

Coverage in any country is defined as the percentage of the national population living in areas where health services have adopted DOTS. "Areas" are the lowest administrative or management units in the country - townships, districts, counties, etc. If an area (with its one or more health facilities) is considered by the NTP to be a DOTS area in 2003, then all the cases registered and reported by the NTP in that area are considered DOTS cases, and the population living within the boundaries of that area counts towards the national DOTS coverage. In some cases, treatment providers who are not following DOTS guidelines (for example private practitioners, or public health services outside the NTP such as those within prisons) notify cases to the NTP. These cases are considered non-DOTS cases, even if they are notified from within DOTS areas. However, when certain groups of patients treated by DOTS services receive special regimens or management (for example nomads placed on long-course treatment), these are considered as DOTS cases. Where possible, additional information about these special groups of patients is provided in the country notes in Annex 2.

Coverage is a crude indicator, which is easy to calculate, and which is most useful during the early stages of DOTS expansion. As a measure of patient access to diagnosis and treatment under DOTS, coverage is an approximation, and usually an overestimate. Where countries are able to provide more precise information about access to DOTS services this information is reported in the country notes of Annex 2. The case detection rate (defined below) is more precise, but also more demanding of data.

## Estimating TB incidence, prevalence and death rates

Estimates of incidence, prevalence and deaths are based on a consultative and analytical process; they are revised annually to reflect new information gathered through surveillance and from special studies, such as prevalence surveys. The details of estimation are described elsewhere.5,6 In brief, estimates of incidence (number of new cases per year) for each country are derived by one or more of four approaches, depending on the available data:



The "Stýblo coefficient" in equation (3) is taken to be a constant, with an empirically derived value in the range 40-60, relating risk of infection (%) to the incidence of smear-positive cases (per 100 000 per year). Given two of the quantities in any of these equations, we can calculate the third, and any of these formulae can be rearranged to estimate incidence, prevalence and death rates. The available data differ from country to country but include case notifications and death records (from routine surveillance and vital registration), and measures of the prevalence of infection and disease (from population-based surveys).

For each country, estimates of incidence for each year in the period 1995–2003 are made as follows. We first select a reference year for which we have a best estimate of incidence; this may be the year in which a survey was carried out, or the year in which incidence was first estimated. We then use the series of case notifications (all forms of TB) to determine how incidence changed before and after that reference year. The time series of estimated incidence rates is constructed from the notification series in two ways: if the rate of change of incidence is roughly constant through time, we fit exponential trends to the notifications; if the rate of change varies (eastern Europe, central Europe and high-HIV Africa), we use a three-year moving average of the notification rates. If the notifications for any country are considered to be an unreliable guide to trend (e.g. because reporting effort is known to have changed), we apply the aggregated trend for all other countries with reliable data from the same epidemiological region. For China, exceptionally, we have used an assessment of the trend in incidence

based on risk of infection derived from tuberculin surveys. For those countries that have no reliable data from which to assess trends in incidence (e.g. for countries such as Iraq, for which data are hard to interpret, and which are atypical within their own regions), we assume incidence is stable. Further details are available at www.who.int/tb.

For countries that have not yet measured HIV infection rates in TB patients directly, an indirect estimate can be obtained from the incidence rate ratio (IRR, the TB incidence rate in HIV-infected people divided by the incidence rate in HIV-uninfected people), as described elsewhere.6 The prevalence of MDR-TB among previously untreated TB patients has also been estimated in a separate exercise,7 supplemented with data from more recent surveys.8

Estimates of incidence form the denominator of the case detection rate. Trends in incidence are determined by underlying epidemiological processes, modified by control programmes. The impact of control on prevalence is determined by the trend in incidence, and by the estimated

<sup>&</sup>lt;sup>5</sup> Dye C et al. Global burden of tuberculosis: estimated incidence, prevalence and mortality by country. Journal of the American Medical Association, 1999, 282: 677-686.

<sup>&</sup>lt;sup>6</sup> Corbett EL et al. The growing burden of tuberculosis: global trends and interactions with the HIV epidemic. Archives of Internal Medicine, 2003, 163:1009-

<sup>&</sup>lt;sup>7</sup> Dye C et al. Worldwide incidence of multidrug-resistant tuberculosis. Journal of Infectious Diseases, 2002, 185:1197-1202.

<sup>8</sup> Anti-tuberculosis drug resistance in the world, Report No.3, WHO/IUATLD Global Project on Anti-Tuberculosis Drug Resistance Surveillance, Geneva, World Health Organization, 2004 (WHO/HTM/TB/ 2004.343).

reduction in the duration of the condition, e.g. smear-positive disease. The impact of control on deaths is determined by the trend in incidence, and by the estimated reduction in case fatality (proportion of incident cases that ever die from TB).5,6

Where population sizes are needed to calculate TB indicators, we use the latest revision of estimates provided by the United Nations Population Division,9 even though these estimates sometimes differ from those made by the countries themselves (some of which are based on more recent census data). The estimates of some TB indicators, such as the case detection rate, are derived from data and calculations that use only rates per capita, and discrepancies in population sizes do not affect these indicators. Where rates per capita are used as a basis for calculating numbers of TB cases, these discrepancies sometimes do make a difference. Some examples of important differences are given in the country notes in Annex 2.

#### Case detection rate

Smear-positive cases are the focus of DOTS programmes because they are the principal sources of infection to others, because sputum smear microscopy is a highly-specific (if somewhat insensitive) method of diagnosis, and because patients with smear-positive disease typically suffer higher rates of morbidity and mortality than smearnegative patients. As a measure of the quality of diagnosis, we calculate the proportion of new sputum smearpositive cases out of all new pulmonary cases, which has an expected value of 65-80% in areas with negligible HIV prevalence.10 However, this report presents the numbers of all TB cases notified, smear-positive and smear-negative pulmonary cases, in addition to those in whom extrapulmonary disease is diagnosed.

The term "case detection", as used here, means that TB is diagnosed (correctly or incorrectly) in a patient, and is reported within the national surveillance system, and then to WHO. The case detection rate is calculated as the ratio of the number of notified smear-positive cases to the number of new smear-positive cases esti-

mated for that year. Detection is presented in two ways - as the case detection rate (countrywide) and as the DOTS case detection rate (by DOTS programmes):

tion rate to coverage estimates the case detection rate within DOTS areas (as distinct from the case detection rate nationwide), assuming that the TB incidence rate is homogene-

(5)case detection rate =

annual new smear-positive notifications (country)

estimated annual new smear-positive incidence (country)

DOTS case detection rate =

annual new smear-positive notifications (DOTS)

estimated annual new smear-positive incidence (country)

The case detection rate and the DOTS case detection rate are identical when a country reports only from DOTS areas. This generally happens when DOTS coverage is 100% but, in some countries where DOTS is implemented in only part of the country, no TB notifications are received from the non-DOTS areas. Furthermore, in some countries where DOTS coverage is 100%, patients may choose to seek treatment from non-DOTS providers, who in some cases notify TB cases to the national authorities.

Both of the above definitions of the case detection rate refer to smearpositive cases, although we also present the detection rate for all forms of TB. The detection rate of all forms is similarly presented in two ways: detection by DOTS programmes, and detection countrywide.

Although these indices are termed "rates", they are actually ratios. The number of cases notified is usually smaller than estimated incidence because of incomplete coverage by health services, under-diagnosis, or deficient recording and reporting. However, the calculated detection rate can exceed 100% if case-finding has been intense in an area that has a backlog of chronic cases, if there has been over-reporting (e.g. double-counting) or over-diagnosis, or if estimates of incidence are too low. If the expected number of cases per year is very low (especially if it is less than one), the case detection rate can vary markedly from year to year due to chance. Whenever this index comes close to or exceeds 100%, we attempt to investigate, as part of the joint planning and evaluation process with NTPs, which of these explanations is correct.

The ratio of the DOTS case detec-

ous across counties, districts, provinces, or other administrative units. Ideally, this ratio would have a value of 70% or more as DOTS coverage increases within any country. Where the value of this indicator is much lower, it is clear that the DOTS programme has been poorly implemented, at least in some parts of the designated DOTS area. Changes in the value of this ratio through time are a measure of changes in the quality of TB control, after the DOTS programme has been established.

#### Treatment success

Treatment success in DOTS programmes is the percentage of new smear-positive patients that are cured (negative on sputum smear examination), plus the percentage that complete a course of treatment, without bacteriological confirmation of cure (Table 2).11 Cure and completion are among the six mutually exclusive outcomes.12 The sum of cases assigned to these outcomes, plus any additional cases registered but not assigned to

<sup>&</sup>lt;sup>9</sup> World population prospects – the 2002 revision. New York, United Nations Population Division, 2003.

<sup>&</sup>lt;sup>10</sup> Tuberculosis handbook, Geneva, World Health Organization, 1998 (WHO/TB/ 98.253).

<sup>&</sup>lt;sup>11</sup>TB control programmes should ensure high treatment success before expanding case detection. The reason is that a proportion of patients given less than a fully-curative course of treatment remain chronically infectious, and continue to spread TB. Thus DOTS programmes must be shown to achieve high cure rates in pilot projects before attempting countrywide coverage.

<sup>12</sup> Treatment of tuberculosis: Guidelines for national programmes. Third edition. Geneva, World Health Organization, 2003 (WHO/CDS/TB/2003.313).

an outcome, adds up to 100% of cases registered (i.e. the treatment cohort).

We also compare the number of new smear-positive cases registered for treatment (for this report, in 2002) with the number of cases notified as smear-positive (also in 2002). All notified cases should be registered for treatment, and the numbers notified and registered should therefore be the same (discrepancies arise e.g. when subnational reports are not received at national level). If the number registered for treatment is not provided we take, as the denominator for treatment outcomes, the number notified for that cohort year. If the sum of the six outcome categories is greater than the number registered (or the number notified), we use this sum as the denominator.

Because the number of patients presenting for a second or subsequent course of treatment, and the outcome of further treatment, are indicative of NTP performance and levels of drug resistance, we have begun to compile these data in this report. We present the numbers of patients registered for re-treatment, and the outcomes of retreatment, for each of three registration types: re-treatment after relapse, failure or default. However, some countries do not yet compile data on cases registered for re-treatment after failure and default separately at national level. Furthermore, some countries do not have outcome data for each of these re-treament case types.

The assessment of outcomes for a given calendar year always lags notifications by one year to ensure that all patients registered during that calendar year have completed treatment. A DOTS country must report treatment outcomes, unless it is newly-classified as DOTS, in which case it would take an additional year to report outcomes from the first cohort of patients treated.

#### Overview of data in annexes

Annex 1 presents data on epidemiology and surveillance, and planning and financing for each of the 22 HBCs. Data on policy and strategy are collected for both DOTS and non-DOTS areas separately.

Annex 2 contains the estimates needed to evaluate MDG Target 8 and indicators 23 and 24. These data include case detection and treatment success rates to monitor DOTS implementation, and incidence, prevalence and death rates to monitor the impact of TB control.

These data are presented, for each of the six WHO regions, as follows:

- TB control policies for each country, stating which technical components of the DOTS strategy have been implemented;
- incidence, prevalence and death rates for 1990 (MDG reference year) and 2003;
- case notifications, detection rates, and DOTS coverage: nationally, and separately for DOTS and non-DOTS programmes. Notifications include new pulmonary cases (smearpositive, smear-negative and laboratory-confirmed), new extrapulmonary and re-treatment cases;
- treatment outcomes for 2002 cohorts: both the new smear-positive and the re-treatment cohorts from DOTS programmes (relapse, retreatment after default and re-treatment after failure are presented separately where possible, as well as all re-treatment cases combined), and the new smear-positive treatment outcomes (where available) from non-DOTS programmes;
- new smear-positive notification rates by age and sex for the whole country;
- new smear-positive notifications (numbers) by age and sex, from DOTS and non-DOTS programmes;
- notification rates and numbers since 1980, for all forms of TB;
- notification rates and numbers since 1995, for new smear-positive cases:
- country notes: remarks that may help to explain data reported by selected countries (e.g. additional breakdown of cases of interest, late-reported data, reasons for incomplete data, discrepancies in estimated population sizes).

The data in Annex 2 are available as Excel spreadsheets from www.who.int/tb.

## **Planning and DOTS** implementation

The information on strategic planning analysed and presented in this report reflects activities from July 2003 to June 2004. Country plans and activities are monitored through several mechanisms, including direct discussion with NTP managers, analysis of a questionnaire on planning and implementation sent by WHO to all HBCs during 2004 (available from www.who.int/tb), collaboration with international technical agencies, monitoring missions, comprehensive programme reviews, GFATM applications, regional NTP managers' meetings, and the annual meeting of the DOTS Expansion Working Group (DEWG) of the Stop TB Partnership. In writing this report, WHO staff worked with NTP managers of the 22 HBCs to:

- assess national TB control activities planned and carried out during 2004, focusing on activities to improve political commitment, expand access to DOTS, strengthen diagnosis, improve treatment outcomes, ensure adequate staffing, and improve programme monitoring and supervision;
- update the country profiles to summarize progress made by the end of 2004 in implementing, or scaling up, national plans for DOTS expansion;
- analyse constraints to reaching the targets for detection and treatment success;
- review and revise the list of partners operating in, or on behalf of, each country;
- assess levels of drug resistance and activities planned to address MDR-TB, including mechanisms of drug-resistance surveillance, MDR-TB diagnosis and treatment policies, and the availability of second-line drugs;
- determine the status of collaborative TB/HIV activities;

determine the status of additional strategies to expand DOTS, and to involve community and health-care providers not currently participating in the provision of DOTS.

#### Planning activities carried out in 2003

In preparation for the 5th DEWG meeting (Paris, France, 27-28 October 2004), NTP managers for the 22 HBCs were asked to summarize what activities had been planned for implementation during 2003, which of those activities were implemented, which were not and why, and what corrective actions were taken so that these activities could be implemented in 2004. The information from these DEWG summary tables, supplemented with additional information provided by NTP managers and by WHO staff, is incorporated into the country profiles (Annex 1).

### Update of country profiles

Country profiles (Annex 1) were updated by incorporating information from the following sources: summary tables prepared for the 5th DEWG; country posters presented by the 22 HBCs at the DEWG meeting; questionnaires submitted by the 22 HBCs; and consultations with, and reviews of, the country profiles by NTP staff and collaborating technical agencies.

#### Constraints and remedial actions

Following the previous analysis of constraints to DOTS expansion and remedial actions proposed,13 this year's report provides an update. Constraints and remedial actions were assessed with information provided at the DEWG meeting, and through personal communications with NTP managers and staff. Special attention was devoted to constraints related to laboratory services and human resources.

## Partnerships and coordination

The list of donors and collaborating organizations was updated in consultation with NTP managers, WHO regional and country offices and partners. Major technical agencies, along with financial partners, are listed in each country profile. The coordination of these numerous agencies is vital for the efficient use of limited

resources within countries, and is facilitated through a formal coordination mechanism, such as the national interagency coordinating committee (NICC).

#### Management of drug resistance

Data on the prevalence of drug resistance are collected through the WHO/ IUATLD Global Project on Anti-tuberculosis Drug Resistance Surveillance (DRS), which began in 1994, and which published its third report in 2004.8 Profiles of the 22 HBCs contain estimates of the national prevalence of MDR-TB among previously untreated TB patients based on survey data for those countries participating in the WHO/IUATLD project. For those countries that have not carried out surveys, figures given in the country profiles are estimates.

WHO develops global policy on the management of MDR-TB and facilitates access to second-line drugs through the Green Light Committee (GLC).14 As part of this process, and under the continuous monitoring of the GLC, several DOTS-Plus pilot projects are evaluating the feasibility and costeffectiveness of using second-line drugs for managing MDR-TB in countries with limited resources. Projects approved by the GLC have access to quality-assured, second-line drugs at reduced prices and benefit from technical support and external monitoring. This report summarizes the number and status of GLC-approved DOTS-Plus projects that had been established by 2004.

#### Collaborative TB/HIV activities

WHO has published an interim policy on collaborative TB/HIV activities15 that outlines the methods and benefits of collaboration between HIV and TB programmes. Three main areas of collaboration are recommended. First, organizational structures should be set up to plan and manage collaborative TB/HIV activities. Second, people infected with HIV should be screened for TB, treated if they have active disease, and offered isoniazid preventive therapy as needed. Third, TB patients should be offered voluntary counselling and testing for HIV infection (VCT); if positive, they should be offered cotrimoxazole preventive therapy and, wherever possible, ART. WHO has also developed a guide for monitoring and evaluating collaborative TB/HIV activities that defines indicators for each of the key activities recommended in the interim policy.<sup>16</sup>

To investigate progress in implementing the recommended collaborative TB/HIV activities, countries were asked, via the standard WHO data collection form, to report on the extent to which TB patients were tested for HIV, assessed for ART and provided with ART during 2003. A supplementary questionnaire (available at www.who.int/tb) was sent to the 41 countries that have the highest incidence rates of TB with HIV coinfection. This questionnaire asked specifically about policy developments between 2002 and 2003. The data obtained from both forms were reviewed at WHO regional offices and headquarters, and any inconsistencies or missing data were discussed with the national correspondent before being included in the analysis.

## Additional strategies for **DOTS** expansion

This report covers three areas:

- PPM initiatives that aim to bring a greater diversity of health-care providers into DOTS programmes, promoting the essential package of patient care and improving reporting and monitoring procedures;
- community participation that improves access to care and fosters a patient-centered approach to the management of TB. While the type and scope of community involve-

<sup>&</sup>lt;sup>13</sup> Global tuberculosis control: surveillance, planning, financing. WHO report 2004. Geneva, World Health Organization (WHO/HTM/TB/2004.331).

<sup>&</sup>lt;sup>14</sup> Gupta R et al. Increasing transparency in partnerships for health - introducing the Green Light Committee. Tropical Medicine and International Health, 2002,

<sup>&</sup>lt;sup>15</sup> Interim policy on collaborative TB/HIV activities. Geneva, World Health Organization, 2004 (WHO/HTM/TB/2004.330; WHO/HTM/HIV/2004.1).

<sup>&</sup>lt;sup>16</sup> A guide to monitoring and evaluation for collaborative TB/HIV activities. Geneva. World Health Organization, 2004 (WHO/ HTM/TB/2004.342; WHO/HIV/2004.09).

ment depends upon location and context, many HBCs regard civil society as an essential partner in providing support to patients and their families:

the feasibility of implementing the Practical Approach to Lung Health (PAL), which several countries are examining, and assessing its potential impact on TB case detection and on the rationalization of drug prescriptions.

In addition to the findings presented in this report, further details of PPM, TB/HIV, PAL and other projects can be found at www.who.int/tb.

## **Financing DOTS expansion**

The financial analysis in this series of annual reports on global TB control has evolved and improved since being introduced in 2002.17 The main developments in this year's report are: (a) to place greater emphasis on the presentation and analysis of trends, with each HBC profile including budget and cost data for four years; and (b) to provide a more complete analysis of data for countries other than the HBCs. The report has eight objectives:

- for each HBC, and for all HBCs combined, to present trends in total NTP budgets and expenditures for the period 2002-2005, with breakdowns by funding source and line
- for each HBC and for all HBCs combined, to present trends in total TB control costs<sup>18</sup> for the period 2002-2005, with breakdowns by funding source and line item:
- for each HBC and for all HBCs combined, to assess trends in NTP budgets and total TB control costs, giving particular attention to where progress has been made and where major funding gaps persist;
- for each HBC, to estimate and compare per patient costs, budgets and available funding for the period 2002-2005 and per patient expenditures for 2002 and 2003;
- for HBCs, to assess the relationship between gross national income (GNI) per capita and (a) per

- patient costs and (b) the fraction of funds contributed by the government;
- for HBCs, to assess whether projected budgets and available funding will be sufficient to achieve the global targets for case detection and treatment success:
- to assess the contribution of the GFATM to funding for TB control;
- for countries other than the HBCs, to quantify NTP budgets and funding gaps in 2004 and 2005.

#### Data collection

We collected data from five main sources: NTPs, the WHO-CHOICE web site,19 costing guidelines developed for the "Disease Control Priorities in Developing Countries" project (DCPP),20 GFATM proposals and databases, and previous WHO reports in this series. In 2004, data were collected directly from countries by means of a two-page questionnaire included in the standard WHO data collection form. NTP managers were asked to complete three tables. The first two tables required a summary of the NTP budget for fiscal years 2004 and 2005 in US\$, broken down by line item and funding source (including a column for funding gaps). The third table requested NTP expenditure data for 2003, broken down by line item and source of funding. The form also requested information about dedicated TB control infrastructure and the way in which general health infrastructure is used for TB control for example, the number of dedicated TB beds that exist, the number of outpatient visits that patients need to make to a health facility during treatment and the average number of days for which patients are hospitalized. We also asked for an estimate of the number of patients that would be treated in 2004 and 2005. We used the WHO-CHOICE web site to identify the average costs, in international dollars (I\$), of a hospital bed-day and an outpatient clinic visit in every country. The costing guidelines for the DCPP and the WHO-CHOICE web site were used to identify the purchasing power parity (PPP) exchange rates required for conversion of I\$ costs to costs in US\$ (for consistency with budget and expenditure data reported on the data collection form).

### Data entry and analysis

High-burden countries. Data entry and analysis focused on the 22 HBCs. We created a standardized spreadsheet, with one worksheet for each country. Additional worksheets were included for summary analyses and for the data required as inputs to the analyses in each country worksheet (e.g. notification data, unit costs for bed-days and outpatient clinic visits, and the typical number of outpatient clinic visits and days in hospital for different types of patient during treatment). For each country worksheet, seven tables were created. These were:

- NTP budget by source of funding for each year 2002-2005, with the funding sources defined according to the 2004 data collection form i.e. government (excluding loans), loans, grants (excluding GFATM), GFATM and budget gap;
- NTP budget by line item for each year 2002-2005, with the line items defined according to the 2004 data collection form i.e. firstline drugs, second-line drugs, dedicated NTP staff, initiatives to increase case detection and cure rates, collaborative TB/HIV activities, buildings/equipment and other;
- NTP expenditures by source of funding for 2002 and 2003, with funding sources as defined for NTP budgets;
- NTP expenditures by line item for 2002 and 2003, with line items defined as for NTP budgets;

<sup>&</sup>lt;sup>17</sup> Global tuberculosis control: surveillance, planning, financing. WHO report 2002. Geneva, World Health Organization, 2002 (WHO/CDS/TB/2002.295).

<sup>&</sup>lt;sup>18</sup> i.e. including costs not reflected in NTP budget data.

<sup>19</sup> www3.who.int/whosis/cea/prices/unit.

<sup>&</sup>lt;sup>20</sup> DCPP guidelines for authors, pp. 74–77, (available at www.fic.nih.gov/dcpp/ authorguide.pdf, accessed 11 January 2005).

- total TB control costs by funding source for each year 2002-2005, with funding sources defined as for NTP budgets;
- total TB control costs by line item for each year 2002-2005, with the line items defined as NTP budget items, hospitalization and clinic vis-
- per patient costs, NTP budget, available funding, expenditures and budget for first-line drugs.

Budget data for 2004 and 2005 were taken from the 2004 data collection form. Budget data for 2002 and 2003 were taken from the 2002 and 2004 annual reports, respectively. Expenditure data for 2002 and 2003 were based on the 2003 and 2004 data collection forms, respectively. Total TB control costs were estimated by adding costs for hospitalization and outpatient clinic visits to either NTP expenditures (for 2002 and 2003) or NTP budgets (for 2004 and 2005).21 Expenditures were used in preference to budgets for 2002 and 2003 because they reflect actual costs, whereas budgets can be higher than actual expenditures (for example, when large budgetary funding gaps exist or the NTP does not spend all the available funding). When expenditures are known for 2004 and 2005, they will be used instead of budget data to calculate, retrospectively, the total cost of TB control in these years. For some HBCs, expenditures were not available for 2002 and 2003. When this was the case, we estimated expenditures based on available funding, which was calculated as the total budget minus the funding gap.

The cost of outpatient clinic visits was estimated in three steps. First, we converted I\$ prices for clinic visits reported on the WHO-CHOICE web site into US\$ prices using the DCPP exchange rates. Second, we multiplied the average number of visits required per patient (estimated on the WHO data collection form) by the average cost (in US\$) per clinic visit, to give the cost per patient treated. Third, we multiplied the cost per patient treated by the number of patients notified (for 2002 or 2003) or the number of pa-

tients that the NTP projects will be treated (for 2004 and 2005). The cost of hospitalization was generally calculated in the same way, replacing the unit cost of a clinic visit with the unit cost of a bed-day. The procedure differed for eight countries that have dedicated TB beds, and where the total cost of these beds is higher than implied by multiplying bed-days per patient by the number of patients treated (this applied to Brazil, Cambodia, India, Nigeria, the Russian Federation, the United Republic of Tanzania, Viet Nam and Zimbabwe). We assumed that all clinic visits and hospitalization are funded by the government.

Per patient costs, budgets, available funding and expenditures were calculated by dividing the relevant total by the number of cases notified (for 2002 and 2003) and the number of patients that the NTP projects will be treated (for 2004 and 2005). Since the total costs of TB control for 2002 and 2003 were based on expenditure data, it is possible for the total TB control cost per patient treated to be less than the NTP budget per patient treated when the funding gap is large or there is an important budgetary under-spend. In addition, for 2002 and 2003, the expenditure per patient was sometimes higher than the available funding per patient. This can occur when some of the NTP budget funding gap is closed following the reporting of budget data to WHO.

All data are reported in nominal prices (i.e. they have not been adjusted for inflation) rather than constant prices (i.e. all data are adjusted to a common year of prices) for two reasons. First, this avoids adjustment of values reported in the 2002–2004 reports in this series, which makes it easier for country staff to review the data for previous years. Second, the adjustment will make only a limited difference to the numbers reported (about 5% to 2002 values and less for other years). However, as data are collected for an increasing number of years, presentation of data in constant prices will be necessary.

Following data entry, text on data sources and assumptions were added. Where there were questions

about the data, these were discussed with NTP staff and the appropriate WHO regional and country office. These discussions were used to produce a final set of charts. Four of these charts appear in the profiles for each country at Annex 1: NTP budget by funding source, NTP budget by line item, total TB control costs by line item, and per patient costs, budgets, available funding, expenditures and budget for first-line drugs. These charts were selected because they illustrate the most important trends in financing, while other data are referred to in the text. A full set of charts and data is available upon request. In some instances, the review process led to revisions to data included in previous annual reports. For this reason, figures sometimes differ from those reported in the 2002, 2003 and 2004 reports.

Finally, we compared the total costs of TB control with total government health expenditures to estimate the percentage of total government health expenditures used for TB control. Total government health expenditures were estimated by multiplying the government health expenditure per capita in US\$ (as estimated in the WHO national health accounts database)22 by population size. We also explored the association between GNI per capita in 2003 and (a) government contributions to total NTP budgets and TB control costs, and (b) the cost per patient treated. Data on GNI per capita were taken from World development indicators 2004.23

Other countries. The data provided by countries other than the HBCs were less complete, and as a consequence our analyses to date are more superficial. We used the data provided on the 2004 data collection form to assess NTP budgets by region, and com-

<sup>&</sup>lt;sup>21</sup> The exception was South Africa, because no data on hospitalization and clinic visits, or on NTP budgets, were provided in the data collection form. Costs were therefore estimated based on recent costing studies, as described in previous WHO reports in this series.

<sup>&</sup>lt;sup>22</sup> www.who.int/nha/country/en/.

<sup>&</sup>lt;sup>23</sup> www.worldbank.org/data/wdi2004/.

pared these with the budgets reported by the HBCs. Only countries that submitted complete data of sufficient quality (e.g. subtotals and totals were consistent by both line item and funding source) were used.

### GFATM contribution to TB control

We assessed GFATM funding for both HBCs and other countries, as announced after the first four rounds of funding. We assessed total approved funding at the end of 2004, how the amounts in signed grant agreements compared with those in the original proposals, disbursements to the end of 2004, the time taken between approval of a proposal and the signature of grant agreements, and the time  $% \left( 1\right) =\left( 1\right) \left( 1\right$ taken between the signing of the grant agreement and the first disbursement of funds.

## Results

## **Progress towards the Millennium Development Goals**

### Countries reporting to WHO

By the end of 2004, 199 (94%) of 211 countries and territories reported case notifications for 2003 and/or treatment outcomes for patients registered in 2002. These countries include 99% of the world's population. WHO received reports from all 22 HBCs.

#### Case notifications and incidence

The 199 countries reporting to WHO in 2003 notified 4.4 million new and relapse cases, of which 1.9 million (44%) were new sputum smear-

positive (Table 3; Figure 1). Among these notifications, 3.7 million were from DOTS areas, including 1.8 million smear-positives. A total of 17.1 million cases, and 8.6 million smearpositives, were notified by DOTS programmes between 1995 and 2003. Based on surveillance and survey data, we estimate that there were 8.8 million new cases of TB in 2003 (140 per 100 000), including 3.9 million (62 per 100 000) smear-positive cases (Table 4; Figure 2).

The African Region (24%), South-East Asia Region (35%), and Western Pacific Region (22%) together accounted for 82% of all notified cases and similar proportions of new smearpositive cases. Because DOTS emphasizes diagnosis by sputum smear microscopy, 47% of all new and relapse cases were smear-positive (45-60% expected) in DOTS areas, compared with 29% elsewhere. Similarly, 58% of new pulmonary cases were smear-positive under DOTS (55–70% expected), compared with 35% elsewhere (Table 3).

The ranking of countries by number of incident TB cases has drawn attention to the 22 HBCs, but the magnitude of the TB burden in individual

TABLE 3 Case notifications, 2003

		NEW AND PSE CASES								RE-TREATMENT CASES EXCLUDING RELAPSE		ΓHER <sup>a</sup>		W PULMONARY MEAR POSITIVE <sup>b</sup>
	DOTS	NON-DOTS	DOTS	NON-DOTS	DOTS	NON-DOTS	DOTS	NON-DOTS	DOTS	NON-DOTS	DOTS	NON-DOTS	DOTS	NON-DOTS
1 India	836 768	236 297	372 088	61 183	305 921	153 503	112 064	20 189	102 542	13 247	_	-	55	29
2 China	553 677	62 191	257 287	10 127	206 493	42 312	27 804	2 964	64 887	2 822	-	-	55	19
3 Indonesia	178 260	-	92 566	-	77 561	-	4 047	-	-	-	-	-	54	-
4 Nigeria	44 184	-	28 173	-	13 276	-	1 525	-	2 151	-	261	-	68	-
5 Bangladesh	88 156	-	53 618	-	24 913	-	7 120	-	-	-	-	-	68	-
6 Pakistan	73 100	-	20 962	-	34 447	-	12 874	-	3 184	-	-	-	38	-
7 Ethiopia	117 600	-	39 698	-	35 141	-	40 883	-	676	-	-	-	53	-
8 South Africa	227 278	42	116 331	33	56 535	5	37 682	4	28 094	8	-	-	67	87
9 Philippines	134 375	-	72 670	-	55 942	-	1 693	-	-	-	-	-	57	-
10 Kenya	91 522	-	38 158	-	37 135	-	13 403	-	1 127	-	2 661	-	51	-
11 DR Congo	84 687	-	53 578	-	9 352	-	18 357	-	1 641	-	387	-	85	-
12 Russian Federation	21 064	102 977	6 322	22 546	12 780	72 252	1 016	3 648	-	22 512	851	4 840	33	24
13 Viet Nam	92 741	-	55 937	_	16 791	-	14 564	-	680	_	-	_	77	-
14 UR Tanzania	61 579	-	24 899	-	21 911	-	12 959	-	378	-	2 708	-	53	-
15 Brazil	16 560	63 554	9 061	30 877	4 795	18 727	1 503	9 081	799	3 663	1 256	2 690	65	62
16 Uganda	41 805	-	20 320	-	16 612	-	3 249	-	-	-	1 096	-	55	-
17 Thailand	54 504	-	28 459	-	17 596	-	6 756	-	_	-	-	-	62	-
18 Mozambique	28 602	-	16 138	-	7 847	-	3 441	-	505	-	-	-	67	-
19 Zimbabwe	53 183	-	14 488	-	28 246	-	8 9 1 6	-	_	-	3 934	-	34	-
20 Myanmar	75 744	-	27 448	-	26 006	-	17 796	-	2 451	-	-	-	51	-
21 Afghanistan	13 808	-	6 510	-	3 440	-	3 254	-	141	-	-	-	65	-
22 Cambodia	28 216	-	18 923	-	4 307	-	4 232	-	79	-	91	-	81	-
High-burden countries	2 917 413	465 061	1 373 634	124 766	1 017 047	286 799	355 138	35 886	209 335	42 252	13 245	7 530	57	30
AFR	1 061 882	10 789	503 217	6 947	319 715	2 513	193 812	1 013	39 548	403	19 902	-	61	73
AMR	142 409	85 142	82 479	43 324	31 761	24 210	19 936	11 835	4 208	4 259	5 909	3 154	72	64
EMR	206 160	3 781	80 822	191	63 703	2 099	51 417	1 488	4 015	-	178	-	56	8
EUR	142 760	195 883	44 673	50 839	64 716	96 592	16 547	12 085	7 334	25 316	22 292	54 198	41	34
SEAR	1 314 983	240 402	610 079	62 799	481 487	155 219	160 093	20 772	107 746	13 408	9 845	583	56	29
WPR	879 827	108 100	431 396	23 336	313 113	63 566	59 422	11 084	66 352	3 563	3 262	3 746	58	27
Global	3 748 021	644 097	1 752 666	187 436	1 274 495	344 199	501 227	58 277	229 203	46 949	61 388	61 681	58	35

Indicates not applicable or not available.

<sup>&</sup>lt;sup>a</sup> Cases not included elsewhere in table.

b Expected percentage of new pulmonary cases that are smear-positive is 65-80%.

FIGURE 1 **Tuberculosis notification rates, 2003** 

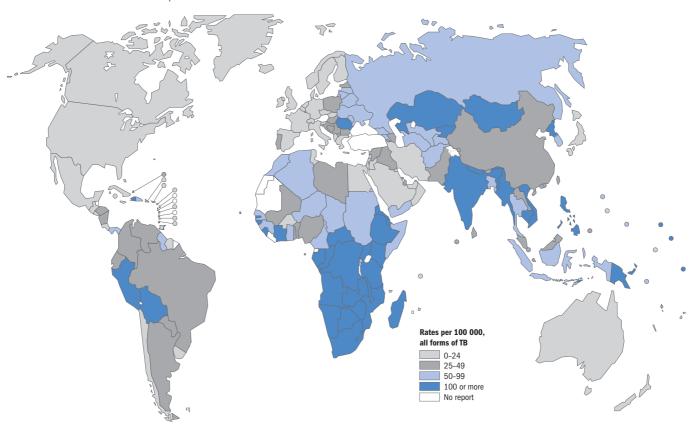


FIGURE 2 Estimated TB incidence rates, 2003

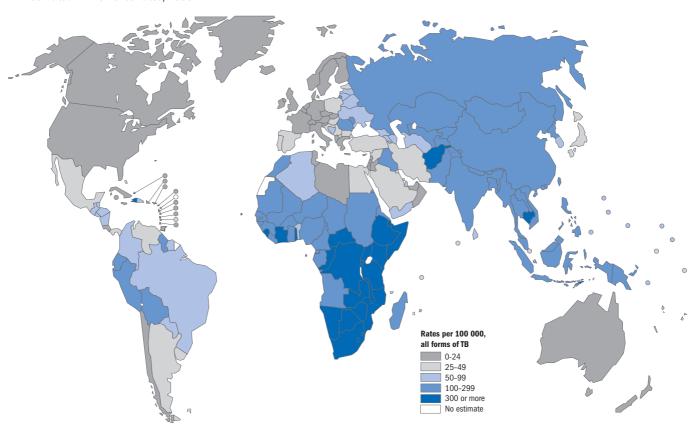
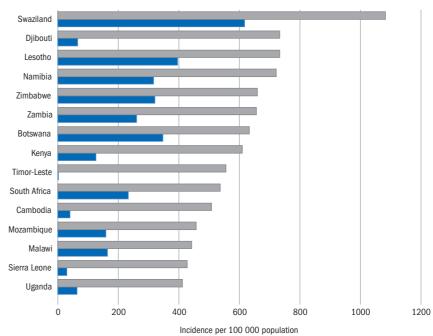


TABLE 4 Estimated TB burden, 2003

	POPULATION 1000s		INCIDENCE				VALENCE	MORTALITY		
		ALL	CASES	SMEAR-PO	SITIVE CASES	ALL FORMS OF TB, INCLUDING IN HIV-INFECTED PEOPL				
		NUMBER 1000s	RATE PER 100 000 POP.	NUMBER 1000s	RATE PER 100 000 POP.	NUMBER 1000s	RATE PER 100 000 POP.	NUMBER 1000s	RATE PER 100 000 POP.	
1 India	1 065 462	1 788	168	798	75	3 086	290	352	33	
2 China	1 304 196	1 334	102	600	46	3 203	246	236	18	
3 Indonesia	219 883	627	285	282	128	1 484	675	143	65	
4 Nigeria	124 009	363	293	156	126	677	546	105	85	
5 Bangladesh	146 736	361	246	162	111	719	490	84	57	
6 Pakistan	153 578	278	181	125	82	551	359	67	43	
7 Ethiopia	70 678	252	356	109	155	377	533	56	79	
8 South Africa	45 026	242	536	98	218	206	458	33	73	
9 Philippines	79 999	237	296	107	133	366	458	39	49	
10 Kenya	31 987	195	610	84	262	283	884	43	133	
11 DR Congo	52 771	195	369	85	160	298	564	43	81	
12 Russian Federation	143 246	161	112	72	50	229	160	29	20	
13 Viet Nam	81 377	145	178	65	80	195	240	19	23	
14 UR Tanzania	36 977	137	371	58	157	194	524	32	86	
15 Brazil	178 470	110	62	49	28	164	92	15	8	
16 Uganda	25 827	106	411	46	179	168	652	25	96	
17 Thailand	62 833	89	142	40	63	130	208	12	19	
18 Mozambique	18 863	86	457	36	190	120	636	24	129	
19 Zimbabwe	12 891	85	659	34	265	85	660	20	153	
20 Myanmar	49 485	85	171	38	76	92	187	12	25	
21 Afghanistan	23 897	80	333	36	150	160	671	22	93	
22 Cambodia	14 144	72	508	32	225	108	762	13	95	
High-burden countries	3 942 338	7 027	178	3 112	79	12 896	327	1 423	36	
AFR	687 405	2 372	345	1 013	147	3 487	507	538	78	
AMR	867 768	370	43	165	19	503	58	54	6	
EMR	518 063	634	122	285	55	1 120	216	144	28	
EUR	878 902	439	50	196	22	577	66	67	8	
SEAR	1 614 648	3 062	190	1 370	85	5 662	351	617	38	
WPR	1 732 104	1 933	112	868	50	4 081	236	327	19	
Global	6 298 890	8 810	140	3 897	62	15 430	245	1 747	28	

FIGURE 3 Fifteen countries with the highest estimated TB incidence rates per capita (all ages, all forms; grey bars) and corresponding incidence rates of HIV-infected TB (among adults aged 15-49 years; blue bars), 2003



countries is better expressed as the incidence rate per capita. Among the 15 countries with the highest estimated TB incidence rates per capita, 12 are in Africa (Figure 3).

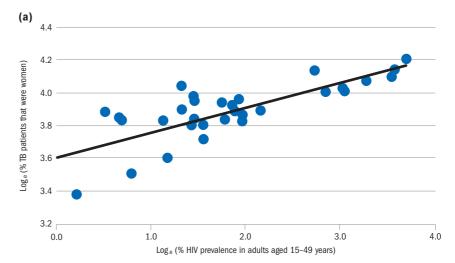
Case notifications from African countries show other patterns that are likely to be associated with HIV infection. First, women aged 15-24 years make up a higher proportion of TB cases in countries with higher rates of HIV infection (Figure 4a). This is consistent with the observation that HIV prevalence tends to be higher in women than men in this age range, and the difference between the sexes is bigger where HIV infection rates are higher. Second, the average age of smear-positive TB cases is typically lower where HIV infection rates are higher, especially for women (Figure 4b). This is another sign that younger rather than older women are more likely to be infected with HIV. Third, the proportion of smear-negative cases among all pulmonary TB cases tends to be higher in African countries with higher rates of HIV infection (data not shown). However, this last association is weak  $(R^2 = 0.16, P = 0.02)$  and could be confounded by the quality of diagnosis if, for example, smear microscopy has become less reliable where the number of HIV-infected TB patients has increased substantially.

Some patterns in the case notification data are striking, but not easy to explain. For example, the number of extrapulmonary TB cases as a proportion of the total reported is consistently different among WHO regions. Between 1995 and 2003, the proportion was lowest in the Western Pacific Region (mostly <5%) and highest in the Eastern Mediterranean Region (20-30%; see Figure 5). We do not know whether these are real epidemiological differences, or due to regional diagnostic biases. Surprisingly, the proportion of cases diagnosed as extrapulmonary disease has not increased in the African Region, despite the growing impact of HIV on the TB epidemic. This raises the question of whether NTPs in Africa are missing extrapulmonary cases.

Using the series of notifications of all TB cases from countries thought to have reliable data, and scaling by

#### FIGURE 4

(a) The proportion of notified TB patients aged 15-24 years that were women, in relation to HIV prevalence in adults aged 15-49 years ( $R^2 = 0.63$ , P < 0.0001). (b) Average age of women with smear-positive TB, in relation to HIV prevalence in adults aged 15-49 years  $(R^2 = 0.46, P < 0.0001)$ . Data are for countries in Africa.



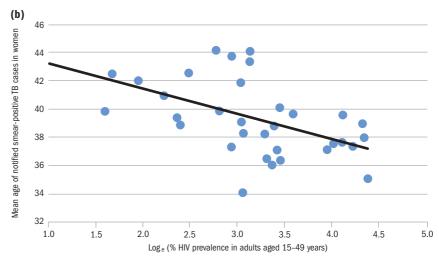


FIGURE 5 The number of extrapulmonary TB cases as a percentage of the total number of cases reported, for each of the six WHO regions

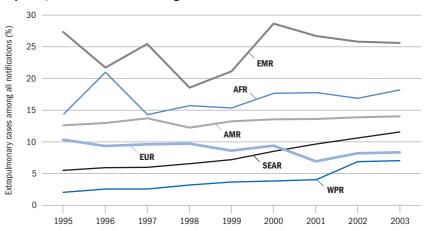
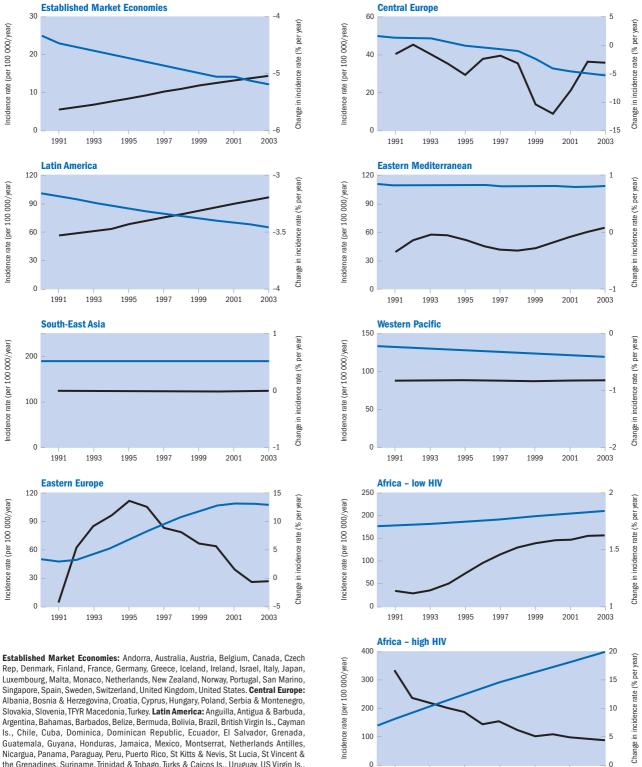


FIGURE 6 Trends in estimated TB incidence rates (all forms; blue lines), and the annual change in incidence rates (black lines), for nine groups of countries, 1990-2003



Luxembourg, Malta, Monaco, Netherlands, New Zealand, Norway, Portugal, San Marino, Guatemala, Guyana, Honduras, Jamaica, Mexico, Montserrat, Netherlands Antilles, the Grenadines, Suriname, Trinidad & Tobago, Turks & Caicos Is., Uruguay, US Virgin Is., Venezuela. **Eastern Mediterranean:** Afghanistan, Bahrain, Egypt, Iran, Iraq, Jordan, Kuwait, Lebanon, Libyan Arab Jamahiriya, Morocco, Oman, Pakistan, Qatar, Saudi Arabia, Syrian Arab Rep., Tunisia, United Arab Emirates, West Bank & Gaza Strip, Yemen. South-East Asia: Bangladesh, Bhutan, DPR Korea, India, Indonesia, Maldives, Myanmar, Nepal, Sri Lanka, Thailand, Timor-Leste. Western Pacific: Amerian Samoa, Brunei Darussalam, Cambodia, China, China Hong Kong SAR, China Macao SAR, Cook Is., Fiji, French Polynesia, Guam, Kiribati, Lao PDR, Malaysia, Marshall Is., Micronesia, Mongolia, Nauru, New Caledonia, Niue, N. Mariana Is., Palau, Papua New Guinea, Philippines, Rep. Korea, Samoa, Solomon Is., Tokelau, Tonga, Vanuatu, Viet Nam, Wallis & Futuan Is. **Eastern Europe:** Armenia, Azerbaijan, Belarus, Bulgaria, Estonia, Georgia, Kazakhstan, Kyrgystan, Latvia, Lithuania, Rep. Moldova, Romania, Russian Federation, Tajikistan, Turkmenistan,

Ukraine, Uzbekistan. Africa - Iow HIV: Algeria, Angola, Benin, Burkina Faso, Cape Verde, Chad, Comoros, Djibouti, Equatorial Guinea, Eritrea, Gambia, Ghana, Guinea, Guinea-Bissau, Liberia, Madagascar, Mali, Mauritania, Mauritius, Niger, Sao Tome & Principe, Senegal, Seychelles, Sierra Leone, Somalia, Sudan, Togo. Africa – high HIV: Botswana, Burundi, Cameroon, Central African Rep., Congo, Côte d'Ivoire, DR Congo, Ethiopia, Gabon, Kenya, Lesotho, Malawi, Mozambique, Namibia, Rwanda, South Africa, Swaziland, Uganda, UR Tanzania, Zambia, Zimbabwe.

1997

1999

2001

2003

1991

1993

1995

the estimated rates of case detection, we have estimated the trends in TB incidence rate (all forms) for nine epidemiologically distinct regions of the world (Figure 6). In six of these regions, the trend in the incidence rate has been downward.

Incidence rates have been increasing for most of the period since 1990 in African countries with low and high rates of HIV infection, and in eastern Europe, although the patterns of change in the three regions are quite different. In African countries with high HIV infection, incidence has been pushed upwards by the spread of HIV, but the rate of increase has fallen from a maximum exceeding 15% per year in the early 1990s (Figure 6). In African countries with lower rates of HIV infection, the rate of increase in TB has never been as high (2-3% per year), but neither are there signs that the increase is slowing. In eastern Europe, the rate of increase reached nearly 15% annually by 1995, but the increase now appears to have been halted, and incidence is once again in decline.

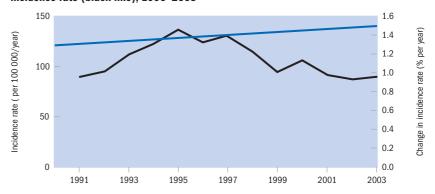
The global trend is obtained by summing the estimated numbers of TB cases across all nine regions (Figure 7). Worldwide, the incidence rate of TB was growing at a maximum of around 1.5% per year in 1995, but less than 1% per year by 2003.

#### TB and HIV

Some countries have carried out surveys of the prevalence of HIV in TB patients, either nationally or locally, and the results have been reported via the data collection form or the supplementary TB/HIV questionnaire. Although the accuracy of the data is not known because, for example, the design of the surveys has not been fully described, a growing number of countries are testing TB patients for HIV infection.

The prevalence of HIV infection in TB patients can be derived from the incidence rate ratio (IRR). IRR is estimated from the relationship between HIV prevalence in adult TB patients and HIV prevalence in the adult population, where both have been measured together (Figure 8). The IRR derived from the national surveys in this set of data is 8.3 (95% CI, 6.1-

FIGURE 7 Trends in the estimated global TB incidence rate (blue line), and the annual change in incidence rate (black line), 1990-2003



#### FIGURE 8

The prevalence of HIV in TB patients as measured in national surveys (blue dots) and subnational surveys (data reported to WHO; black dots), plotted against the prevalence of HIV in adults aged 15-49 years (data from UNAIDS). The incidence rate ratio is 8.3 (6.1–10.8; P = 0.0036) for the national survey data and 8.4 (7.9–10.0; P =0.0029) for the subnational surveys. The countries are: BFA Burkina Faso; BOT Botswana; BUU Burundi; CAE Cameroon; CAF Central African Republic; CAM Cambodia; CNG Congo; COD DR Congo; DJI Djibouti; ETH Ethiopia; GHA Ghana; HAI Haiti; IVC Côte d'Ivoire; KEN Kenya; LES Lesotho; MAL Malawi; MOZ Mozambique; NIE Nigeria; RWA Rwanda; SOA South Africa; TAN UR Tanzania; ZIM Zimbabwe.

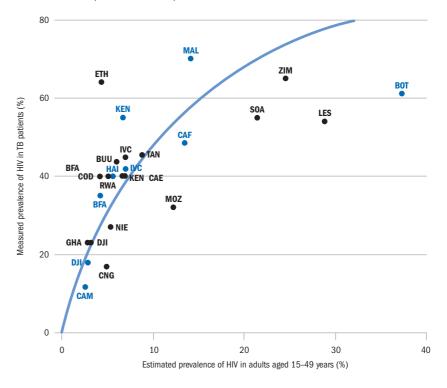


FIGURE 9
Estimated HIV prevalence in TB cases, 2003

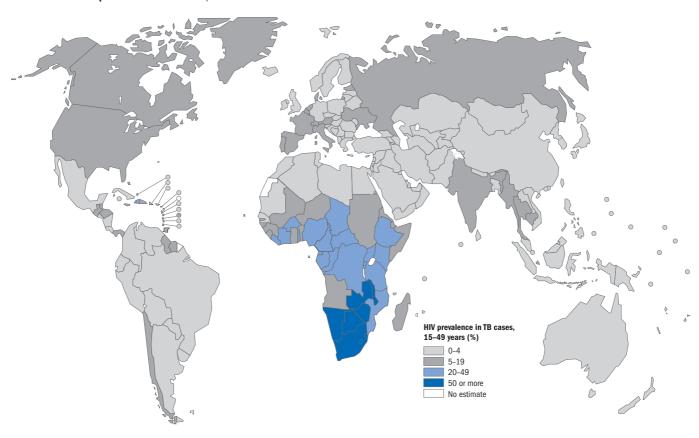


FIGURE 10

Number of countries implementing DOTS (out of a total of 211 countries), 1991–2003

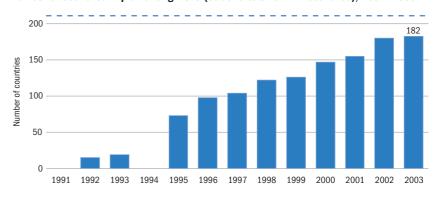
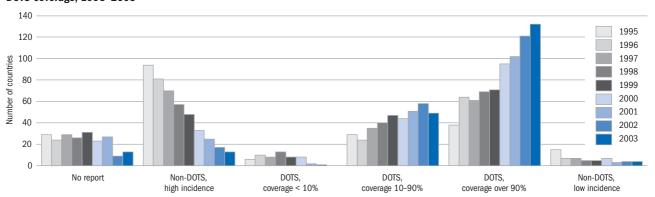


FIGURE 11 **DOTS coverage, 1995–2003** 



10.8), which is higher than, but not significantly different from, the previously published estimate of 6.0 (3.1- $8.0).^{24}$ 

Further HIV surveys among TB patients will give, once the data have been validated, better direct measures of the TB/HIV association for the countries surveyed and, through the IRR, better indirect estimates for countries that do not yet test TB patients for HIV infection, thereby improving the distribution map in Figure 9.

### DOTS coverage

The total number of countries implementing DOTS increased by two during 2003, bringing the total to 182 out of 211 (Figure 10). All 22 HBCs have had DOTS programmes since 2000; many of these programmes have been established for much longer.

DOTS coverage within countries has steadily increased since 1995 (Figure 11; Table 5). By the end of 2003, 77% of the world's population lived in counties, districts, oblasts and provinces of countries that had adopted DOTS. Coverage was reported to be more than 70% in all regions except Europe (Figure 12).

### Case detection

The 4.4 million cases of TB (new and relapse) notified in 2003 represent half (50%) of the 8.8 million estimated new cases; the 1.9 million new smearpositive cases notified also account for half (50%) of the 3.9 million estimated (Table 3, Table 4). In parallel with trends in case notifications, the detection rate of all TB cases, from DOTS and non-DOTS programmes, has remained stable since 1995, while the detection rate of smear-positive cases has slowly increased (Figure 13). Therefore, the proportion of all cases diagnosed as smear-positive has been rising.

DOTS programmes detected an estimated 43% of all new and relapse cases, and 45% of new smear-positive cases, in 2003. The detection rate achieved by DOTS programmes has

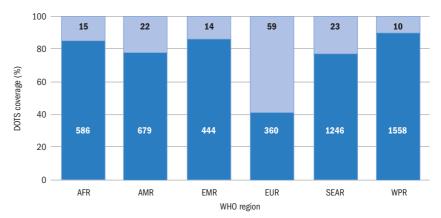
TABLE 5 Progress in DOTS implementation, 1995-2003

			PERCENT	AGE OF PO	PULATION C	OVERED B	Y DOTS		
	1995	1996	1997	1998	1999	2000	2001	2002	2003
1 India	1.5	2	2.3	9	13.5	30	45	51.6	67.2
2 China	49	60.4	64.2	63.9	64	68	68	77.6	91
3 Indonesia	6	13.7	28.3	80	90	98	98	98	98
4 Nigeria	47	30	40	45	45	47	55	55	60
5 Bangladesh	40.5	65	80	90	90	92	95	95	99
6 Pakistan	2	8	-	8	8	9	24	45	63
7 Ethiopia	39	39	48	64.4	63	85	70	95	95
8 South Africa	_	0	13	22	66	77	77	98	99.5
9 Philippines	4.3	2	15	16.9	43	89.6	95	98	100
10 Kenya	15	100	100	100	100	100	100	100	100
11 DR Congo	47	51.4	60	60	62	70	70	70	75
12 Russian Federation	-	2.3	2.3	5	5	12	16	25	25
13 Viet Nam	50	95	93	96	98.5	99.8	99.8	99.9	100
14 UR Tanzania	98	100	100	100	100	100	100	100	100
15 Brazil	_	0	0	3	7	7	32	25	33.6
16 Uganda	_	0	100	100	100	100	100	100	100
17 Thailand	_	1.1	4	32	59	70	82	100	100
18 Mozambique	97	100	84	95	-	100	100	100	100
19 Zimbabwe	_	0	0	100	11.6	100	100	100	100
20 Myanmar	_	59	60	60.3	64	77	84	88.3	95
21 Afghanistan	_	_	12	11	13.5	15	12	38	53
22 Cambodia	60	80	88	100	100	99	100	100	100
High-burden countries	24	32	36	43	46	55	61	68	79
AFR	43	47	56	62	56	70	70	81	85
AMR	12	48	50	55	65	68	73	73	78
EMR	23	11	18	33	51	66	72	77	86
EUR	5.4	8.2	17	22	23	26	32	40	41
SEAR	6.7	12	16	30	36	50	61	66	77
WPR	43	55	57	58	57	67	68	77	90
Global	22	32	37	44	47	57	62	69	77

O Indicates that a report was received, but the country had not implemented DOTS.

FIGURE 12

DOTS population coverage by WHO region, 2003. The shaded portion of each bar shows the DOTS coverage as a percentage of the population. The numbers in each bar show the population (in millions) within (dark portion) or outside (light portion) DOTS areas.

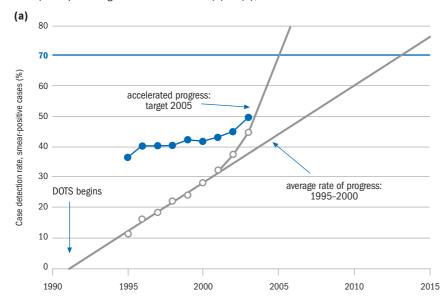


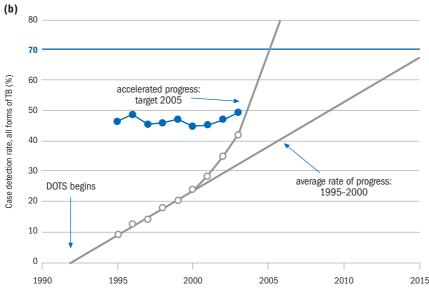
 $<sup>^{\</sup>rm 24}\,\mbox{Corbett EL}$  et al. The growing burden of tuberculosis: global trends and interactions with the HIV epidemic. Archives of Internal Medicine, 2003, 163:1009-1021.

Indicates that no report was received.

### FIGURE 13

Progress towards the 70% case detection target. (a) Open circles mark the number of smear-positive cases notified under DOTS 1995-2003, expressed as a percentage of estimated new cases in each year. The solid line through these points indicates the average annual increment from 1995 to 2000 of about 134 000 new cases, compared with the increment from 2002 to 2003 of 324 000 cases; the steeper line represents a higher annual increment of approximately 488 000 cases per year needed to reach the 70% target by 2005. Closed circles show the total number of smear-positive cases notified (DOTS and non-DOTS) as a percentage of estimated cases. (b) As (a), but for all forms of TB.





been rising more quickly than the overall case detection rate, and has accelerated since 2000. The 7.5% increase in DOTS case detection between 2002 and 2003, an additional 324 000 smear-positive cases, is the largest annual increase so far reported. If this rate of increase is maintained, the estimated detection rate will be 60% in 2005. To reach the 70% target by 2005, DOTS programmes must find and treat an extra 488 000 cases in each of the two remaining years.

Because case detection under DOTS has increased faster than the overall rate of case detection, the proportion of notified smear-positive cases that were notified by DOTS programmes has also increased, reaching 90% in 2003. DOTS programmes have continued to recruit largely from the pool of patients that would have been detected anyway in the public

Although more cases are recruited to DOTS programmes each year, the case detection rate within DOTS areas (measured by the ratio of case detection to population coverage) has changed little, averaging 52% worldwide between 1996 and 2003 (Figure 14). There are signs of a slow rise in the HBCs, from 35% in 1995 to 56% in 2003, due mostly to improvements in Bangladesh, India, Indonesia, Myanmar and the Philippines.

Smear-positive case detection rates by DOTS programmes in 2003 were lowest in the European Region (23%) and highest in the African Region, Region of the Americas and Western Pacific Region (all 50%; see Figure 15, Table 6). The rate of improvement in case detection by DOTS pro-

FIGURE 14 Smear-positive case detection rate within **DOTS** areas for high-burden countries (blue) and the world (grey), 1995-2003. DOTS case detection rate divided by DOTS coverage, expressed as percentage.

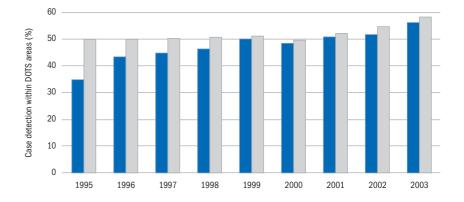


FIGURE 15 Smear-positive case detection rate by DOTS programmes, by WHO region, 1995–2003. Heavy line shows global DOTS case detection rate.

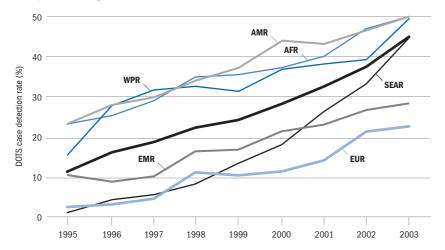


TABLE 6 Case detection rate of new smear-positive cases (%), 1995-2003

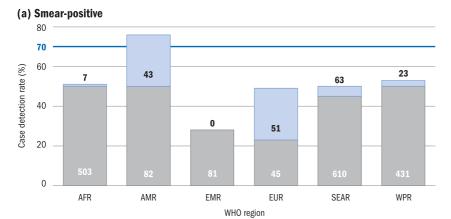
_				DOTS	PROGRAM	IMES							WH	OLE COUN	TRY			
	1995	1996	1997	1998	1999	2000	2001	2002	2003	1995	1996	1997	1998	1999	2000	2001	2002	200
1 India	0.3	0.9	1.1	1.7	7.1	12	24	31	47	38	41	38	38	46	46	50	50	54
2 China	15	28	32	32	29	31	31	30	43	22	34	39	33	33	34	34	32	45
3 Indonesia	1.4	4.6	7.5	12	18	19	20	27	33	13	*	*	*	*	19	*	*	*
4 Nigeria	12	12	12	12	14	14	14	13	18	*	*	*	*	*	*	17	15	*
5 Bangladesh	6.7	14	18	23	23	23	25	29	33	15	21	23	26	25	25	26	29	*
6 Pakistan	1.0	1.8	-	3.8	2.0	2.8	5.2	13	17	2.5	*	_	14	5.5	*	9.2	13	*
7 Ethiopia	16	21	23	25	26	35	35	36	36	*	*	*	*	27	*	*	*	*
8 South Africa	_	_	6.1	22	70	75	81	105	118	41	69	80	91	93	91	95	106	118
9 Philippines	0.4	0.5	3.2	10	20	48	57	62	68	96	87	80	68	71	65	*	*	*
10 Kenya	53	54	50	54	52	44	47	46	46	*	*	*	*	*	48	*	*	k
11 DR Congo	43	49	46	57	56	53	58	57	63	47	*	*	57	*	*	*	*	*
12 Russian Federation	_	0.5	1.0	1.0	1.7	4.6	5.2	6.9	8.8	72	69	64	60	29	35	34	37	40
13 Viet Nam	30	60	79	83	84	83	84	88	86	60	77	*	86	84	*	*	88	k
14 UR Tanzania	55	54	51	52	50	47	46	43	43	*	*	*	*	*	*	*	*	*
15 Brazil	_	_	_	4.1	3.9	7.5	8.0	10	18	79	79	78	80	78	79	75	82	81
16 Uganda	_	_	58	58	58	50	46	46	44	49	54	*	*	*	*	*	*	*
17 Thailand	_	0.3	5.0	21	39	46	73	65	72	55	46	35	*	*	*	*	*	*
18 Mozambique	56	50	48	48	_	45	44	45	45	*	*	*	*	47	*	*	*	k
19 Zimbabwe	_	_	_	51	49	46	46	47	42	50	54	57	*	*	*	*	*	*
20 Myanmar	-	26	26	29	32	48	56	65	73	26	28	28	*	*	*	58	*	k
21 Afghanistan	_	_	2.0	5.9	5.3	9.0	14	19	18	_	_	*	*	*	*	*	*	*
22 Cambodia	40	34	44	47	53	49	47	55	60	*	42	*	*	*	*	*	*	*
High-burden countries	8.5	14	17	20	23	27	31	35	44	33	37	37	37	39	39	41	43	48
AFR	24	26	29	35	36	37	40	47	50	38	43	41	45	45	43	44	47	50
AMR	23	28	30	34	37	44	43	46	50	72	72	77	77	76	74	75	76	76
EMR	11	9.1	10	17	17	22	23	27	28	20	25	24	30	27	23	26	27	28
EUR	2.6	3.4	4.5	11	11	12	14	22	23	63	62	57	57	45	47	43	42	49
SEAR	1.5	4.2	5.6	8.2	14	18	27	33	45	30	30	30	30	37	39	42	45	49
WPR	16	28	32	33	31	37	38	39	50	37	45	48	44	44	43	43	43	52
Global	11	16	18	22	24	28	32	37	45	37	40	40	40	42	42	43	45	5(

<sup>-</sup> Indicates not available.

st No additional data beyond DOTS report, either because country is 100% DOTS, or because no non-DOTS report was received.

FIGURE 16

Proportion of estimated new smear-positive (a) and of all estimated new cases (b) notified under DOTS (grey portion of bars) and non-DOTS (blue portion of bars), 2003. Figures indicate the number of cases (in thousands) represented by each portion of each bar.



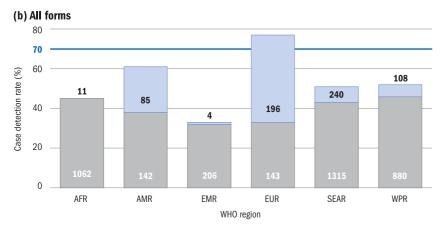
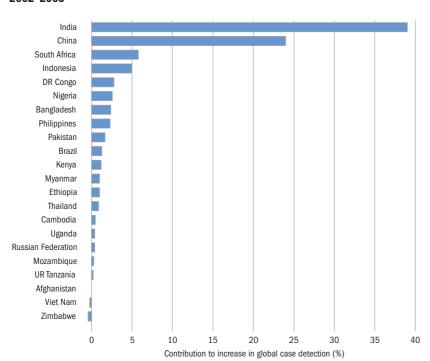


FIGURE 17 Contributions to the global increase in case detection made by high-burden countries, 2002-2003



grammes has been roughly the same in all WHO regions except the South-East Asia Region, where the acceleration in case-finding has been visible since 1998, driven mainly by DOTS expansion in India.

In the Region of the Americas, European Region and South-East Asia Region, significant numbers of smearpositive cases were reported, as usual, from outside DOTS programmes (Figure 16a). In the Region of the Americas, the estimated proportion of smear-positive cases detected from all sources exceeded 70%. Thus, the target for case detection would have been reached in this region if all patients in whom TB had been diagnosed had been treated under DOTS. There were similar differences among regions in the detection rates of all TB cases (Figure 16b).

Of the additional smear-positive cases reported by DOTS programmes in 2003 (compared with 2002), 63% were in India (39%) and China (24%; Figure 17). Although China and India have made big improvements in case detection, these two countries still account for an estimated 36% of all undetected smear-positive cases (Figure 18). They are among eight countries that together account for two thirds (67%) of all undetected cases in 2003. In order of importance, these are: India, China, Indonesia, Nigeria, Bangladesh, Pakistan, Ethiopia and the Russian Federation.

### Outcomes of treatment

More than 1.4 million new sputum smear-positive cases were registered for treatment in DOTS programmes in 2002, approximately the same number that were notified that year (Table 7, Annex 2). Discrepancies between the numbers of cases notified and registered for treatment were small globally, by region and for most HBCs, the largest differences being in Kenya and the Philippines (where about 10% of notified cases were not registered for treatment) and Afghanistan (where 20% of cases registered for treatment were not notified).

The cure rate among all cases registered under DOTS was 73%, and a further 9% completed treatment (no laboratory confirmation of cure), giv-

FIGURE 18  $\label{lem:countries} \textbf{Smear-positive TB cases undetected by DOTS programmes in six high-burden countries},$ 2003. Figures indicate the percentage of all cases missed globally which are missed by each country.

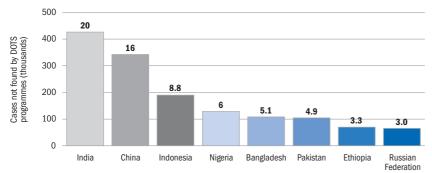


TABLE 7 Treatment outcomes for new smear-positive cases, DOTS strategy, 2002 cohorta

						TREATN	IENT OUTCO	MES (%) <sup>b</sup>			Ç	6 ESTIMATED CASES
	NOTIFIED®	REGISTER NUMBER	RED <sup>d</sup> %	CURED	COMPLETED TREATMENT	DIED	FAILED	DEFAULTED	TRANS- FERRED	NOT EVAL <sup>e</sup>	TREATMENT SUCCESS (%)	SUCCESSFULLY TREATED UNDER DOTS
1 India	245 135	244 859	100	86	1.0	4.4	2.5	6.1	0.4	0.0	87 †	27
2 China	180 239	180 239	100	90	2.5	1.3	0.9	1.0	1.8	2.6	93 †	28
3 Indonesia	76 230	76 230	100	72	15	2.0	1.3	4.9	2.6	3.0	86 †	24
4 Nigeria	19 596	20 559	105	69	10	1.7	6.7	11	1.5	0.0	79	11
5 Bangladesh	45 741	46 811	102	81	2.9	4.7	0.7	7.0	3.5	0.2	84	25
6 Pakistan	15 331	14 314	93	65	13	3.0	1.2	14	3.8	0.8	77	9.1
7 Ethiopia	36 541	36 541	100	59	17	6.6	0.7	5.0	9.8	1.7	76	27
8 South Africa	97 656	98 090	100	54	14	8.5	1.3	13	9.3	0.0	68	71
9 Philippines	65 148	59 453	91	77	11	2.6	1.1	5.3	2.9	0.5	88 †	49
10 Kenya	34 337	30 966	90	65	14	4.9	0.4	8.9	6.5	0.0	79	33
11 DR Congo	44 518	45 013	101	71	7.8	6.6	1.0	7.8	4.5	1.7	78	45
12 Russian Federation	5 179	5 171	100	64	2.7	13	9.1	7.2	4.1	0.0	67	4.6
13 Viet Nam	56 698	56 590	100	90	1.8	3.4	0.8	1.5	2.0	0.0	92 †	81
14 UR Tanzania	24 136	24 136	100	76	4.3	11	0.3	4.1	4.4	0.1	80	35
15 Brazil	4 835	4 606	95	46	29	6.2	0.3	7.8	11	0.0	75	6.9
16 Uganda	19 088	19 098	100	30	31	6.2	0.4	19	6.6	7.3	60	27
17 Thailand	25 593	26 559	104	69	5.3	11	1.7	9.5	4	0.0	74	50
18 Mozambique	15 236	15 236	100	77	1.4	11	1.3	7.4	2.3	0.3	78	35
19 Zimbabwe	15 941	15 941	100	62	5.6	11	0.1	6.6	15	0.0	67	32
20 Myanmar	24 162	23 922	99	71	10	5.4	1.9	9.2	2.4	0.0	81	52
21 Afghanistan	6 509	7 780	120	60	27	4.0	1.8	4.7	2.6	0.0	87 †	20
22 Cambodia	17 258	17 396	101	89	3.0	3.8	0.3	2.4	1.1	0.0	92 †	51
High-burden countries	1 075 107	1 069 510	99	76	7.0	4.6	1.5	6.2	3.3	1.0	83	29
AFR	442 729	437 873	99	60	13	7.2	1.4	11	6.6	1.4	73	31
AMR	82 312	78 180	95	64	17	4.8	1.0	5.4	3.4	4.3	81	38
EMR	74 450	74 799	100	72	12	3.1	1.4	7.8	2.6	1.5	84	22
EUR	43 112	40 307	93	63	13	6.2	6.2	5.9	2.2	3.1	76	16
SEAR	449 615	451 162	100	81	4.3	4.4	2.1	6.3	1.5	0.5	85 †	28
WPR	340 666	339 754	100	84	6.2	2.4	1.0	2.3	2.3	1.5	91 †	35
Global (DOTS)	1 432 884	1 422 075	99	73	8.7	4.8	1.6	6.7	3.4	1.4	82	30

Cases diagnosed during 2002 and treated/followed-up through 2003.

See Table 2 and accompanying text for definitions of treatment outcomes.

For AMR and EUR, the regional total of notified cases includes the number of laboratory confirmed (as opposed to smear-positive) cases for one country: USA and Israel, respectively.

If the number registered was provided, this (or the sum of the outcomes, if greater) was used as the denominator for calculating treatment outcomes. If the number registered was missing, then the number notified (or the sum of the outcomes, if greater) was used as the denominator. The number used as the denominator is shown in column labelled "Registered".

Eval: evaluated.

Estimated: estimated number of cases for 2002 (as opposed to notified or registered).

<sup>†</sup> Treatment success ≥85%.

TABLE 8
Treatment outcomes for new smear-positive cases, non-DOTS strategy, 2002 cohort<sup>a</sup>

	TREATMENT OUTCOMES (%) <sup>a</sup>											
	NOTIFIED	REGISTERED®	REGST'D (%)	CURED	COMPLETED TREATMENT <sup>a</sup>	DIED	FAILED	DEFAULTED	TRANS- FERRED	NOT EVAL	TREATMENT SUCCESS (%)	
1 India	150 698	41 368	27	41	17	1.5	1.4	30	9.3	0.0	58	
2 China	14 733	13 681	93	85	7.4	1.1	1.0	2.4	1.2	2.2	92 †	
3 Indonesia	_	_	_	_	_	_	_	_	_	_	_	
4 Nigeria	2 340	-	-	_	_	_	_	_	_	-	-	
5 Bangladesh	1 070	_	_	_	_	_	_	_	_	_	_	
6 Pakistan	934	-	-	_	-	_	_	_	_	-	-	
7 Ethiopia	_	_	_	_	_	_	_	_	_	_	_	
8 South Africa	1 143	1 239	108	59	6.2	5.4	2.4	13	8.2	5.6	65	
9 Philippines	_	_	_	_	_	_	_	_	_	_	_	
10 Kenya	-	-	_	_	_	_	_	_	_	-	-	
11 DR Congo	_	_	_	_	_	_	_	_	_	_	_	
12 Russian Federation	22 686	_	_	_	_	_	_	_	_	-	_	
13 Viet Nam	_	_	_	_	_	_	_	_	_	_	_	
14 UR Tanzania	-	_	_	_	_	_	_	_	_	-	_	
15 Brazil	36 536	24 246	66	24	57	5.9	0.3	12	0.6	0.0	81	
16 Uganda	_	_	-	_	-	-	-	-	-	-	-	
17 Thailand	_	_	_	_	_	_	_	_	_	_	-	
18 Mozambique	_	_	_	-	_	_	_	_	_	-	-	
19 Zimbabwe	_	_	_	_	-	_	_	_	_	_	_	
20 Myanmar	_	_	_	-	-	_	_	_	-	-	-	
21 Afghanistan	_	_	_	_	-	-	-	-	-	-	_	
22 Cambodia	_	_	_	_	_	_	_	_	-	-	-	
High-burden countries	203 140	80 534	35	44	27	2.8	1.0	20	5.3	0.5	71	
AFR	11 714	9 332	80	42	22	10	7.8	10	8.5	0.8	63	
AMR	49 603	30 985	62	29	49	5.8	0.5	13	2.0	0.5	78	
EMR	1 323	334	25	28	49	1.5	0.6	15	4.8	0.0	78	
EUR	40 450	8 622	21	47	21	4.4	4.5	6.6	0.7	15	69	
SEAR	157 115	43 784	28	42	17	1.6	1.5	28	9.0	0.1	59	
WPR	31 442	18 982	60	64	9.9	1.9	1.4	2.0	12	8.7	74	
Global (non-DOTS)	291 647	112 039	38	43	25	3.7	2.0	16	6.9	2.9	68	

<sup>-</sup> Indicates not available.

ing a reported, overall treatment success rate of 82%. An estimated 30% of all smear-positive cases arising in 2002 were treated successfully by DOTS programmes. In non-DOTS areas, the quality of reporting was worse: only four HBCs provided data for the 2002 cohort (Table 8).

By WHO region, the documented treatment success rates by DOTS programmes varied from 73% in the African Region to 85% in the South-East Asia Region and 91% in the Western Pacific Region, the latter two regions having met the 85% target (Table 7, Figure 19). Fatal outcomes were most common in the African Region (7%), where a higher fraction of cases are HIV-positive, and in the European Region (6%), where a higher fraction of cases are drug resistant

(eastern Europe), or occur among the elderly (western Europe). Treatment interruption (default) was most frequent in the African Region (11%) and the Eastern Mediterranean Region (8%). Transfer without follow-up was also especially high in the African Region (7%). Treatment failure was conspicuously high in the European Region (6%), mainly because a high proportion of patients in eastern Europe are recorded as failures.

DOTS treatment success exceeded 85% in seven HBCs (Table 7). It was under 70% in the Russian Federation, South Africa, Uganda and Zimbabwe. Treatment results for individual African countries once again point to the effects of HIV: cohort death rates were 9% or more in Mozambique, South Africa, the United Republic of

a See notes for Table 7.

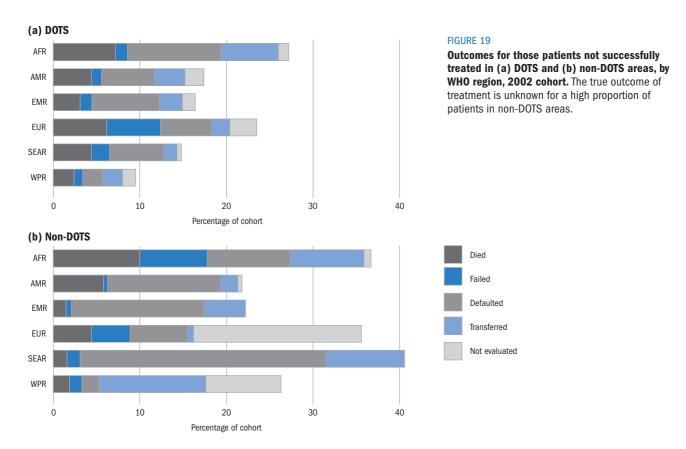


TABLE 9 Treatment success for new smear-positive cases (%), 1994-2002 cohorts<sup>a</sup>

_				DOTS	PROGRAM	MMES							WHO	OLE COUN	NTRY			
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1994	1995	1996	1997	1998	1999	2000	2001	2002
1 India	83	79	79	82	84	82	84	85	87	*	25	21	18	27	21	77	54	83
2 China	94	96	96	96	97	96	95	96	93	91	93	94	95	95	95	93	95	92
3 Indonesia	94	91	81	54	58	50	87	86	86	*	*	*	*	*	*	*	*	*
4 Nigeria	65	49	32	73	73	75	79	79	79	*	*	*	*	*	*	*	*	*
5 Bangladesh	73	71	72	78	80	81	83	84	84	*	*	63	73	77	79	81	83	*
6 Pakistan	74	70	_	67	66	70	74	77	77	69	*	_	*	23	*	*	*	*
7 Ethiopia	74	61	73	72	74	76	80	76	76	*	*	71	*	*	74	*	*	*
8 South Africa	_	-	69	73	74	60	66	65	68	78	58	61	68	72	57	63	61	68
9 Philippines	80	_	82	83	84	87	88	88	88	88	60	35	78	71	*	*	*	*
10 Kenya	73	75	77	65	77	78	80	80	79	*	*	*	*	*	79	*	*	*
11 DR Congo	71	80	48	64	70	69	78	77	78	72	74	48	64	*	*	*	*	*
12 Russian Federation	_	65	62	67	68	65	68	67	67	_	*	57	*	*	*	*	*	*
13 Viet Nam	91	91	90	85	93	92	92	93	92	*	89	89	85	92	92	*	*	*
14 UR Tanzania	80	73	76	77	76	78	78	81	80	*	*	*	*	*	*	*	*	*
15 Brazil	_	_	_	_	91	89	73	67	75	70	17	20	27	40	78	71	55	80
16 Uganda	_	_	33	40	62	61	63	56	60	_	44	*	*	*	*	*	*	*
17 Thailand	_	_	78	62	68	77	69	75	74	58	64	*	58	*	*	*	*	*
18 Mozambique	67	39	54	67	_	71	75	77	78	*	*	55	65	_	*	*	*	*
19 Zimbabwe	_	-	_	_	70	73	69	71	67	52	53	32	69	*	*	*	*	*
20 Myanmar	_	66	79	82	82	81	82	81	81	77	67	79	*	*	*	*	*	*
21 Afghanistan	_	_	_	45	33	87	86	84	87	_	_	_	*	*	86	85	*	*
22 Cambodia	84	91	94	91	95	93	91	92	92	*	*	*	*	*	*	*	*	*
High-burden countries	87	83	78	81	83	81	84	84	83	83	53	50	56	62	60	81	72	83
AFR	59	62	57	63	70	69	72	71	73	60	60	56	64	70	68	71	70	72
AMR	77	77	81	81	80	83	81	83	81	65	50	51	58	67	79	77	72	80
EMR	82	87	86	79	76	83	83	83	84	79	79	66	73	56	79	81	83	84
EUR	68	69	72	72	76	77	77	75	76	67	67	58	72	63	75	75	72	75
SEAR	80	74	77	72	72	73	83	84	85	66	33	31	29	40	34	79	63	83
WPR	90	91	93	93	95	94	92	93	91	87	80	72	91	92	91	90	91	90
Global	77	79	77	79	81	80	82	82	82	75	57	54	60	64	64	80	73	81

<sup>-</sup> Indicates not available.

<sup>\*</sup> No additional data beyond DOTS report, either because country is 100%, or because no non-DOTS report was received.

Tanzania and Zimbabwe. But programme performance also remains poor in some African countries. For example, more than 15% of patients were lost to follow-up in Ethiopia, Kenya, South Africa, Uganda and Zimbabwe. Large proportions of patients completed treatment without confirming cure (a final, negative sputum smear) in Ethiopia (17%) and Uganda (31%). Uganda reported the lowest proportion of successful treatments among the 22 HBCs (60%). The aggregated treatment results for the European Region are strongly influenced by the performance in the Russian Federation, where 13% of patients died, 9% failed treatment and 11% were lost to follow-up.

A comparison of treatment results

for nine consecutive cohorts (1994-2002) shows that the overall success rates have been 80% or more in DOTS areas since 1998 (Table 9). Treatment success rates have been persistently poor outside DOTS programmes in all regions, principally because large fractions of cases are not registered or evaluated.

In DOTS areas, about 250 000 cases were registered for re-treatment in 2002 (Table 10; Annex 2). Some patients remain on treatment (included with those "not evaluated"), but the latest data give an overall treatment success rate of 72%. When the three registration types (re-treatment after relapse, failure and default) are distinguished and compared with new TB patients, three patterns

emerge. First, the comparative success of re-treatment was consistent with expectations: lower on average for re-treatment (72%) than for new cases (82%), but higher for relapses (71%), intermediate for defaulters (68%) and lowest for failures (58%). The rank order relapse > default > failed held for six out of eight HBCs that provided data, and for five out of six WHO regions. Second, patients who defaulted from their first course of treatment tended to default when treated again (17% of patients that were re-treated after default failed to complete the subsequent course of treatment, compared with 11% among all re-treated patients and 7% of patients on their first course of treatment). This was true in all six WHO

TABLE 10 Re-treatment outcomes for smear-positive cases, DOTS strategy, 2002 cohorta

					TREAT	MENT OUTCOMES	S (%) <sup>a</sup>		
	REGISTERED <sup>a</sup>	CURED	COMPLETED TREATMENT <sup>a</sup>	DIED	FAILED	DEFAULTED	TRANS- FERRED	NOT EVAL	TREATMENT SUCCESS (%)
1 India	84 078	68	3.3	7.2	6.1	14	0.7	0.0	72
2 China	46 932	83	5.2	2.6	3.8	2.0	1.0	0.0	88 †
3 Indonesia	3 731	60	17	2.2	3.3	5.4	3.1	8.4	78
4 Nigeria	2 373	63	11	4.7	7.6	11	1.1	2.5	73
5 Bangladesh	4 360	66	2.6	4.3	1.9	10	2.9	12	69
6 Pakistan	2 871	33	43	4.5	1.8	11	5.2	1.9	76
7 Ethiopia	1 716	52	8.6	6.9	2.8	5.3	2.4	22	60
8 South Africa	28 755	43	10	11	2.1	17	10	6.8	53
9 Philippines	_	_	-	-	-	-	-	-	_
10 Kenya	2 476	65	12	10	0.4	6.7	5.8	0.0	77
11 DR Congo	4 618	61	5.8	9.3	3.6	9.8	7.3	3.3	67
12 Russian Federation	962	37	9.4	12	26	8.6	8.1	0.0	46
13 Viet Nam	6 079	79	5.5	4.9	5.4	2.3	2.4	0.0	85 †
14 UR Tanzania	2 081	71	6.4	13	0.7	4.6	3.7	0.5	77
15 Brazil	640	36	24	7.0	1.1	18	13	0.0	60
16 Uganda	2 555	28	27	10	0.9	16	5.3	13	55
17 Thailand	1 990	55	6.4	17	6.7	9.3	5.7	0.0	62
18 Mozambique	1 721	65	1.2	12	2.2	9.4	4.0	5.7	67
19 Zimbabwe	1 371	58	4.7	20	0.9	7.6	8.7	0.0	63
20 Myanmar	8 036	65	10	7.8	3.9	10	3.7	0.2	75
21 Afghanistan	_	_	-	_	_	_	_	-	_
22 Cambodia	875	86	2.9	5.8	1.5	2.7	1.1	0.0	89 †
High-burden countries	208 220	66	6.5	6.8	4.5	11	2.9	2.4	73
AFR	59 574	49	11	10	2.4	14	7.7	6.7	59
AMR	7 635	64	6.8	5.8	4.3	12	4.3	2.3	71
EMR	8 825	51	23	4.4	3.5	10	5.3	3.2	74
EUR	12 551	42	12	10	12	11	2.6	10	54
SEAR	106 423	68	4.3	7.1	5.6	13	1.3	0.8	72
WPR	57 071	81	5.6	3.1	4.0	2.3	1.8	1.9	87 †
Global	252 079	64	7.2	6.9	4.7	11	3.2	3.1	72

Indicates not available

<sup>†</sup> Treatment success ≥85%.

See notes for Table 7.

regions. Third, the regional distribution of adverse re-treatment outcomes resembled the pattern observed for new cases. Thus, countries in the African Region reported high death rates (10%), and many patients were lost to follow-up (28%). Countries in the European Region reported high rates of death (10%) and treatment failure (12%).

### Trends in case detection and treatment success: overview of national DOTS programmes

Data on both treatment success and case detection were provided by 177 DOTS countries. Case detection exceeded 50%, and treatment success exceeded 70%, in 75 countries (Figure 20). They include the HBCs Cambodia, the Democratic Republic of the Congo, Myanmar, the Philippines, Thailand and Viet Nam. Of these countries, 22 appear to have reached the WHO targets, but together the 75 countries accounted for only 17% of all new smear-positive cases in 2003. Viet Nam was still the only member of the current group of HBCs to have reached targets for both case detection (>70%) and treatment success (>85%), although Myanmar and the Philippines are close to these targets (Figure 21). Three HBCs - Brazil, the Russian Federation and Uganda - had low rates of both case detection (<50%) and treatment success (<70%). More details of progress in each of the 22 HBCs can be found in the profiles (Annex 1).

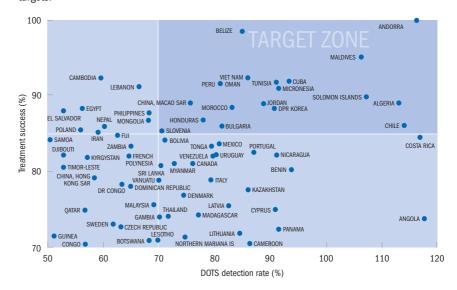
Of 161 countries that provided data for both the 2001 and 2002 cohorts, 91 (57%) showed higher treatment success rates for the 2002 cohort, and 68 of 172 (40%) improved case detection by more than 5%. Annex 2 tabulates case detection and treatment success rates by country over the nine years for which there are data.

### Trends in prevalence and death rates

The trends in prevalence and mortality for each region are calculated from the trend in incidence (Figure 6) and from estimates of the duration of illness (e.g. time smear-positive) and the case-fatality rate. Summing estimates from across the regions gives the global trends in prevalence and

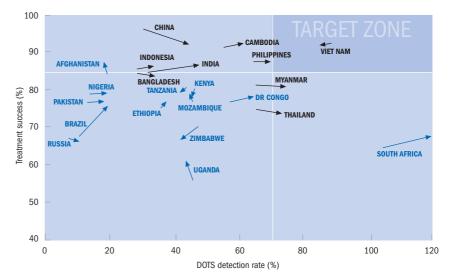
### FIGURE 20

DOTS status in 2003: countries close to targets. 75 countries reported treatment success rates 70% or over and DOTS detection rates 50% or over. 22 countries (including Kiribati, Wallis & Futuna Islands and Marshall Islands, out of range of graph) have reached both



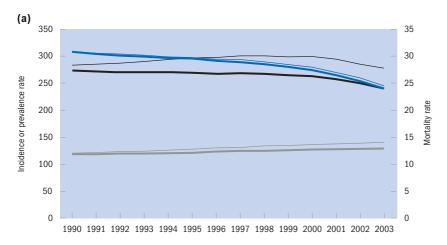
#### FIGURE 21

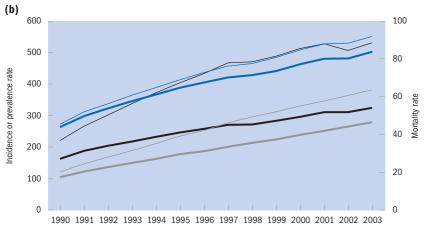
DOTS progress in high-burden countries, 2002–2003. Treatment success refers to cohorts of patients registered in 2001 or 2002, and evaluated by the end of 2002 or 2003, respectively. Countries in AFR, AMR, EMR and EUR are shown in blue; those in SEAR and WPR are shown in black.



### FIGURE 22

(a) Estimated global TB incidence (per 100 000 population per year; grey), prevalence (per 100 000 population; blue), and mortality rates (per 100 000 population per year; black), 1990-2003, including (thin lines) or excluding (thick lines) TB patients coinfected with HIV. (b) As for (a), but for African countries with high rates of HIV infection ( $\geq$ 4% in adults aged 15–49 years).





deaths, which are shown in Figure 22a with and without the contribution of HIV-positive TB patients, and in comparison with the trends in incidence (Figure 7). Although the global incidence rate was still increasing in 2003, prevalence and death rates had already begun to fall. Excluding HIVpositive TB patients, the incidence rate increased 0.6% between 2002 and 2003, prevalence fell by 5.7% and mortality by 3.5%. When TB patients coinfected with HIV are included, the incidence rate increased by 1.0%, and prevalence and mortality fell by 5.5% and 2.5%, respectively. The differential effects of HIV on incidence, prevalence and mortality are also visible, and magnified, in the trends for countries of eastern and southern Africa, where all three indicators were still

increasing in 2003 (Figure 22b). Among the nine regions defined in Figure 6, TB prevalence and death rates increased between 2002 and 2003 only in the two African regions; as for incidence rates, prevalence and deaths were falling or stable in the other seven regions.

### **Planning and DOTS implementation**

### TB control in the context of the health-care system

Country profiles incorporate information from the summary planning tables that were prepared for the 2004 DEWG meeting and from the questionnaires submitted by all 22 HBCs. The health systems of many countries are still undergoing reform and restructuring. However, all HBCs except the Russian Federation and Thailand reported that TB control functions are fully integrated with essential national health services. The MoH generally provides support for TB control through a specific technical unit, although in Bangladesh, South Africa and Thailand this function needs to be strengthened. All HBCs have a national plan for TB control and many will conduct, during 2005, a new planning exercise for the next five years. A total of 15 HBCs have prepared, or are developing, a plan for human resource development (HRD) reflecting their specific needs in the context of the health system.

### Constraints and remedial actions

In summary tables and questionnaires, countries reported the following constraints:

- 1. Shortage or inadequate capacity of staff. This remains a major constraint identified by 18 HBCs. The problem is being addressed by various means: situation assessments, development of HRD plans, intensification of training and supervision, redistribution of staff and appointment of new staff. All HBCs except Brazil, Nigeria, South Africa and Zimbabwe plan to implement projects funded by the GFATM in 2005; a shortage of managers, or inadequate managerial capacity, will almost certainly hinder these projects.
- 2. Inadequate central management capacity. Insufficient capacity at the highest levels delays the implementation of national plans, as reported in Bangladesh, Mozambique, South Africa, Uganda, the United Republic of Tanzania and Zimbabwe. Support from technical partners has been the main temporary remedial action.
- 3. Inadequate infrastructure. Lack of transportation infrastructure (roads and vehicles), poor communication networks, unreliable or non-existent electricity supplies, inadequate buildings and equipment, and weak primary health-care systems all impede TB control. A total of 12 HBCs reported deficiencies in at least one of these areas.

- 4. Weak political commitment. Ethiopia, Mozambique, Nigeria, South Africa, Thailand and Zimbabwe reported limited commitment to TB control from central and peripheral levels. Remedial actions include providing better support to local government following decentralization, forming provincial task forces, expanding international support through high-level advocacy missions and advocacy for TB control in civil society, especially in support of patients infected with HIV. Brazil, China and the Russian Federation reported significant progress on legislation to support TB control.
- 5. Weak laboratory services. The main obstacles are summarized in Table 11.
- 6. Nearly all HBCs had a secure supply of anti-TB drugs in 2003, thanks in large part to the Global TB Drug Facility (GDF). Mozambique reported drug shortages, but these will be rectified following a successful application to the GDF. South Africa experienced

- drug shortages due to the phasing in of the new drug combinations and a complete stock-out of streptomycin when the sole supplier stopped production.
- 7. Poor monitoring and evaluation. Timely and reliable data are essential for monitoring trends and for planning corrective actions. The Russian Federation and South Africa have addressed problems reported in 200313 by establishing standardized recording and reporting systems. China has introduced a new Internet-based reporting system, but the data being collected need validation.
- 8. Insufficient funds. A lack of money is no longer one of the major constraints identified by most HBCs. The governments of the wealthier HBCs make large contributions to TB control, international donors have increased their investments and the GFATM has begun to bridge financial gaps. As a result, some NTPs now have sufficient funding to expand DOTS programmes. However, some of the
- HBCs did report shortfalls in their 2004 budgets. Some of these countries still report gaps (see section on Financing DOTS Expansion), and others have problems in distributing funds from local or central governments to programmes (e.g. Nigeria). Mozambique has inadequate funds to pay salaries, and Zimbabwe is heavily affected by the country's general financial crisis.
- Poor access to remote areas. Access to geographically remote and politically unstable areas is a challenge in Afghanistan, Bangladesh, the Philippines and Uganda. The NTP in Viet Nam, having reached the targets for DOTS implementation, has made service provision in remote areas an important part of consolidating programme success.
- 10. Low public awareness. Limited knowledge about TB and its treatment, and the stigma of having TB (and perhaps also HIV infection), both hamper efforts to detect and treat TB suspects. A

TABLE 11 Status of TB laboratory services, high-burden countries, 2003-2004

	POPULATION PER DIAGNOSTIC FACILITY	INSUFFICIENT EQUIPMENT OR SHORTAGE OF SUPPLIES	SHORTAGE OF STAFF (OR EXISTING STAFF IN NEED OF TRAINING)	QUALITY ASSURANCE	CULTURE	DRUG SUSCEPTIBILITY TESTING (DST)	NATIONAL REFERENCE LABORATORY (NRL)
1 India	100 000		Υ	limited, planned	Υ	Υ	Υ
2 China	500 000		Υ	limited	Υ	Υ	Υ
3 Indonesia	80 000		Υ	limited	Υ	Υ	N, one acting
4 Nigeria	150 000	Υ	Y	limited, planned			Y
5 Bangladesh	230 000			limited	limited	limited	Υ
6 Pakistan	100 000	Υ	Υ	N	limited	limited	Y
7 Ethiopia	180 000		Υ	limited	Υ	Υ	Υ
8 South Africa	unknown <sup>a</sup>			Υ	Υ	Y	private
9 Philippines	75 000		to be trained	limited	limited	limited	Y
10 Kenya	54 000		Υ	Υ	Υ	Y	Υ
11 DR Congo	70 000	Υ	Υ	limited	limited	limited	Υ
12 Russian Federation	25 000		to be trained	Υ	Υ	Y	N
13 Viet Nam	125 000			Υ	limited	limited	Υ
14 UR Tanzania	65 000		Υ	weak	limited	limited	Υ
15 Brazil	45 000		to be trained	limited	Υ	Υ	Υ
16 Uganda	60 000		Υ	planned	Υ	Y	Υ
17 Thailand	65 000		to be trained	Υ	Υ	Υ	Υ
18 Mozambique	60 000	Υ	Υ	N	limited	limited	Υ
19 Zimbabwe	120 000	Υ	Υ	limited	limited	limited	Υ
20 Myanmar	200 000		Υ	weak	limited	limited	Υ
21 Afghanistan	100 000	Υ	Υ	N	limited	limited	N
22 Cambodia	75 000		to be trained	limited	Υ	N	Υ

<sup>&</sup>quot;Y" or "N" indicates whether the problem exists or the activity is undertaken.

<sup>&</sup>quot;Limited" indicates that the activity is restricted to certain areas of the country.

<sup>&</sup>lt;sup>a</sup> Laboratory services contracted out to National Health Laboratory Service in eight of nine provinces.

### FIGURE 23

The number of TB cases that were tested for HIV for every 100 TB cases that were notified in 2003 for countries that reported more than 1000 cases and tested more than 1% of notified cases. The countries are: ARM Armenia; AUS Australia; AZE Azerbaijan; BOT Botswana; BRA Brazil; CAM Cambodia; CNG Congo; CUB Cuba; DJI Djibouti; ELS El Salvador; GEO Georgia; GHA Ghana; GUT Guatemala; HOK Hong Kong SAR; HON Honduras; IVC Côte d'Ivoire; KOR Republic of Korea; LIY Libyan Arab Jamahiriya; LVA Latvia; MAL Malawi; MEX Mexico; NAM Namibia; NIC Nicaragua; PAN Panama; PNG Papua New Guinea; POR Portugal; SOA South Africa; SUD Sudan; SYR Syrian Arab Republic; UZB Uzbekistan; VTN Viet Nam.

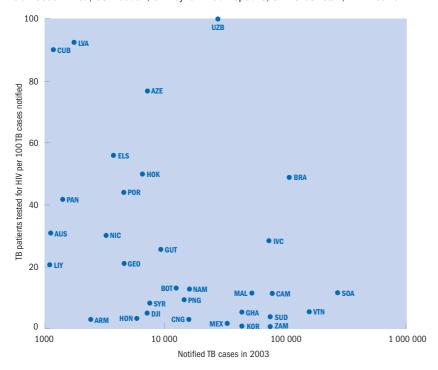
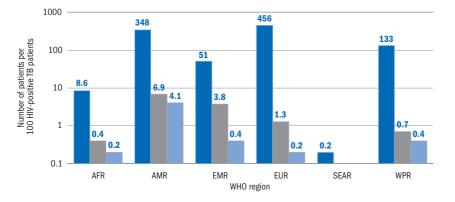


FIGURE 24

The number of TB patients tested for HIV (blue), assessed for ART (grey) and starting ART (pale blue) for every estimated 100 HIV-positive TB patients in each WHO region



total of 13 HBCs have plans to launch or intensify advocacy and communication campaigns.

### Intensified support and action in countries

During 2004, the DEWG launched ISAC, an emergency initiative to reach targets for DOTS implementation by 2005 and to generate further momentum towards the MDG targets for 2015. The goal of ISAC is to rapidly increase managerial capacity for TB control at central and intermediate levels of administration. Participating countries include China, India, Indonesia, Kenya, Pakistan, Romania, the Russian Federation and Uganda.

### Partnerships, coordination and advocacy

All HBCs have some mechanism for coordinating TB control activities. Most countries have an NICC that meets regularly to share information on planning and progress. This has served as a model for the creation of Global Fund Country Coordination Mechanisms (CCMs). However, in some countries (e.g. Ethiopia, Viet Nam) the CCM has become the main coordinating body. During 2004, Indonesia, Pakistan and Uganda formed and launched national partnerships to Stop TB, in order to establish collaborations among various stakeholders (NTP, WHO, technical and financial partners, NGOs, and patients' associations), and to share human and financial resources to address more effectively some of the constraints hindering NTP performance. NGOs are actively collaborating with NTPs to improve service coverage in 20 HBCs (Annex 1).

Most of the HBCs recognize the need for improved advocacy and communication on TB control. Ethiopia, India, Kenya, Pakistan, South Africa and Viet Nam reported that such activities were intensified in 2004. Cambodia, the Democratic Republic of the Congo, Indonesia, Nigeria, the Philippines, the United Repbulic of Tanzania and Uganda are planning advocacy and communication campaigns for 2005.

### Management of drug resistance

Among the HBCs, Kenya (pilot site in Nairobi), the Philippines (pilot site in Manila) and the Russian Federation (Archangelsk, Ivanovo, Orel and Tomsk oblasts) have DOTS-Plus pilot projects approved by the GLC. The projects in Kenya and the Philippines are supported financially by the GFATM, as is one of the four projects in the Russian Federation (Tomsk). In 2005, applications to the GLC are expected from Bangladesh, Myanmar, the Philippines, the United Republic of Tanzania and Viet Nam.

By December 2004, the GLC had approved 30 DOTS-Plus pilot projects for a total of 10 133 MDR-TB patients in 23 countries.25 However, only three HBCs - Brazil, the Russian Federation and South Africa - have national policies for the diagnosis and treatment of MDR-TB, and manage MDR-TB under the NTP. Even in the few countries that do have policies, MDR-TB treatment often fails to meet acceptable standards in practice. Second-line drugs are available in almost all HBCs, and are locally produced in Bangladesh, Brazil, China, Kenya, India, Indonesia, the Philippines, Pakistan, the Russian Federation, South Africa, Thailand and Viet Nam. In many countries, substandard MDR-TB treatment is available in the private sector or at specialized health centres, often for a fee.

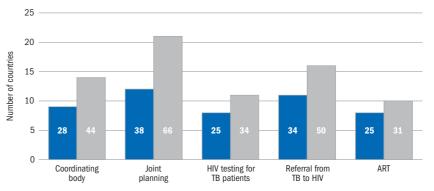
The planning of activities related to MDR-TB is described in the individual profiles of the 22 HBCs (Annex 1).

### Collaborative TB/HIV activities

Among the 199 countries that completed the WHO data collection form, 49% have a national policy of offering HIV testing to TB patients, the first step in accessing appropriate prevention and care services for HIV-positive TB patients. However, in 2003, only 3% of 4.4 million notified TB cases were reported to have been tested for HIV. Of these 199 countries, 46 (23%)

### FIGURE 25

Progress in developing national policy on collaborative TB/HIV activities, 2002 (blue bars) and 2003 (grey bars). Data are for the 32 countries with high numbers of people with TB who are also HIV-infected which returned the supplementary questionnaire. The bars show the number of countries that have a coordinating body, carry out joint planning for TB and HIV control activities, have a policy of testing TB patients for HIV, have a system for referring TB patients for HIV care and support and that provide ART in the public sector. The numbers in each bar give the percentage of the countries that fall into each category.



indicated that HIV-positive TB patients were routinely assessed for their eligibility for ART. Only 1349 TB patients were reported to have started ART in 2003.

Figure 23 shows the breakdown of HIV testing rates among TB patients by country, for all countries that notified more than 1000 TB patients in 2003 and tested more than 1% of them for HIV. Even in Brazil, where ART is provided free of charge in the public sector, only half of the notified TB patients were reported to have been tested for HIV. Among other countries that notified more than 10 000 cases each year, Côte d'Ivoire tested less than 30%. Botswana, Cameroon, Malawi, Namibia and South Africa tested about 10%. For other high-incidence countries, the proportions tested were still lower. Some countries with low TB incidence rates, including Cuba and Latvia, tested most of their TB patients.

Regional differences in HIV testing, and in assessment for and provision of ART, are shown in Figure 24, where the denominator is the estimated number of new HIV-positive TB patients in 2003. Under normal circumstances, several TB patients would have to be tested in order to detect one HIV-positive patient. Thus the number of patients tested for HIV should be several times greater than the number estimated to be HIV positive, but the number tested exceeded the estimated number only in the Re-

gion of the Americas, European Region and Western Pacific Region.

By region, as by country, the number of HIV-positive TB patients who were assessed for ART is much smaller than the number who were tested for HIV. In the Region of the Americas, seven HIV-positive TB patients were assessed for ART for every 100 estimated to be HIV-positive, and most of these were in Brazil. In the African Region, the region worst affected by HIV/AIDS, only four patients were assessed for ART for every 1000 HIV-positive TB patients. The percentage of HIV-positive TB patients reported to have started ART was still lower. The Region of the Americas performed better than other regions, but fewer than four out of 1000 HIV-positive TB patients started ART.

Of the supplementary questionnaires sent to 41 high-incidence countries, 32 were returned to WHO. The data show that, between 2002 and 2003, TB/HIV collaboration had improved (Figure 25). However, implementation and recording and reporting remain weak. For example, few countries were able to report the exact numbers of TB patients or HIV-infected people that were benefiting from these activities.

### Additional strategies for DOTS expansion

Public public and public private mix for DOTS (PPM). During 2004, 13 HBCs reported improved links between the

<sup>&</sup>lt;sup>25</sup> Bolivia, Costa Rica, El Salvador, Egypt, Estonia, Georgia, Haiti, Honduras, Jordan, Kenya, Kyrgyzstan, Latvia, Lebanon, Malawi, Mexico, Nepal, Nicaragua, Peru, Philippines, Romania, Russian Federation, Syrian Arab Republic and Uzbekistan.

TABLE 12 Additional strategies for DOTS expansion planned or implemented, high-burden countries

	PPM (PUBLIC-PUBLIC AND PUBLIC-PRIVATE MIX)	COMMUNITY TB CARE	PRACTICAL APPROACH TO LUNG HEALTH (PAL)
1 India	scale-up	nationwide	
2 China	referral, planned	some areas	
3 Indonesia	pilot	pilot, scale-up	planned
4 Nigeria	planned	planned	planned
5 Bangladesh	scale-up	nationwide	
6 Pakistan	pilot		
7 Ethiopia	pilot	pilot	
8 South Africa	with mining companies	community health workers	
9 Philippines	scale-up	some areas, scale-up	
10 Kenya	scale-up	pilot, scale-up	
11 DR Congo	planned	pilot, scale-up	planned
12 Russian Federation		through NGOs	planned
13 Viet Nam	planned	IEC, scale-up	planned
14 UR Tanzania	pilot	nationwide	planned
15 Brazil			
16 Uganda	planned		
17 Thailand	pilot	some areas, scale-up	planned
18 Mozambique		some areas, scale-up	planned
19 Zimbabwe	pilot	some areas, with TB/HIV	
20 Myanmar	pilot		
21 Afghanistan	planned	Kabul only	
22 Cambodia	planned	pilot	planned

NTP and other health-care providers (Table 12); 14 have established better collaborations with medical colleges; 10 have PPM pilot projects in various stages of implementation. Seven more HBCs are planning PPM initiatives for 2005, and four (India, Kenya, Myanmar and the Philippines) are attempting to implement PPM initiatives nationally. China, India, Indonesia, Kenya and Pakistan have specific plans to scale-up collaborations between NTP and non-NTP public hospitals.

Community TB care. A total of 17 HBCs reported some form of community contribution to TB care. In Bangladesh, India and Uganda, NGOs and community groups have played a vital part in expanding access to TB treatment. Communities are involved in TB care in limited parts of Afghanistan, Cambodia, China, the Democratic Republic of the Congo, Ethiopia, Indonesia, Kenya, Mozambique, the Philippines, South Africa, the United Republic of Tanzania and Viet Nam. Nigeria is planning to carry out a pilot study in 2005.

TABLE 13 Budget and expenditure data received, all countries, 2005

	NUMBER OF	REPORTS		BUDGET DATA			EXPENDITURE DATA				
	COUNTRIES	RECEIVED	COMPLETE	PARTIAL	NONE	COMPLETE	PARTIAL	NONE	TREATED QUANTIFIED		
AFR	46	35	26	6	3	23	5	7	33		
AMR	44	29	15	6	8	12	6	11	21		
EMR	22	17	5	7	5	6	5	6	15		
EUR	52	16	5	9	2	8	5	3	16		
SEAR	11	8	7	1	0	7	0	1	7		
WPR	36	29	12	10	7	13	9	7	28		
Global	211	134	70	39	25	69	30	35	120		

TABLE 14 Budget and expenditure data received, high-burden countries, 2005

Global	22	21	19	2	1	17	2	3	20
WPR	4	4	4	0	0	4	0	0	4
SEAR	5	5	4	1 <sup>d</sup>	0	4	1 <sup>d</sup>	0	4
EUR <sup>c</sup>	1	1	1	0	0	1	0	0	1
EMR	2	2	1	1 <sup>b</sup>	0	1	1 <sup>b</sup>	0	2
AMR	1	1	1	0	0	1	0	0	1
AFR	9	8	8	0	1	6	0	3 <sup>a</sup>	8
	COUNTRIES	RECEIVED	COMPLETE	PARTIAL	NONE	COMPLETE	PARTIAL	NONE	TREATED QUANTIFIED
	NUMBER OF	REPORTS		BUDGET DATA			XPENDITURE DATA		NO. PATIENTS TO BE

Kenya, South Africa and Uganda.

Afghanistan.

Data for the Russian Federation were prepared by WHO staff (Moscow office). See country profile for further details.

Practical Approach to Lung Health (PAL). Cambodia, the Democratic Republic of the Congo, Indonesia, Mozambique, Nigeria, Russian Federation, the United Republic of Tanzania, Uganda and Viet Nam are planning to investigate the feasibility of implementing PAL in 2005. These studies will investigate how the syndromic approach to diagnosis and treatment can influence TB case detection and the rationalization of drug prescription practices.

### **Financing DOTS expansion**

### Data received

Financial data were received from 134 out of 211 (64%) countries (Table 13), more in total than for 2004 (123 countries), but with fewer reports from the European Region. Complete budget data were provided by 70 countries (compared with 77 in 2004), and 69 provided complete expenditure data (down from 74 in 2004). Fewer complete reports were provided by the European Region and the Western Pacific Region, perhaps because more data were requested for the present report (in particular, two years of budget data rather than one). The main improvement in reporting was in the African Region, where the number of complete budget and expenditure forms increased by 37% and 44% respectively, probably because WHO

regional office staff intensively followed up on data collection with NTPs.

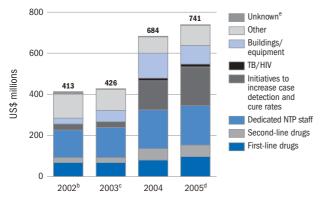
Data were received from all 22 HBCs except South Africa (Table 14), providing the most complete set of data since financial monitoring was introduced by WHO in 2002. Complete budget data were provided for 19 countries (up from 17 in 2004); data were missing for South Africa and only partially complete for Afghanistan and Thailand. Complete expenditure data were provided for 17 countries (up from 15 in 2004); Kenya, South Africa and Uganda provided no data, and Afghanistan and Thailand provided incomplete data. A total of 20 countries made projections of the number of cases they would treat in 2004 and 2005, compared with the 16 countries that provided projections for 2004 in last year's report. Again, the main improvement in reporting was in the African Region. Countries in the Western Pacific Region as well as India submitted, on time, exemplary data that required minimal follow-up.

### Total NTP budgets and funding in HBCs

NTP budgets in 18 of the 22 HBCs have increased during the period 2002-2005, sometimes by substantial amounts (Figure 26, Figure 27, Table 15 columns 3 and 4). The total combined budgets for 2004 and 2005 are US\$ 684 million and US\$ 741 million, respectively, compared with around US\$ 400 million in both 2002 and 2003. The main reason why the budgets for 2004 and 2005 are higher than in previous years is that countries are aiming to detect and treat more patients. This is associated with large proposed spending increases on initiatives to increase case detection and cure rates (US\$ 160 million 2002-2005, of which US\$ 39 million is in China), investment in buildings and equipment (US\$ 63 million 2002-2005), and dedicated NTP staff (US\$ 57 million 2002-2005). Important increases are also budgeted for first and second-line drugs (both up by about US\$ 30 million 2002–2005). Relatively small budgets were reported for collaborative TB/HIV activities.

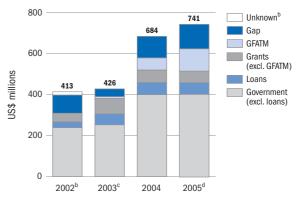
The countries with by far the largest budgets for 2005 (Table 15 column 2) are the Russian Federation (US\$ 316 million) and China (US\$ 158 million), followed by India and Indonesia (both around US\$ 45 million). Other countries reported budgets of around or less than US\$ 20 million. In absolute terms, China reported the largest budgetary increase between 2002 and 2005 (an additional US\$ 60 million; Table 15, column 3). China is committed to achieving the 70% case detection target in 2005, and the budgetary increases reflect plans to achieve this.

FIGURE 26 Total NTP budgets by line item 2002-2005, 21 high-burden countries, a 2002-2005



- Data not available for South Africa.
- <sup>b</sup> Estimates assume budget 2002 equal to expenditure 2002 (Ethiopia), budget 2003 (Afghanistan, Bangladesh, Mozambique and Uganda) and expenditure 2003 (Russian Federation and Zimbabwe)
- <sup>c</sup> Estimates assume budget 2003 equal to expenditures 2003 for Mozambique, Russian Federation and 7imbabwe
- Budget data for UR Tanzania based on 2004 data.
- "Unknown" applies to Mozambique in 2002 and Afghanistan 2002-2005, as breakdown by line item not available.

FIGURE 27 Total NTP budgets by source of funding, 21 high-burden countries, a 2002-2005



- a Data not available for South Africa.
- <sup>b</sup> Estimates assume budget 2002 equal to expenditure 2002 (Ethiopia), budget 2003 (Afghanistan, Bangladesh, Mozambique and Uganda) and expenditure 2003 (Russian Federation and Zimbabwe). "Unknown" applies to DR Congo and Nigeria, as breakdown by funding source not available.
- Estimates assume budget 2003 equal to expenditures 2003 for Mozambique, Russian Federation and Zimbabwe.
- <sup>d</sup> Budget data for UR Tanzania based on 2004 data.

TABLE 15 NTP budgets and available funding, high-burden countries, 2005

	TOTAL NTP BUDGET	CHANGE FROM 2002	CHANGE FROM			VAILABLE FUNDIN (US\$ MILLIONS)	IG		CHANGE II		E FUNDING SIN		SOURCE
	(US\$ MILLIONS)	(US\$ MILLIONS)	2002	GOVERNMENT (EXCL. LOANS)	LOANS	GRANTS (EXCL. GFATM)	GFATM	GAP	GOVERNMENT (EXCL. LOANS)	LOANS	GRANTS (EXCL. GFATM)	GFATM	GAP
1 India	46	10	29	5	11	6	8	15	-1	-13	0.7	8	15
2 China	158	60	61	98	14	4	21	21	45	14	2	21	-22
3 Indonesia	43	9	25	24	0	4	15	0	17	0	1	15	-25
4 Nigeria <sup>a</sup>	12	3	35	2	0	3	0	7	0.5	0	-2	0	-0.05
5 Bangladesh <sup>b</sup>	22	15	210	3	6	3	8	1	-0.1	6	-0.6	8	1
6 Pakistan	19	14	257	8	0	3	0	9	5	0	2	0	7
7 Ethiopia <sup>c</sup>	7	2	40	0.6	0	1	5	0	-0.6	0	-3	5	0
8 South Africad	-	-	-	-	-	-	-	-	-	-	-	-	-
9 Philippines	8	0.5	6	3	0	4	2	0	-0.7	0	4	2	-4
LO Kenya	14	9	177	3	0	0	3	8	2 e	0	-3	3	7
11 DR Congo <sup>a</sup>	11	4.2	64	0.6	0	5	2	3	-0.4	0	-0.3	2	-0.8
12 Russian Federation <sup>f</sup>	316	178	129	220	25	2	30	39	90	25	-6	30	39
13 Viet Nam	12	0.4	4	9	0	0.9	2	0.3	0.5	-2	-0.1	2	0.3
14 UR Tanzania <sup>g</sup>	9	3	59	1	0	5	0.2	2	1	0	0.6	0.2	1
15 Brazil	21	8	59	19	0	2	0	0	6	0	2	0	0
L6 Uganda <sup>b</sup>	6	1	22	0.7	0	0.8	0.9	4	0.6	-1	0.3	0.9	0.7
17 Thailand	5	-1	-23	3	0	0	2	0	-3	0	0	2	0
18 Mozambique <sup>b</sup>	7	-1	-10	0.8	0.2	2	4	0.1	0.5	0.2	-0.5	4	-5
19 Zimbabwe <sup>f</sup>	11	9	536	0.5	0	2	0	9	0.4	0	0.04	0	9
20 Myanmar	5	2	86	0.4	0	1	3	0.6	-0.03	0	1	3	-2
21 Afghanistan <sup>b</sup>	3	-0.3	-10	0.3	0	2	0.7	0	0	0	0.5	0.7	-2
22 Cambodia	7	3	60	0.7	0	4	1	1	-0.6	-0.7	3	1	0.01
ligh-burden countries	741	328	59 h	402	56	55	109	119	161	28	3	109	20

The increase in the Russian Federation could be larger (US\$ 178 million), if it is assumed that the 2002 budget was similar to reported expenditures (budget data were not reported for 2002). In 2003, the Russian Federation developed an ambitious five-year plan to expand DOTS and to upgrade TB control in general, covering the period 2003-2007. For other countries, the budget differences between 2002 and 2005 are all US\$ 15 million or less.

In relative terms, the biggest budget increases are for Bangladesh and Pakistan (both more than 200%), followed by Kenya at 177% (Table 15, column 4). Six countries reported changes of 50-100% (Brazil, Cambodia, China, the Democratic Republic of the Congo, Myanmar, the United Republic of Tanzania), and four of 2550% (Ethiopia, India, Indonesia, Nigeria). When compared with expenditures rather than budgets, the increases since 2003 are enormous for Zimbabwe (536%) and large for the Russian Federation (129%).

These large budget increases have been accompanied by big improvements in available funding for NTPs (Table 15 columns 5-8 and 10-13; Figure 27). For all HBCs, available funding has increased by about US\$ 300 million since 2002, reaching US\$ 622 million in 2005. In 2005, HBC governments will provide 62% of the required funding (including loans), the GFATM 15% and grants from other sources 7%, leaving a gap equivalent to 16% of the reported budgets. However, sources of funding vary among the 22 HBCs (Figure 28), with a few countries relying mostly on government funding but most relying extensively on grants from the GFATM and other sources. Most of the increased funding since 2002 is from governments (an increase of US\$ 189 million since 2002, including loans, almost all of which is in China and the Russian Federation) and the GFATM (US\$ 109 million for 17 HBCs in 2005 compared with no contribution in 2002). There has been virtually no change in grant funding from sources other than the GFATM.

Despite this progress in securing additional funding, there is a large funding gap of US\$ 119 million in 2005 (Table 15 column 9), which is higher than the gaps reported for 2003 and 2004. In absolute terms, the largest funding gaps are those reported by China, India, Pakistan, the Russian Federation and Zimbabwe

Available funding compared with 2003, as no funding breakdown was provided in 2002; thus total of changes in available funding by source (US\$ 321 million) does not equal the total shown in column 3 (US\$ 328 million).

Comparisons are with budget for 2003, as budget data for 2002 not available.

Comparisons are with expenditure for 2002, asf budget data for 2002 not available.

No data were provided by the NTP.

May include some loan funding.

Comparisons are with expenditure for 2003, as budget data for 2002 or 2003 not available. Figures for Russian Federation for 2005 based on estimates prepared by WHO staff (Moscow office). See country profile for further details.

Latest available data are for 2004.

Median value

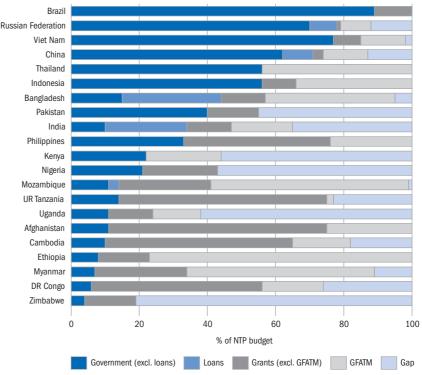
(US\$ 93 million, or 78% of the total gap). The shortfall in India is associated with the end of the existing World Bank credit (1998-2004), but the balance is expected to be made up through additional grants and a new World Bank credit to be negotiated in 2005. Gaps in the other four countries are linked to the development in 2003 and/or 2004 of much more ambitious plans to expand and improve TB control. Proportionally, the largest gaps are in Kenya, Nigeria, Uganda and Zimbabwe (all more than 50% of the total NTP budget; Figure 28). These gaps are large enough to seriously constrain progress in TB control in these countries.

Further details, including charts showing trends in NTP budgets by funding source and line item for each year 2002-2005, are provided in the country profiles (Annex 1).

### Total costs of TB control and funding in HBCs

NTP budgets include only part of the resources needed for TB control. In particular, they do not include the costs associated with general health services staff and infrastructure, which are used when TB patients are hospitalized or make outpatient clinic visits for directly observed treatment (DOT) and monitoring. For the 22 HBCs combined, the total costs of TB con-

FIGURE 28 Sources of funding for NTP budgets, 21 high-burden countries, a 2005

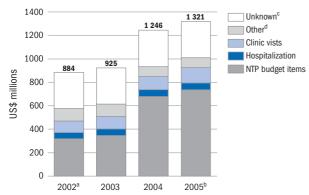


No data available for South Africa. Figures for Russian Federation based on estimates prepared by WHO staff (Moscow office). See country profile for further details

trol are projected to be US\$ 1.2 billion and US\$ 1.3 billion in 2004 and 2005, respectively, compared with actual costs of around US\$ 900 million in 2002 and 2003 (Figure 29, Figure 30, Table 16). These increases in projected costs are because of the large increases in planned NTP spending (described above) and because of the higher costs of clinic visits and hospitalization that are associated with treating more patients.

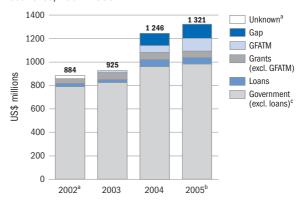
The largest costs are for the Russian Federation and South Africa,

FIGURE 29 Total TB control costs by line item, 22 high-burden countries, 2002-2005



- $^{\mathrm{a}}$  Costs assumed to be as for 2003 for Afghanistan, Bangladesh, Mozambique, Russian Federation, Uganda and Zimbabwe,
- Estimate for LIR Tanzania is based on 2004 data
- Total TB control costs for Thailand and South Africa could not be broken down as for other countries, so the total is presented as "Unknown". Estimates for South Africa are based on recent costing studies.
- "Other" includes costs for hospitalization and fluorography in the Russian Federation not reflected in the NTP budget.

FIGURE 30 Total TB control costs by funding source, 22 high-burden countries, 2002-2005



- <sup>a</sup> Costs assumed to be as for 2003 for Afghanistan, Bangladesh, Mozambique, Russian Federation, Uganda and Zimbabwe. "Unknown" applies to DR Congo and Nigeria, as breakdown of NTP budget by funding source not available.
- Estimate for UR Tanzania is based on 2004 data
- Estimates of total TB control costs (2002-2005) for South Africa are based on costing studies and all costs are assumed to be funded by the government

TABLE 16 Total TB control costs and available funding, high-burden countries, 2005

	TOTAL COST	CHANGE FROM 2002 <sup>a</sup>	CHANGE FROM			VAILABLE FUNDIN (US\$ MILLIONS)	IG		CHANGE IN		E FUNDING SINC	E 2002, <sup>b</sup> BY	SOURCE
	(US\$ MILLIONS)	(US\$ MILLIONS)	2002	GOVERNMENT (EXCL. LOANS)	LOANS	GRANTS (EXCL. GFATM)	GFATM	GAP	GOVERNMENT (EXCL. LOANS)	LOANS	GRANTS (EXCL. GFATM)	GFATM	GAP
1 India	89	28	45	48	11	6	8	15	5	-13	0.7	8	15
2 China	158	97	159	98	14	4	21	21	45	14	2	21	-22
3 Indonesia	50	28	125	31	0	4	15	0	20	0	1	15	-25
4 Nigeria	21	10	90	12	0	3	0	7	4	0	-2	0	-0.05
5 Bangladesh	27	17	169	9	6	3	8	1	2	6	-0.7	8	1
6 Pakistan	26	21	434	15	0	3	0	9	9	0	2	0	7
7 Ethiopia <sup>c</sup>	9	1	12	3	0	1	5	0	-1	0	-3	5	0
8 South Africad	300	-	-	300	-	-	-	-	-	-	-	-	-
9 Philippines	30	9	41	25	0	4	2	0	6	0	4	2	-4
10 Kenya	18	11	170	6	0	0	3	8	-0.7	0	-3	3	7
11 DR Congo	34	14	70	24	0	5	2	3	7	0	0	2	-0.8
12 Russian Federation <sup>e</sup>	399	154	63	303	25	2	30	39	66	25	-6	30	39
13 Viet Nam	28	3	13	25	0	0.9	2	0.3	0.2	-2	-0.1	2	0.3
14 UR Tanzania <sup>f</sup>	21	5	31	13	0	5	0.2	2	2	0	0.6	0.2	1
15 Brazil	46	9	25	44	0	2	0	0	7	0	2	0	0
16 Uganda	7	5	198	1	0	0.8	0.9	4	0.7	-1	0.3	0.9	0.7
17 Thailand	11	3	31	9	0	0	2	0	0.6	0	0	2	0
18 Mozambique	9	5	139	3	0.2	2	4	0.1	0.7	0.2	-0.5	4	-5
19 Zimbabwe <sup>e</sup>	15	9	158	5	0	2	0	9	0.4	0	0.04	0	9
20 Myanmar	6	4	232	1	0	1	3	0.6	0.3	0	1	3	-2
21 Afghanistan	3	-1	-34	0.3	0	2	0.7	0	0	0	0.5	0.7	-2
22 Cambodia	12	4	63	5	0	4	1	1	0.3	-0.7	3	1	0.0
ligh-burden countries	1321	437	70 <sup>g</sup>	982	56	55	109	119	175	28	3	109	20

which together account for US\$ 700 million of the total cost of US\$ 1.3 billion estimated for 2005. South Africa is a middle-income country, and the high costs are mainly explained by the higher prices for items such as hospitalization and outpatient visits, compared with those typical in lowincome countries. The Russian Federation staffs and runs an extensive network of TB hospitals for treatment, has a large budget for second-line drugs to treat many MDR-TB patients and still carries out mass population screening by fluorography. China and India have the third and fourth highest costs, estimated at US\$ 158 million and US\$ 89 million respectively in 2005. Seven additional countries have total costs of US\$ 25-50 million in 2005, three have costs of around US\$ 20 million and the rest have costs of US\$ 15 million or less.

The countries with the largest projected absolute increases in annual costs are the Russian Federation (US\$ 154 million since 2003) and China (US\$ 97 million since 2002). Increases of around US\$ 20-30 million since 2002 are estimated for Bangladesh, India, Indonesia and Pakistan. The changes for other HBCs are around or below US\$ 10 million. The biggest proportional increases are for Myanmar and Pakistan (both more than 200%), while increases are in the range 100–200% for seven additional countries.

Funding for the general health services staff and infrastructure used by TB patients during clinic visits and hospitalization is assumed to be funded by governments. This assumption, together with the implicit assumption that health systems have sufficient resources to support the treatment of growing numbers of patients in 2004 and 2005, means that the resources available for TB control are estimated to have increased from almost US\$ 900 million in 2002 to US\$ 1.2 billion in 2005 (Figure 30). The contribution by HBC governments to the total cost of TB control in 2005 is 79% on average, which is larger than their contribution to NTP budgets (Figure 31). This high average figure conceals important variation among countries; many HBCs are dependent on grants to cover more than one third

a TB control costs in 2002 were estimated using expenditure rather than budget data wherever possible. For countries that did not provide expenditure data for 2002 (Kenya and UR Tanzania), available funding was used as a proxy. Where neither budget nor expenditure data were available for 2002 (Afghanistan, Bangladesh, Mozambique, Russian Federation, Uganda and Zimbabwe), comparisons are with 2003.

b The sum of changes in available funding is different from the total change in TB control costs (column 3) when expenditures are lower than available funding. A further reason is that changes are calculated with respect to 2003 when a breakdown of funding was not available for 2002 (DR Congo and Nigeria).

Comparisons are with expenditure data for 2002.

No data were provided by the NTP; the cost per patient was estimated using recently published costing studies and multiplied by the number of patients notified in 2003 to give the estimated total cost.

Comparisons are with expenditure data for 2003. Figures for Russian Federation for 2005 based on estimates prepared by WHO (Moscow office). See country profile for further

f Latest available data are for 2004.

g Median value.

of the total costs of TB control, or to close large funding gaps. The share of the total costs provided by HBC governments is closely related to average income levels (Figure 32), although Viet Nam stands out as a low-income country with a very high government contribution (90%).

For all HBCs, the estimated gap between the funding already available and the total cost of TB control is US\$ 119 million in 2005, i.e. the NTP budget gap reported above. Further details, including charts that show trends in total TB control costs by line item for each year 2002-2005, are provided in the country profiles.

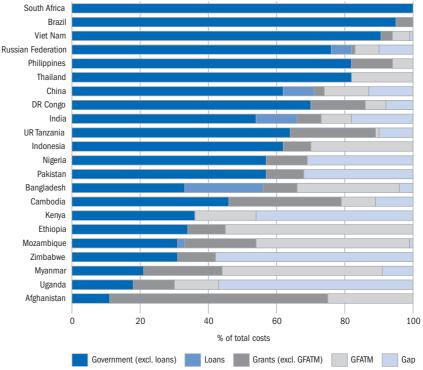
### Per patient costs and budgets

There is much variation among countries in budgets and costs per patient (Table 17). The budgets for first-line drugs are lowest in India, Myanmar and the Philippines (US\$ 11-17 per patient). In most countries, the budget is in the range US\$ 20-35, but higher in Bangladesh, Brazil, Indonesia, Mozambique and the Russian Federation. Higher budgets in Bangladesh and Mozambique are explained by the creation of large buffer stocks, which distort the average value in 2005. In Indonesia, the drug budget has increased to allow use of fixed-dose combinations (FDCs).

The budget per patient, including all line items, is lowest in India, at US\$ 34. The budget is also relatively low in Ethiopia and the Philippines (both around US\$ 50) and in Myanmar (US\$ 68). Most other countries (n =13) have budgets in the range US\$ 100-200 per patient. The only low-income country with a budget above US\$ 200 per patient is Mozambique. The Russian Federation has by far the highest budget per patient, for reasons explained above (the figure for South Africa may also be high, but no data are available).

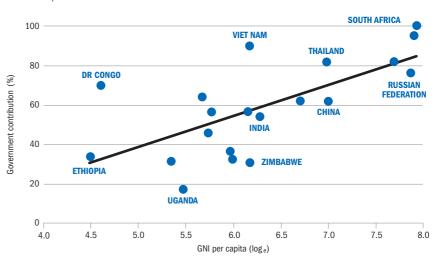
The total cost per patient treated in 2005 is lowest in India (US\$ 66), below US\$ 100 in Ethiopia and Myanmar, and below US\$ 150 in Bangladesh and Uganda. It is in the range US\$ 150-300 in 11 countries,26 and

FIGURE 31 Sources of funding for total TB control costs, 22 high-burden countries, a 2005



Figures for Russian Federation based on estimates prepared by WHO staff (Moscow office) and recent costing studies. See country profile for further details

FIGURE 32 Government contribution to total TB control costs by GNI per capita, 20 high-burden countries,<sup>a</sup> 2005



<sup>&</sup>lt;sup>a</sup> No information on GNI per capita available for Afghanistan or Myanmar. Figure for Russian Federation based on estimates prepared by WHO staff (Moscow office) and recent costing studies. See country profile for further details

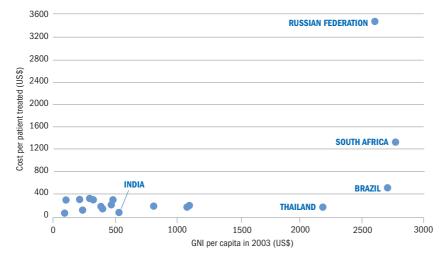
<sup>&</sup>lt;sup>26</sup>This assumes that if data for Afghanistan were available, the cost would be in this range.

TABLE 17 Total costs and NTP budgets per patient, high-burden countries, 2005

	2005 (US\$)			CHANGES FROM 2002 (FACTOR <sup>a</sup> )				
	FIRST-LINE DRUGS BUDGET	NTP BUDGET	TOTAL COST	FIRST-LINE DRUGS BUDGET	NTP BUDGET <sup>b</sup>	TOTAL COST		
1 India	11	34	66	1.1	1.0	1.1		
2 China	20	188	188	1.2	1.4	1.4		
3 Indonesia	47	155	182	1.5	1.3	1.3		
4 Nigeria	29	159	291	0.5	1.2 °	1.0		
5 Bangladesh <sup>c</sup>	45	116	146	2.2	1.5	1.3		
6 Pakistan	23	153	213	0.4	3.3	2.3		
7 Ethiopia	34	49	68	1.3	1.1	0.9		
8 South Africa	-	-	1320 <sup>d</sup>	-	-	-		
9 Philippines	17	48	174	0.7	0.9	1.0		
10 Kenya	31	142	173	0.8	2.7	2.1		
11 DR Congo	14	94	297	0.4	1.0	1.0		
12 Russian Federa	ation <sup>e</sup> 70	2748	3472	1.0	2.5 ℃	1.8		
13 Viet Nam	31	129	299	0.9	1.1	1.2		
14 UR Tanzania <sup>f</sup>	21	133	320	0.5	1.6	1.2		
15 Brazil	47	239	516	1.0	1.4	1.1		
16 Uganda <sup>c</sup>	20	111	120	0.4	2.3	2.1		
17 Thailand	25	72 <sup>g</sup>	173	0.3	0.6	1.0		
18 Mozambique <sup>c</sup>	75	241	311	3.4	3.3	2.2		
19 Zimbabwe <sup>c</sup>	35	192	266	1.1	5.8	2.4		
20 Myanmar	13	68	80	0.7	3.2	2.5		
21 Afghanistan <sup>c</sup>	_	104	_	_	0.3	_		
22 Cambodia	27	186	311	0.7	1.4	1.1		
High-burden coun (median value)	tries 28	133	213	0.9	1.4	1.2		

- Indicates not available.
- Calculated as 2005 value divided by 2002 value.
- Calculated as NTP budget per patient projected in 2005 divided by NTP expenditure per patient notified in 2002. Available funding was used as a proxy for expenditures for India, Kenya, Thailand, Uganda, UR Tanzania and Viet Nam.
- Comparisons are with 2003 data.
- Estimate is based on notifications for 2003, as no projections for 2005 available.
- Figures for 2005 based on estimates prepared by WHO staff (Moscow office). See country profile for further details. Comparison for first-line drugs is with 2004.
- Latest available data are for 2004. Thus comparison is of 2004 with 2002.
- g This figure is artificially low because it is based on the central level budget only (see country profile).

FIGURE 33 Cost per patient treated by GNI per capita, 20 high-burden countries, 2005



<sup>&</sup>lt;sup>a</sup> No information on GNI per capita available for Afghanistan or Myanmar. Figure for Russian Federation based on estimates prepared by WHO staff (Moscow office) and recent costing studies. See country profile for further details.

slightly more than US\$ 300 in three countries. There are three countries with much higher costs: Brazil, the Russian Federation and South Africa. Their higher costs are not surprising given their middle-income status and associated higher prices for inputs such as staff (Figure 33), as well as the extensive use of hospitalization in the Russian Federation.

Budgets and costs are generally stable (notably during a period of rapid DOTS expansion in India) or increasing.

Further details, including charts that show five per patient indicators (costs, budgets, available funding, expenditures and first-line drugs budget) for each year 2002-2005, are provided in the country profiles (Annex 1).

### Expenditures in comparison with budgets and available funding

For countries that have received large increases in funding, the challenge now is to spend the extra money, and to translate extra spending into improved case detection and treatment success rates. The ability to spend available money can be assessed by comparing expenditures with available funding and budgets (Table 18). Complete sets of data on budgets, funds and expenditures are available for 15 HBCs in 2003 (the most recent year for which expenditure data are currently available). Expenditures were generally less than available funding. In India, costs proved to be lower than anticipated, and all planned activities were implemented. The capacity of the Tanzanian NTP to spend available money is discussed in Annex 1. For other countries, more work is needed to understand the reasons why expenditures are lower than available funding. The findings will have implications for programmes that are now benefiting from large influxes of new money. It is too early to say if large increases in spending can be translated into improved programme performance.

### Budgets, funds and targets

Countries can be categorized according to whether the number of patients to be treated is consistent with meeting the 2005 targets, treatment success rates, the extent to which the

budget for the projected number of patients is funded, how the budget per patient has changed through time and whether there is evidence that the additional funding can be effectively absorbed (Table 19). India, Myanmar, the Philippines and Viet Nam are in the best financial position to reach the targets (or to maintain the programme at target levels in the case of Viet Nam). Cambodia and China are well placed to do so if they can make up the remaining funding shortfalls. Indonesia appears to have the funding required to achieve targets, and Bangladesh may come close. However, it is unclear how many more cases will actually be detected and successfully treated as a result of the additional funds now available in these two countries. For the remaining 14 HBCs, the planned programmes of treatment are less than required to meet the targets for case detection (11 countries) and/ or it is not clear if they are sufficient to meet the target for treatment success, although five of these countries report no or negligible shortfalls in funding.

### GFATM contribution to TB control

High-burden countries. The GFATM is the single most important source of grant funding for HBCs, and several countries are relying on the GFATM to fund more than one third of their budgets. After four rounds of proposals, the total value of approved proposals (which, with four exceptions, cover five years) is US\$ 818 million (Table 20). The amounts included in the two-year grant agreements<sup>27</sup> total US\$ 218 million, and are sometimes lower than the amounts in years 1 and 2 of the original proposals (75% for those proposals for which both the original request for years 1 and 2 and the grant agreement amount are available; the biggest discrepancy is for Indonesia).

By the end of 2004, US\$ 116 million had been disbursed. For each country, we can compare the actual and expected rates of disbursal, where the expected rate assumes that disbursements should be spread evenly

TABLE 18 Budgets, available funding and expenditures (US\$ millions), 15 high-burden countries, 2003

	BUDGET	AVAILABLE FUNDING	EXPENDITURES	
1 India	42	42	25	
2 China	95	87	80	
3 Indonesia	32	29	21	
4 Nigeria	13	6	6	
5 Bangladesh	7	7	7	
6 Pakistan	6	6	3	
7 Ethiopia	11	11	8	
8 Philippines	7	6	4	
9 DR Congo	10	7	5	
10 Viet Nam	11	11	11	
11 UR Tanzania	5	5	4	
12 Brazil	16	16	14	
13 Mozambique	8	3	2	
14 Myanmar	5	1	1	
15 Cambodia	6	3	2	
Total	273	239	193	

Categorization of high-burden countries according to financial criteria, 2005

CATEGORY	CRITERIA	COUNTRIES
I	Projected number of cases to be treated in 2005 sufficient to meet 70% case detection target  Treatment success rate achieved or close to being achieved for 2002 cohort  Budget per patient treated stable or increasing  No or minor funding gap, or funding gap likely to be filled  Demonstrated ability to absorb any additional funds required to achieve targets	India Myanmar Philippines Viet Nam
lla	As for I, except that funding gap needs to be filled	Cambodia China
IIb	As for I, except that ability to absorb large increases in funding and translate them into improved detection and treatment success rates is currently unproven	Indonesia
llc	As for Ilb, but projected number of cases treated only within 10% of case detection target	Bangladesh
III	Projected cases not in line with case detection target, and/or unclear if treatment success target can be achieved, but no (or negligible) reported funding gap	Brazil Ethiopia Mozambique South Africa Thailand
IV	Projected cases not in line with case detection target (all except DR Congo) and/or treatment success, and large funding gap	Afghanistan DR Congo Kenya Nigeria Pakistan Russian Federation Uganda UR Tanzania Zimbabwe

<sup>&</sup>lt;sup>27</sup> Signature of grant agreements is needed before any disbursements can take place.

TABLE 20 GFATM financing, high-burden countries, as of end 2004

		TOTAL BUDGET APPROVED <sup>a</sup>	TOTAL YEAR 1 AND 2 BUDGETS (AS IN PROPOSALS)	GRANT AGREEMENT	TOTAL DISBURSEMENTS BY END 2004 (AS OF 20-DEC-2004)	OF GRANT	2004 AS % AGREEMENT EXPECTED		DATE GRANT	TIME BETWEEN BOARI APPROVAL AND SIGNATURE OF GRANT	AGREEMENT SIGNATURE AND FIRST
	GRANT	(US\$ MILLIONS)	(US\$ MILLIONS)	(US\$ MILLIONS)	(US\$ MILLIONS)	RATE OF DISBURSAL	RATE OF DISBURSAL	DATE BOARD APPROVAL	AGREEMENT SIGNATURE	AGREEMENT (MONTHS)	(MONTHS)
1 India	G1 <sup>b</sup>	9	6	6	4	76	96	22-Apr-02	30-Jan-03	9	6
	G2°	29	13	7	2	27	42	13-Jan-03	12-Feb-04	13	2
	G3 <sup>c,d</sup>	15	3	3	0	0	8	15-0ct-03	15-0ct-04	12	2+
	G4	27	7	Not signed yet				28-Jun-04		6+	
2 China	G1	48	25	25	25	100	96	22-Apr-02	30-Jan-03	9	2
	G2	56	28	Not signed yet				28-Jun-04		6+	
3 Indonesia		70.7	55	22	16	73	96	22-Apr-02	27-Jan-03	9	2
4 Bangladesh	G1°	42	17	11	5	41	21	15-0ct-03	07-Jul-04	9	0.6
	G2°	18	8	5	2	44	13	15-0ct-03	24-Aug-04	10	0.2
5 Pakistan	G1°	4	2	2	0.7	31	67	13-Jan-03	06-Aug-03	7	4
	G2°	13	7	6	2	33	8	15-0ct-03	12-0ct-04	12	1
6 Ethiopia <sup>c</sup>		21	_	11	7	59	88	22-Apr-02	18-Mar-03	11	5
7 South Africa	G1 <sup>d</sup>	70	14	2	2	100	67	22-Apr-02	08-Aug-03	16	4
	G2c,d	_	_	12	9	76	67	22-Apr-02	08-Aug-03	16	4
	G3c,d	72	27	27	13	48	67	22-Apr-02	08-Aug-03	16	4
	G4 <sup>d</sup>	25	8	Not signed yet				13-Jan-03		23+	
8 Philippines		11	3	3	3	77	75	13-Jan-03	11-Jun-03	5	0.6
9 Kenya		11	5	5	2	50	75	13-Jan-03	23-Jun-03	5	2
10 DR Congo <sup>b, c</sup>		8	6	6	6	90	75	13-Jan-03	18-Jun-03	5	1
11 Russian Federation	G1(Tomsk) <sup>c</sup>	11	6	6	2	28	8	15-0ct-03	14-0ct-04	12	2
	G2	92	54	Not signed yet				28-Jun-04		6+	
12 Viet Nam <sup>c</sup>		10	3	3	0.4	16	58	13-Jan-03	15-0ct-03	9	5
13 UR Tanzania	G1 <sup>c,d</sup>	88	25	24	7	30	13	15-0ct-03	06-Sep-04	11	2
	G2 (Zanziba	r) <sup>c</sup> 2	1	1	0.7	70	95	13-Jan-03	07-Sep-04	20	2
14 Uganda <sup>c</sup>		6	7	5	1	25	38	13-Jan-03	15-Mar-04	14	1
15 Thailand		13	7	7	3	45	79	22-Apr-02	18-May-03	13	2
16 Mozambique <sup>c</sup>		18	12	9	0	0	38	13-Jan-03	02-Apr-04	15	9+
17 Myanmar <sup>c</sup>		17	7	7	2	34	17	13-Jan-03	13-Aug-04	19	1
18 Afghanistan <sup>b</sup>		3	2	Not signed yet				28-Jun-04		6+	
19 Cambodia		7	3	3	1	45	58	13-Jan-03	14-0ct-03	9	2
Total		818	361	218	116	45°	67°			11e	<b>2</b> e

<sup>-</sup> Indicates not available.

over the two years following the date on which the agreement is signed (Table 20, column 7). China is the only country where all funds in the two-year grant agreement have been disbursed within the expected period of two years. For eight countries and 10 grants, disbursements are better than expected.28 For five countries, disbursements are within 20% of the expected value, and for nine countries disbursements are around 25% or more below the expected value. One example is Ethiopia, a country that is largely dependent on GFATM funds. Another is Mozambique, which is also highly dependent on the GFATM, but which has to date received no funds at all, even though the grant agreement was signed in April 2004. The GFATM web site notes that initial disbursements are often small, given the need for strengthening programme capacity and preparation of procurement plans.29 Furthermore, low disbursement rates appear to be associated with the principal recipient; for eight countries and nine grants where the disbursements are below the expected value, the GFATM web site notes that assessments of the principal recipient are pending.

The initial delay in disbursement is caused mainly by the time taken to sign the grant agreement after proposal approval. Once grant agreements are signed, disbursements are usually made within 2 months (the exception is currently Mozambique), compared with delays of between 5 and 23 (median value 11) months between grant approval and signature.

Other countries. After four rounds of proposals, 60 non-HBCs have ap-

Total budget requested is for five years, unless otherwise stated.

Total budget requested is for three years.

<sup>&</sup>lt;sup>c</sup> Assessment of principal recipient pending.

TB/HIV grant.

Median value.

<sup>&</sup>lt;sup>28</sup> However, the figure for Myanmar is misleading, because the NTP is a sub-recipient that had not received any funds by the end of 2004.

<sup>&</sup>lt;sup>29</sup> http://www.theglobalfund.org/en/ funds\_raised/commitments/.

proved proposals with a total value of US\$ 400 million. The amounts included in the two-year grant agreements<sup>30</sup> total US\$ 153 million, of which US\$ 65 million had been disbursed by the end of 2004. Disbursements are generally similar or higher than expected values, except in the European and South-East Asia Regions. A summary table with the same indicators as those shown for the HBCs is available upon request.

The regional distribution of GFATM grants for HBCs and other countries is shown in Figure 34.

### NTP budgets by WHO region, HBCs and other countries

NTP budgets and sources of funding by WHO region in 2005 are shown for both HBCs and non-HBCs in Figure 35, based on the 55 countries that submitted data of sufficient quality. Total budgets and sources of funding are dominated by the HBCs in the South-East Asia Region and the Western Pacific Region, because the HBCs account for almost all TB cases in these regions. While non-HBCs account for a large share of cases in the European Region, Eastern Mediterranean Region and Region of the Americas, we received insufficient data to make an assessment of total budgets and funding sources, or to make any useful comparisons between HBCs and non-HBCs. For the African Region, we had budget data for countries that account for 79% of TB cases.31 Non-HBCs add substantially to the HBC budget totals (US\$ 128 million versus US\$ 77 million for HBCs alone). Proportional to budgets, funding gaps are smaller in non-HBCs in the African Region, with relatively higher funding contributions from the GFATM and governments.

FIGURE 34 Regional distribution of GFATM lifetime budgets, as of end 2004

Total TB and TB/HIV: US\$ 1.2 billion Total TB/HIV: US\$ 296 million

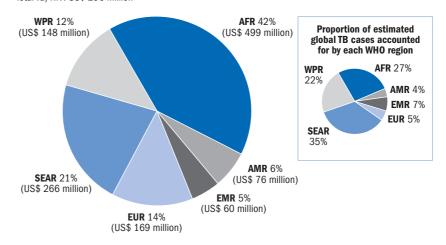
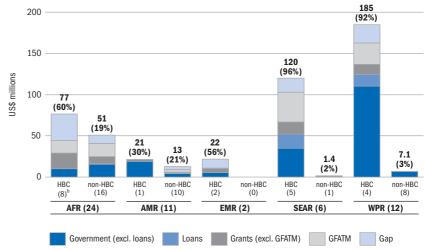


FIGURE 35

### Regional distribution of total NTP budgets by source of funding, 20 high-burden and 35 non high-burden countries, 2005

Figures in parentheses show the percentage of all estimated global TB cases in the region accounted for by the countries included in the bar.



The European Region is excluded because the much higher budget in the Russian Federation makes it difficult to illustrate patterns in other regions. Complete data were received from three countries other than the Russian Federation: Estonia, Latvia and Republic of Moldova. For these three countries funds come entirely from the government with the exception of a GFATM grant for the Republic of Moldova.

Excludes South Africa

<sup>30</sup> Signature of grant agreements is needed before any disbursements can take place.

<sup>31</sup> If data for South Africa were available, the figure would be 89%.

## Discussion

### **Progress towards the Millennium Development Goals**

Within the framework of the United Nations MDGs, TB control is guided by five principal indicators, two that quantify DOTS implementation (case detection, treatment success) and three that could measure the impact of DOTS on the epidemic (incidence, prevalence, mortality). The MDG framework is a stimulus to think beyond the 2005 targets for DOTS implementation, and to consider the benefits of TB control up to and beyond 2015. Long-term thinking underpins sustainable TB control, and is vital for planning the trajectory towards TB elimination.

This is the first in this series of annual reports to evaluate changes in incidence, prevalence and deaths from 1990, the MDG reference year, through to 2003. Based primarily on trends in case notifications, the TB incidence rate was, by 2003, falling or stable in seven out of the nine regions of the world defined in Figure 6. Incidence rates in eastern Europe (mostly countries of the former Soviet Union) and Africa (countries with low and high HIV rates) increased during the 1990s, but appear to have peaked in Europe around 2001, and have since fallen. There is no persuasive method of predicting when peak incidence rates will be reached, and at what levels, in Africa, but the rates of increase slowed markedly during the 1990s.

Because these adverse regional effects are diminishing, the rate of increase in global incidence is also slowing, after growing most rapidly in the mid 1990s. The global incidence rate reached 140 per 100 000 population in 2003 (8.8 million new cases, including those who are HIV-positive), but was still increasing at 1.0% annually. This assessment is, however, dependent on the assumptions we have made about trends in HBCs. For example, the series of case notifications for India suggest that the incidence rate is falling, but the preferred assumption, until further evidence be-

comes available, is that incidence is stable (this is conservative with respect to case detection and the impact of DOTS). If the incidence rate is actually falling in India at 2.4% per year, as indicated by case notifications in 1992-2003, then the global incidence rate would also be falling, al-

The trend in global TB incidence has been little affected, so far, by DOTS programmes. Chemotherapy is more likely to have reduced TB prevalence and deaths, but neither of these indicators is measured routinely in HBCs. The calculations presented here, which are derived from estimates of incidence, duration and case fatality, suggest that the global prevalence rate fell from 309 to 245 per 100 000 between 1990 and 2003 (including HIV-positive TB patients), and was falling at 5% per year in 2003. The TB death rate (including deaths among HIV-positive TB patients) was also falling in 2003, but more slowly at 2.5% per year. Prevalence and deaths, like incidence, have been rising in Africa, and most steeply in African countries with the highest rates of HIV infection. In our assessment, incidence, prevalence and death rates are falling or stable in five out of the six WHO regions, and in seven of the nine regions of the world shown in Figure 6.

Treatment success in the 2002 cohort was reported to be 82% of 1.4 million registered cases, close to the 85% target, but no higher than in the previous two annual cohorts. The overall rate of treatment success is strongly influenced by data from the three countries that have the largest numbers of new cases annually -China, India and Indonesia. All three submitted data indicating that the 85% target had been exceeded in 2002. These are impressive results, achieved while treating hundreds of thousands of patients, but success rates reported to be higher than 90% (e.g. China) need to be kept under review.

Of greater concern are the low cure

rates in the European Region and the African Region in 2002. The European Region reported the highest rate of treatment failure, probably linked to the high levels of drug resistance in countries of the former Soviet Union. It also reported the second highest death rate on treatment, which is likely to be associated both with drug resistance and with the high proportion of elderly patients in Western Europe. The African Region reported the highest death rates in TB patients, undoubtedly associated with HIV coinfection. But the success rate of African DOTS programmes was also low because they lost 19% of patients through default, transfer between treatment centres or by failing to record any outcome of treatment (patients "not evaluated"). Such losses to follow-up were also high in the Region of the Americas, Eastern Mediterranean Region and European

The target for treatment success under DOTS refers only to new smearpositive cases, but information about patients presenting for re-treatment, including the outcomes of treatment for these patients, is also indicative of programme performance. The WHO data collection form for the 2002 cohort asked DOTS programmes to distinguish between re-treatment after relapse, default and failure, both for cases reported in 2003 and for patients undergoing re-treatment during 2002. Although it is probable that the case definitions (Table 2) are not strictly observed (and many re-treated patients were not classified in data submitted to WHO), some of the findings deserve comment. First, the comparative success of treatment for different classes of patients was consistent with expectations: lower on average for re-treated than for new cases; and among re-treated patients, higher for relapses, intermediate for defaulters and lowest for failures. Moreover, patients who defaulted during their first course of treatment tended to default from a second or subsequent course of treatment.

Third, the regional distribution of adverse re-treatment outcomes resembled the pattern observed for new cases: African countries reported high death rates and many patients were lost to follow-up; European countries reported high rates of death and treatment failure. The accuracy of reporting needs to be verified, but these data should help to identify TB patients who, for example, are less likely to comply with treatment (persistent defaulters) and those who are more likely to be infected with HIV (especially in Africa), or who are carrying drug-resistant bacilli (especially in eastern Europe).

Although DOTS programmes have diagnosed and treated more than 17 million cases since 1995, the global DOTS case detection rate was still only 45% in 2003, well below the 70% target. However, the detection rate increased by 8% between 2002 and 2003, faster than at any time since recording began in 1995. If detection continues to increase at this rate, the estimated global case detection rate will be approximately 60% by 2005.

While the acceleration in case-finding during 2003 exceeded expectations, most of the additional patients (63%) were reported by just two countries: China and India. If this pace of expansion is to be maintained or accelerated, other HBCs must contribute more. Approximately 1.8 million smear-positive cases were notified by DOTS programmes in 2003. According to our estimates, another 1.4 million new patients, undetected by DOTS programmes in 2003, were living in just eight HBCs, including China and India. Together, these 3.2 million patients account for more than 70% of new cases arising in 2003. Therefore, intensive case-finding in these countries would contribute greatly to meeting the global target of 70% case detection.

In some regions of the world, large numbers of patients are reported from outside DOTS areas. The 70% target could be reached in the Region of the Americas by ensuring that more than 43 000 smear-positive patients currently reported by non-DOTS programmes are diagnosed and treated under DOTS, the majority in Brazil. In

the European Region, the Russian Federation reported more than 100 000 patients from the 75% of the country not yet covered by DOTS, 23 000 of which were new smearpositive patients. China and India reported an additional 70 000 new smear-positive patients from non-DOTS areas in 2003.

By contrast, DOTS programmes in other HBCs including Bangladesh, Ethiopia, Indonesia, Nigeria and Pakistan will have to recruit patients that are not yet seen and reported by public health surveillance systems. These unreported patients undoubtedly exist because they are found, for example, during population-based prevalence surveys. Some never receive TB treatment; some are treated in public and private clinics and hospitals that are not linked to ministries of health. To ensure that these patients have access to DOTS services, TB control programmes will need to embark on new activities and establish new collaborations, many of which will be specific to the structure of local health services.

Considering both of the targets for DOTS implementation, Viet Nam was still the only member of the current group of HBCs to have reached 70% case detection and 85% treatment success by 2003. However, Cambodia, Myanmar and the Philippines were all close to achieving the targets. Although these Asian countries have different problems to solve, they should, with China and India, be able to meet the targets by 2005.

Smear-positive patients are the focus of the DOTS strategy, but many DOTS programmes also routinely treat smear-negative patients, with pulmonary or extrapulmonary disease. For the countries that report smearnegative patients, the numbers may be less accurate than for smearpositive disease because diagnosis is more difficult. In this context, the remarkable differences between regions in the proportions of patients reported with extrapulmonary disease need further investigation. The exceptionally high extrapulmonary case-load in the Eastern Mediterranean Region (20-30%) might be due to over-diagnosis, but it might also be a real and unexplained epidemiological phenomenon.

The establishment of the MDGs presents a challenge, not just for the implementation of DOTS and other means of TB control, but also for the measurement of epidemiological impact. Ideally, all countries would count new cases and deaths via a comprehensive routine system of surveillance and vital registration, and estimate the prevalence of disease and infection by population-based surveys. In reality, countries will have to select some methods of measurement in preference to others. Some guidance on the advantages and disadvantages of different epidemiological measurements is given in Table 21.

### **Planning and DOTS** implementation

All HBCs have a strategic plan for DOTS expansion and, during 2005, many will begin a new planning cycle for the next five years. However, the transition from planning to implementation, and then to the improvement of coverage, case detection and treatment success has been slower than anticipated in several countries. The success of some NTPs in raising funds for TB control (and particularly from the GFATM) has not been followed by productive spending.

Among the obstacles to DOTS expansion, five are of overriding importance: shortages of trained staff, lack of political commitment, weak laboratory services, and the inadequate management of MDR-TB and of TB in people infected with HIV.

The acute shortage of adequately trained staff affects the distribution and quality of services. This workforce crisis is felt particularly in the underperformance of central management, and through failings in the laboratory network. To remedy the problem, the HBCs need, at the very least, strong and clear policies for recruiting, retaining and motivating staff. One way to secure political commitment to solve this and other problems is by strengthening national and international partnerships. A consistent message about the importance of TB control, delivered from various constituencies, is a basis for effective advocacy and communication.

### TABLE 21 Advantages and disadvantages of various epidemiological measurements for TB control

Text in blue refers to attributes of the indicator; regular text refers to attributes of the measurement technique.

MEASURE	ADVANTAGES	DISADVANTAGES
Prevalence of infection	Risk of infection changes relatively quickly in response to control (but prevalence, from which risk is calculated, changes slowly).	Measures infection, not disease burden; not an MDG indicator.
From tuberculin surveys	Relatively cheap and logistically straightforward.	Results often hard to interpret where infection rates are low and where BCG coverage is high or where exposure to environmental mycobacteria is high; measures average risk of infection over past 5–10 years; Stýblo 1:50 rule for indirectly estimating disease incidence may not be applicable under chemotherapy, or where HIV infection rates are high.
Prevalence of disease	Component due to duration of illness changes relatively quickly in response to control; MDG indicator.	Component due to incidence changes slowly in response to control.
From population-based surveys	Accurate measure of bacteriologically confirmed disease; should change quickly in response to control; surveys useful where routine surveillance data are poor, and are a platform for related investigations e.g. of interactions between patients and health system.	Costly; logistically complex (especially with radiography), therefore cannot be measured annually; does not easily lead to an estimate of TB incidence (denominator of WHO case detection rate), because duration is hard to assess.
Incidence of disease	Direct measure of denominator of WHO case detection rate; MDG indicator.	Changes slowly following reductions in transmission.
From case notifications	Direct measure of incidence; absolute incidence can be assessed from routine case reports where case detection judged to be high; trends can be judged from series of routine case reports, if measured consistently; every country now has a surveillance system, reporting annually or sub-annually.	Case detection mostly low in high-burden countries (underestimates incidence), and may vary through time (inaccurate trends).
From consecutive prevalence surveys	Direct measure of incidence.	Costly; logistically complex; requires ≥2 surveys with carefully judged survey interval and follow-up of individual patients.
TB mortality	Direct measure of TB burden accounting for a high proportion of DALYs; case fatality falls quickly in a new control programme; MDG indicator.	Component due to incidence changes slowly in response to control; hard to reduce case fatality further in low-burden countries.
From observations on patient cohorts	Direct observation of number of patients dying.	Deaths observed are those in cohort only, not in the population at large, and not beyond the period of cohort follow-up; deaths among defaulters and transfers usually unknown; TB not always the cause of death for patients on TB treatment.
From product of incidence and case-fatality rate	Simple and widely applicable.	Relies on accurate measures of incidence (above) and case fatality; case fatality measurable in observed DOTS cohorts, but not among patients treated elsewhere or untreated. Approximate at best.
From vital (death) registrations (VR)	Direct measure of TB deaths and trends; can be reported annually or sub-annually.	VR does not yet exist in many high-burden countries (notably in Africa and Asia); typically underestimates TB deaths; sensitivity and specificity untested.
From verbal autopsy (VA)	Review of registered deaths can improve accuracy of cause of death statistics.	Sensitivity and specificity of VA not fully evaluated; where no death registration system exists, laborious to compile deaths from a rare disease, and requires large sample sizes.

Besides the staff shortages, many laboratories participating in DOTS programmes have insufficient equipment and supplies, and limited procedures for quality assurance. All these essential elements need to be in place before laboratories take on the larger tasks of culturing M.

tuberculosis and testing for drug sensitivity, as will be required to integrate DOTS-Plus projects within DOTS programmes. To help improve capacity in HBCs, the DEWG has established a subgroup concerned with laboratory strengthening.

In addition to the deficiencies in

laboratories, the lack of national policies on MDR-TB management, the widespread availability of drugs of uncertain quality and the large numbers of MDR-TB patients treated outside the NTP together suggest that the treatment of drug-resistant TB is often inadequate. The high proportions of re-treatment cases reported by NTPs are also a signal that drugresistant forms of TB could be common in some populations where no surveys have yet been done. There are several remedies. WHO is in the process of expanding drug resistance surveillance and DOTS-Plus components within the context of regular TB control programmes. WHO is also working to establish a long-term competitive market for quality-assured drugs by leading a project to pre-qualify second-line drugs worldwide. GFATM grants are also being used to stimulate demand for drugs from reliable manufacturers. The Fund has selected the GLC as the mechanism for second-line drug procurement, and for monitoring approved projects.

The management of drug-resistant TB will be aided by a better understanding of the scale and distribution of the problem. Surveillance of drug resistance must be expanded to the five HBCs for which no data are yet available,8 and to other countries suspected to have high prevalence rates of MDR-TB. Information about new TB patients will be supplemented by data on patients presenting for re-treatment, including the systematic notification of all categories of re-treatment cases, the reporting of treatment outcomes and representative drug resistance surveys.

During 2003, very few TB patients had access to VCT and to ART. The numbers that actually have access to these services are probably somewhat higher than reported, but cannot be accurately known until TB/HIV monitoring systems are substantially improved. HIV/AIDS programme staff are increasingly aware of the fact that people infected with HIV are at high risk of developing active TB, while their counterparts in TB control programmes are seeing the impact of HIV on TB case-load, and on death rates in cohorts of TB patients on treatment. There has, until now, been little collaboration between TB and HIV/AIDS control programmes, but many such programmes are beginning to adopt elements of the WHO interim policy on collaborative TB/HIV activities.15 Even with the imperfect data presented in this report, it is clear that much closer collaborations of this kind are needed to develop and improve access to prevention, treatment and support services, for both TB and HIV/ AIDS patients.

Notwithstanding these weaknesses, this report has also identified a series of positive developments in DOTS implementation. The contributions to TB control of NGOs and community groups are clear expressions of the growing commitment of civil society. The work of these groups puts patients at the centre of the DOTS strategy, and improves access to TB services in remote areas and among disadvantaged and marginalized populations. NGOs are increasingly recognized as essential members of national partnerships for TB control. This recognition is helping not only to coordinate routine activities but also to develop a collaborative approach to solving the problems faced by NTPs. Some African countries are planning to involve community groups in collaborative TB/HIV activities. PPM projects are showing a measurable impact on case detection in several Asian countries, and may prove to be a mechanism for expanding TB control services in African cities.

With the significant influx of resources for TB control (from the GFATM, banks and bilaterals), especially to HBCs, some additional, catalytic funding is needed to ensure that NTPs have the technical capacity to make the best use of the new grants and loans. To satisfy this need, Stop TB partners launched a new initiative in 2003 - ISAC - an extraordinary effort to push towards the 2005 targets in selected countries, including China, India and Indonesia. The technical work under way aims to facilitate the access of patients to DOTS services, for example by expanding the geographical coverage of DOTS, by involving a greater diversity of public and private health-care providers, by strengthening in-country advocacy and social mobilization, and through partnership building and collaborative TB/ HIV activities.

### **Financing DOTS expansion**

There has been a big increase in NTP budgets and a big improvement in the funding available for TB control since 2002, with particularly large increases between 2003 and 2004. The total reported NTP budgets for the 22 HBCs in 2005 are US\$ 741 million, of which US\$ 622 million is available and US\$ 119 million is a funding gap. The total estimated costs of TB control32 are projected to be US\$ 1.3 billion, of which US\$ 1.2 billion is already available. With the exception of large additional government contributions in China, Indonesia and the Russian Federation, almost all of the extra funding for TB control since 2002 is from GFATM grants. The GFATM now plays a major role in the financing of TB control, contributing more than one third of the budget in several HBCs, and over half in a few.

As usual, the summary statistics conceal important variations among countries. Our analyses suggest that in 2005, the HBCs fall into four categories. In the first are four countries (India, Myanmar, the Philippines and Viet Nam) that have budgets consistent with reaching the 2005 targets, and which are likely to have minimal or no funding shortfall. India has continued to expand rapidly with fullyfunded budgets over the period 2002-2004, which, in 2003, provided more than enough money for planned activities. The Indian Revised National TB Control Programme has also maintained a constant budget per patient treated during the rapid expansion of DOTS. In the second are four countries that are close to being in this group, but which need to make up funding shortfalls (China, Cambodia), or where it is unclear how many more cases will actually be detected and successfully treated as a result of the substantial additional funds now available (Bangladesh, Indonesia). China stands out as having developed much larger budgets for 2004 and 2005 compared with previous years, for mobilizing a substantial increase in domestic and external financing to fund these budgets and for being the first HBC to secure full disbursement of a

<sup>32</sup> i.e. NTP budgets plus the cost of hospitalization and outpatient clinic visits of TB patients that are usually not included in NTP budgets.

two-year GFATM grant. In the third group are five countries that report no, or negligible, funding gaps for 2005, but whose plans are not sufficient to reach the targets for case detection (e.g. Ethiopia) or there are doubts about whether existing plans will ensure achievement of the treatment success target (e.g. South Africa). The nine countries in the final group need special attention because they report large funding gaps and, in addition, do not expect to treat enough patients to reach the case detection target (eight countries) and/or there are doubts about whether they can reach the treatment success target. Among these nine countries, Nigeria and Zimbabwe are the only low-income HBCs not to have secured GFATM funding to date. Pakistan's funding shortfall is a consequence, in large part, of planning for accelerated DOTS expansion in 2005.

The funding gap of US\$ 119 million identified by all NTPs for 2005 is higher than reported in 2003 and 2004, but may still be an underestimate. The budget gap is the difference between the funds needed to carry out planned activities and the funds actually available. If the activities planned by NTPs for 2005 are a realistic assessment of what can achieved, the budget gaps reported are arguably an accurate reflection of the funding gap. However, the activities required to meet the 2005 case detection target are greater than planned in 12 countries, and while Brazil and South Africa may already detect more than 70% of all TB cases it is unclear whether they are budgeting sufficient resources to reach the target for treatment success. In this sense, the funding gaps are underestimates, although Brazil and South Africa are relatively wealthy middle-income countries that should be able to find any necessary resources from domestic budgets (Brazil has already increased its NTP budget by 50% since 2002). Apart from the question of whether NTPs are budgeting enough to meet targets, further reasons why the NTP budgets and associated funding gaps could be considered too low are the generally limited budgets for collaborative TB/ HIV activities, especially in African

countries, and the typically small or non-existent budgets for second-line drugs to treat MDR-TB patients (the Russian Federation is a notable exception).

For NTPs to carry out their activities as planned, they must actually receive the funds promised or anticipated. The establishment of the GFATM has not so far caused a decline in grant funding from other sources, and the Fund is thus apparently providing additional money. Nevertheless, its central financing role in several countries, and its smaller but nonetheless important contribution in others, means that the rate at which funds are made available in countries is of considerable importance. If these funds are not received by the NTP, gaps will replace expected GFATM contributions. This is a concern for some countries, where delays in receiving expected disbursements are already evident. The most important example is Mozambique, which had not received funds by the end of 2004, even though its proposal was approved in January 2003. Removing the obstacles to disbursement should be a priority, particularly in countries where GFATM grants contribute a large share of planned budgets.

For those countries that have secured large additional grants or loans, the key question now is whether the NTP can spend the money effectively. In 2003, expenditures were lower than the funding available, and in that year the total amount of money available was much lower than in 2004 or 2005. The most obvious need is for additional staff, particularly those with general and financial management skills. This need has already been recognized in several countries, and additional funds have been sought through the ISAC initiative. For example, China's ISAC proposal includes a budget to support the recruitment of new staff at provincial level. Bangladesh has also identified a need for additional staff at central level, following its successful application to the

When countries succeed in mobilizing additional funds, the new money must be translated into better programme performance. For most countries it is too early to say whether or not this is happening, because the biggest budgetary and funding increases have mostly been in 2004. However, it is striking that India's TB control programme is both relatively low cost and very effective. As data become available for more years, it will be possible to assess the relative cost-effectiveness of TB control in the 22 HBCs, and the reasons for variation among countries.

Some HBCs still have difficulties in providing financial data. South Africa has not yet been able to complete the financial section of the WHO data collection form. A major part of the explanation is that budgeting for TB control is decentralized in South Africa. Decentralization has also affected the completeness of data available for Afghanistan and Thailand. Most NTPs find it more difficult to provide data on expenditures than budgets. Similarly, expenditures are not yet available on the GFATM web site, although the Fund does provide an impressive volume of data on budgets, grant agreements and disbursements. Efforts to follow up data were intensified in the African Region in 2004, and resulted in major improvements in the quantity and quality of data collected. Similar efforts are now needed in other regions, both for the HBCs and other countries.

In summary, financing for global TB control has improved since 2002, dramatically in some countries. Some HBCs now have sufficient funds, but must show that they can spend them effectively; some have no apparent shortfall, but should verify that their budgets are sufficient to meet targets; some have an obvious funding gap, and must focus on raising the money needed to improve programme performance.

# Profiles of high-burden countries

# Afghanistan

Afghanistan has undertaken a programme of health service reconstruction. With help from international partners, funds have been mobilized to create an NTP and to start DOTS activities. The DOTS strategy is included in the country's basic package of health-care services (BPHS). Afghanistan has brought together many NGOs in a common effort to deliver DOTS services, and their involvement has been critical in carrying out programme activities. Along with the general health system, TB control services face several impediments, notably an inadequate number of health facilities. continuing insecurity in many areas and staff shortages at all levels. Nevertheless, the NTP has made substantial progress in recent years. DOTS coverage has increased slowly and treatment success rates have been close to or above the global target for four consecutive years. Improving the currently low case detection rate will

require improvements in the security situation. In Afghanistan, more women than men seek treatment from the public TB control programme; it is possible that more men than women are treated by private practitioners. Private physicians and other healthcare providers including community volunteers are being encouraged to engage in DOTS.

### System of TB control

Over the past two decades, the health service infrastructure collapsed; reconstruction is hampered by the dangers of working in regions where the central government is not fully in control. Nevertheless, progress has been made in rebuilding the general health system, including TB services. The DOTS strategy is a component of the BPHS, and since 2002 the NTP has been strengthened at all levels.

The NTP consists of a central unit under the MoPH General Directorate

of Health Care and Promotion, which is responsible for the overall implementation and management of the NTP, and for policy development. The National TB Institute (NTI) at the central level in Kabul supports training, technical assistance, operational research and laboratory activities. In provinces and districts, TB coordinators supervise and monitor DOTS activities in general health service facilities on four levels: health posts, basic health centres, comprehensive health centres and district hospitals.

There is no national reference library (NRL) in Afghanistan, but the NTI is upgrading its activities so that it can function as an NRL. Additionally, there are eight regional and 144 district laboratories for diagnostic activities.

### Surveillance and monitoring

Surveillance was improved with the introduction of the DOTS strategy in the late 1990s and, although the 2003 estimate of 53% DOTS coverage is probably optimistic, there was a steady rise in the number of smearpositive cases diagnosed between 1997 and 2002. With these improvements, the estimated case detection rate was 19% in 2002 and 18% in 2003. Although the case detection rate is not expected to be much higher than this, a tuberculin skin-test survey carried out in Kabul in 20001 suggests that the national incidence rate of 150 smear-positive cases per 100 000 population could be an overestimate. This is one aspect of case detection that needs further scrutiny in Afghanistan. Another is the unusual finding, noted in Global Tuberculosis Control 2004, that many more women seek treatment from the DOTS programme than men, especially among young adults. Operational research to address this issue is almost complete,

### **PROGRESS IN TB CONTROL IN AFGHANISTAN**

### **Indicators**

DOTS treatment success, 2002 cohort	87%
DOTS detection rate, 2003	18%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	8%
Government contribution to total TB control costs, including loans, 2004	NA
Government health spending used for TB control, 2004	NA

### **Major achievements**

- Formation of the organizational structure and terms of reference of the central NTP unit under the General Directorate of Health Care and Promotion of the MoPH
- Definition of the structure and roles of the TB laboratory network, including the National TB Institute and provincial and district laboratories
- Revision of the national TB guidelines and translation into Dari and Pashtu languages
- Training of over 900 health personnel on DOTS implementation and expansion

### **Major planned activities**

- Expand DOTS and integrate TB control activities into the basic package of essential health services
- Improve capacity of NGOs and other partners, and involve the private sector and the community in TB activities
- Provide adequate supplies and equipment for laboratories throughout the country in a timely manner
- Establish an EQA system for smear microscopy

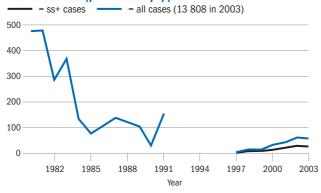
NA indicates not available.

<sup>&</sup>lt;sup>1</sup> Dubuis M et al. A tuberculin skin test survey among Afghan children in Kabul. International Journal of Tuberculosis and Lung Disease 2004, 8:1065-1072.

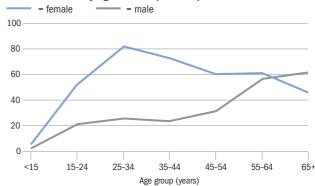
### **AFGHANISTAN**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	23 896 943	DOTS coverage (%)	15	12	38	53
Global rank (by est. number of cases)	21	Notification rate (all cases/100 000 pop)	33	46	60	58
Incidence (all cases/100 000 pop/year)	333	Notification rate (new ss+/100 000 pop)	14	21	28	27
Incidence (new ss+/100 000 pop/year)	150	Detection of all cases (%)	10	14	18	17
Prevalence (all cases/100 000 pop)	671	Case detection rate (new ss+, %)	9.0	14	19	18
TB mortality (all cases/100 000 pop/year)	93	DOTS case detection rate (new ss+, %)	9.0	14	19	18
TB cases HIV+ (adults aged 15-49, %)	0.0	DOTS case detection rate (new ss+)/coverage (%)	60	117	50	34
New cases multidrug resistant (%)	7.3	DOTS treatment success (new ss+, %)	86	84	87	_

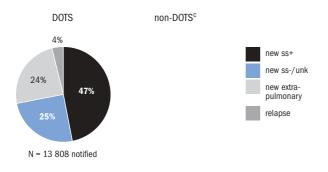
### Notification rate (per 100 000 pop)



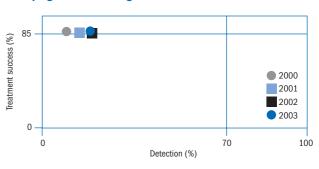
### Notification rate by age and sex (new ss+)<sup>b</sup>



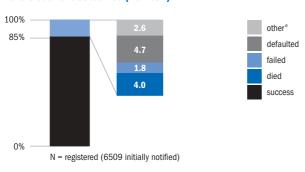
### **Case types notified**



### **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

Absence of a graph indicates that the data were not available or applicable.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

and the results will be available in early 2005.

Treatment success among DOTS patients registered in 2002 was 87%, and has exceeded the 85% target in three of the last four annual cohorts. With respect to monitoring progress towards the Millennium Development Goals, the focus in Afghanistan is still on assessing TB burden and trends, and on evaluating DOTS implementation.

### Improving programme performance

Given the dangers of working in some provinces, national and international experts are sometimes unable to carry out supervision and monitoring visits. Nevertheless, a network of 46 national TB experts has been established, including an NTP manager, a deputy NTP manager, a National Surveillance Officer, a National Logistics Officer, the NTI director and a deputy, eight regional coordinators and 32 provincial coordinators. National TB guidelines have been revised and translated into the Dari and Pashtu languages.

More than 900 health personnel have been trained to provide DOTS services since early 2002. Once the organization of the NTP is complete, the priorities will be to further develop the HR development strategy and to increase training of staff at all levels. A national workshop on HR development for TB control was conducted by the NTP and WHO in March 2004 to revise the basic curricula for all health personnel through the development of appropriate learning materials and training schedules. Five medical schools are preparing training material and courses for all disciplines and are introducing DOTS into the undergraduate curriculum.

Reconstruction of health services has taken place through contracting NGOs to provide basic health services, including TB control, in geographically defined areas. Contracts have been made with 30 NGOs.

In September 2003, the NTP and WHO, in agreement with other partners, procured anti-TB drugs in bulk through the GDF; this supply should cover the needs for 2004 and part of 2005. To maintain regular supplies to

all regions, a national warehouse of anti-TB drugs and laboratory consumables was set up at the NTI, and a computer programme for calculating drug needs and requests has been developed. There are no data on drug resistance, DST is not performed and second-line drugs are not avail-

Other areas in which programme performance needs to be improved include diagnostic and laboratory services, links with other health-care providers and links with the community. The need for collaborative TB/HIV activities is unclear, given the lack of information about the prevalence of HIV.

### Diagnostic and laboratory services

Diagnostic and laboratory services in Afghanistan face major difficulties because of inadequate laboratory equipment and supplies, limited numbers of trained staff and high staff turnover. In 2004, microscopes, reagents and other laboratory materials, including microscopy slides and sputum containers, were purchased and distributed with support from donors. Once basic infrastructure is developed, the priorities will be training of staff, the establishment of an EQA system, and regular monitoring and supervision.

### TB/HIV coordination

No data are available on the prevalence of HIV in the general population or in TB patients. A rapid appraisal of the HIV situation is planned, which will provide an estimate of the prevalence of HIV in the general population and among various vulnerable groups.

### Links with other health-care providers

The NTP regards the involvement of private sector providers as an important component of DOTS implementation and expansion. Many patients are treated privately, but private physicians are not yet involved in DOTS services. The NTP plans to establish a PPM-DOTS task force and to develop PPM guidelines. Progress in including all relevant public sector providers in DOTS has been made, and public hospitals, medical colleges, prison health-care services and army health facilities are now involved in many areas.

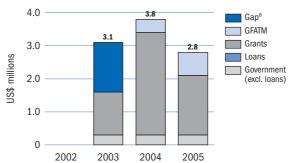
### Links with the community

There is community involvement in TB control activities in Kabul City, where around 10 000 widows have been trained to assist with health education. In Nemruz Province, local people help with TB case referral.

### **Partnerships**

An interagency coordination committee (ICC) for TB control has been established and holds regular meetings in Kabul. A country coordination mechanism (CCM) to facilitate support from the GFATM also exists, and meets monthly to address technical and operational issues. WHO and JICA are the main technical partners, and several NGOs including the Anti-TB Association, CARE International, COOPI, GMS, LEPCO, MEDAIR and MSF are providing additional technical assist-

### NTP budget by source of funding



<sup>&</sup>lt;sup>a</sup> Funding gaps exist in 2004 and 2005. However, they have not been quantified due to the rapidly changing social, economic and security situation and are therefore not shown in the graph. See text for further details.

### **AFGHANISTAN**

ance. CIDA, the Government of Italy and USAID are the major funding partners.

### **Budgets and expenditures**

Budget and expenditure data are limited for Afghanistan. The NTP budget has been approximately US\$ 3-4 million in each year 2003–2005. Almost all funding is provided by grants, including from the GFATM. It is extremely difficult to estimate either the total funds needed or the funding gap because of the highly volatile situation in the country. Although a funding gap of US\$ 1.5 million was reported for 2003 (see Global Tuberculosis Control 2004), the general situation has deteriorated and it is likely that in 2004 and 2005 the funding gap is much greater. This is illustrated by the funding gap for the BPHS, which includes TB control. For the years 2004 and 2005, the funding gap for the BPHS is US\$ 49 million and US\$ 43 million respectively, and is expected to increase to US\$ 70 million in 2006.

A breakdown of the NTP budget by line item is not available for any year 2003-2005, although expenditures on drugs were reported to be about US\$ 1 million in 2003 (equivalent to about US\$ 74 per patient treated), and relatively large investments in infrastructure were made in the same year.

# Bangladesh

Bangladesh adopted the DOTS strategy in 1993. Since then, the NTP has expanded to cover nearly all of the country. For many years, NGOs have been largely responsible for delivering DOTS services and have had a formal involvement in the NTP since 1994. Their collaboration has been instrumental in promoting DOTS and achieving high DOTS coverage. Participation of NGOs in programme delivery continues to be an enormous asset, while the government ensures coordination and sustainability of TB control. With TB control a government priority, recognized as an essential service to be delivered by the health system, the NTP needs to build capacity and strengthen programme management. This is now a matter of urgency as there has been a large increase in funding, mainly from the GFATM, and the amount of money available for TB control almost tripled in 2004 and 2005. Thanks to this encouraging financial position, ambitious plans have been made to dramatically increase case detection and to accelerate a comprehensive programme to

strengthen laboratories. With many and diverse partners from the public and private sectors, clear central leadership will be crucial to ensure coordination, to maintain momentum and to undertake the expanded activities now made possible through the additional funding.

### **System of TB control**

The NTP is recognized as a priority in the revised Health, Nutrition and Population Sector Programme. Under the guidance of the Director-General of Health Services, the NTP manager is responsible for the NTP at central level. At the subnational level, the NTP is integrated into the divisional, district and upazila (subdistrict) general health services. Chest disease clinics, located in district capitals and metropolitan cities, support the NTP by offering diagnostic and treatment services for surrounding areas and serving as referral centres for entire districts. NGOs provide NTP services at upazila level in collaboration with the government; some have their own health-care infrastructure. At the

peripheral level, health inspectors and assistants, medical assistants, village doctors and NGO community health workers provide basic services such as identification and referral of TB suspects, provision of DOT, tracing of defaulters and various behaviourchange communication activities.

The NTP has established a network of nearly 600 sputum microscopy centres, each one covering a population of about 230 000, on average. There is one NRL, which is part of the central public health laboratory, and 45 intermediary laboratories in chest disease clinics. Peripheral laboratories are found in upazila health complexes, in private urban facilities, medical colleges and in health services for special population groups including health services in prisons, the police and industry.

### **Surveillance and monitoring**

The incidence rate of TB in Bangladesh is uncertain because the estimate is based on a 40-year-old tuberculin survey and on local prevalence surveys that may not be nationally representative. Between 1980, when WHO records began, and the introduction of DOTS in 1993, the case notification rate appeared to be in slow decline, despite some variation. Since 1994, there has been a significant rise in the average age of TB patients, allowing for demographic changes, and the notification rates for men are higher in older age groups. Together, these observations suggest that the TB incidence rate is falling, and this assumption underpins the projected year-on-year changes in the estimated smear-positive incidence rate for Bangladesh.

The smear-positive case detection rate increased rapidly after the introduction of DOTS, stabilized between 1998 and 2001 at around 23%, but has recently increased again, reaching 33% in 2003. Most of these gains have been made as the role of upazila health complexes in case-finding has increased, in addition to chest

### **PROGRESS IN TB CONTROL IN BANGLADESH**

### Indicators

DOTS treatment success, 2002 cohort	84%
DOTS case detection rate, 2003	33%
NTP budget available, 2004	94%
Government contribution to NTP budget, including loans, 2004	28%
Government contribution to total TB control costs, including loans, 2004	43%
Government health spending used for TB control, 2004	3%

### **Major achievements**

- Expansion of DOTS and initiation of PPM-pilot projects in Dhaka City
- Introduction of DOTS in prisons, academic institutions and workplaces
- Sustained strong collaboration between the government and NGOs
- Revision of national guidelines, incorporating new treatment regimens with FDCs, and laboratory guidelines
- Expansion of EQA for smear microscopy to most microscopy centres

### **Major planned activities**

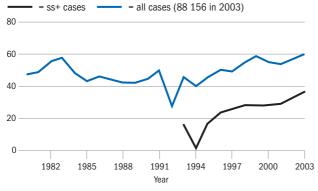
- Create new microscopy centres in populations of more than 300 000
- Provide basic training for newly appointed technicians and refresher training for all laboratory staff
- Implement activities according to GFATM project proposal, in order to improve case detection

NA indicates not available.

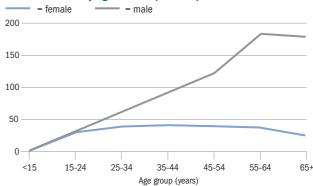
## **BANGLADESH**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	146 736 131	DOTS coverage (%)	92	95	95	99
Global rank (by est. number of cases)	5	Notification rate (all cases/100 000 pop)	55	54	57	60
Incidence (all cases/100 000 pop/year)	246	Notification rate (new ss+/100 000 pop)	28	29	33	37
Incidence (new ss+/100 000 pop/year)	111	Detection of all cases (%)	22	22	23	24
Prevalence (all cases/100 000 pop)	490	Case detection rate (new ss+, %)	25	26	29	33
TB mortality (all cases/100 000 pop/year)	57	DOTS case detection rate (new ss+, %)	23	25	29	33
TB cases HIV+ (adults aged 15-49, %)	0.1	DOTS case detection rate (new ss+)/coverage (%)	26	26	30	33
New cases multidrug resistant (%)	1.4	DOTS treatment success (new ss+, %)	83	84	84	_

#### Notification rate (per 100 000 pop)



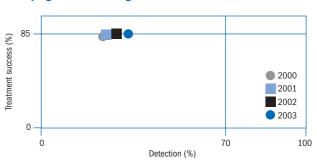
#### Notification rate by age and sex (new ss+)b



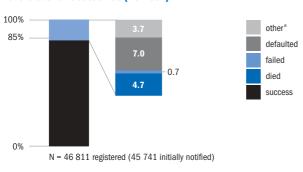
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

disease clinics, which were the dominant source of patients in 1995. Since 2000, the DOTS programme has also reported more patients from metropolitan areas. Despite these improvements, and notwithstanding uncertainty concerning the true incidence rate, case detection by the DOTS programme is still low. Treatment success was 84% for the 2002 cohort and has been 80% or more since 1998. Default (7%) was the most important reason why treatment success was still below the 85% target in 2002. Stimulated by the need to make a better assessment of the scale of the TB problem, and to provide a baseline for evaluating the epidemiological impact of DOTS, the NTP has drawn up plans to carry out a national disease prevalence survey.

#### **Improving programme** performance

In 2002, DOTS was expanded to Dhaka city. In 2003, national guidelines were updated to strengthen the implementation of DOTS, including the control of childhood TB. Laboratory manuals have been revised and distributed throughout the country; specific guidelines for involving private practitioners and delivering DOTS services in workplaces are being developed. In view of proposed DOTS expansion activities funded by the GFATM, there is a need to strengthen capacity at the central level. Additional management capacity and technical assistance are urgently needed if the planned activities are to be implemented on schedule.

Collaboration with NGOs and additional partners in the metropolitan city centres has been expanded. With the increasing number of partners, strong supervision and standardized systems for referral, recording and reporting need to be developed. With different NGOs working in the same area, the supervision, structure and accountability between NGOs, the NTP and the Chief Health Officer in metropolitan city areas also need to be addressed.

A TB control steering committee was established to support, direct and monitor procedures and activities to ensure that NTP and global targets are

reached. In late 2003, as noted above, international partners assisted the government in developing a plan for a national prevalence survey in Bangladesh.

Short-course treatment for all TB cases has been further standardized with the introduction of new treatment regimens and FDCs. The new treatment regimens follow WHO recommendations and are more consistent with private sector prescription practices, which may facilitate increased referral of patients. They also simplify drug management at all levels. The difficulties of ensuring drug quality and an uninterrupted drug supply have been alleviated by the successful application for funding by the NTP to the GDF. There is no national policy on the management of MDR-TB, and MDR-TB cases are not treated within the NTP. However, the Damien Foundation Bangladesh (DFB) treats all confirmed MDR-TB cases in the areas it covers. The National Institute of Diseases and Chest Hospitals also treats MDR-TB. Some second-line drugs are produced in the country.

A budget for both DRS and DOTS-Plus will be included in the country's application to the fifth round of the GFATM. Should the GFATM application be approved, Bangladesh will apply to the GLC for reduced-price quality-assured second-line drugs and for technical assistance in implementing sound MDR-TB control measures.

Three other areas in which programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

EQA is becoming a routinely accepted standard in many NGO-supported areas in Bangladesh, and NGOs are offering their services to the government to expand EQA. A major challenge for the NTP is to refocus the NRL on training, EQA, expansion of culture services and drug susceptibility testing, in addition to routine microscopy work. Future laboratory priorities include basic training for newly appointed technicians and refresher training for all laboratory staff on smear microscopy and quality assurance. Diagnostic services will be expanded by establishing new microscopy centres in upazilas with population coverage greater than 300 000. By 2005, EQA for smear microscopy should be available in all urban and rural diagnostic centres.

#### TB/HIV coordination

The HIV prevalence in the adult population (aged 15-49 years) and the proportion of HIV-positive patients among adult TB cases are still low at 0.01% and 0.1%, respectively, according to the latest UNAIDS and WHO estimates. A similar figure for HIV prevalence among TB cases was found in Dhaka in 1999. There is as yet little collaboration between the NTP and the national HIV/AIDS programme.

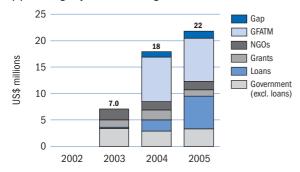
#### Links with other health-care providers

Most DOTS implementation in Bangladesh has been done by NGOs, and during 2004 their involvement has increased. The main NGO partners include the Bangladesh Rural Advancement Committee (BRAC) and DFB, who together cover most of the rural districts in the country; urban areas are covered mainly by other NGOs. There are a number of PPM-DOTS initiatives in Bangladesh. Several private chest physicians in Dhaka have become involved in DOTS services, and the participation of more private practitioners is needed. DFB is expanding its cadre of private "village doctors", who are currently responsible for the detection of about 10% of patients and the provision of DOT to 45% of patients in DFB areas. BRAC has started similar initiatives in periurban areas, while in rural areas they deliver DOT through a network of community workers. Recently, the NTP and collaborating NGOs have begun to include medical colleges, prison health services and the private corporate sector in DOTS activities.

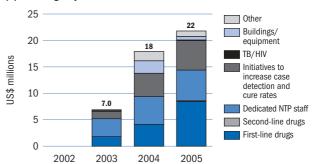
#### **Partnerships**

Several technical partners participate in TB control activities in Bangladesh, led by BRAC and the DFB. Thanks to the joint efforts of the partners, the CCM has made a successful application for funding from the GFATM. Financial support is also provided by

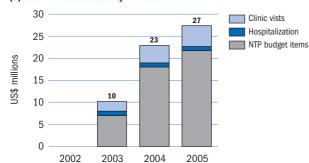
#### (a) NTP budget by source of funding



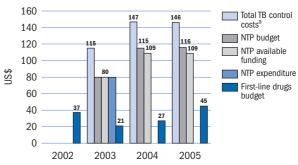
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



<sup>&</sup>lt;sup>a</sup> Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

CIDA, the World Bank and other partners through general funding for the health sector.

#### **Budgets and expenditures**

The TB control budget data reported to WHO include both a budget for the NTP and the budgets for the two major NGOs that are responsible for DOTS implementation in most of Bangladesh (i.e. BRAC and DFB). The budgets for both 2004 and 2005 are substantially higher than in previous years, at about US\$ 20 million compared with US\$ 7 million in 2003. This reflects an ambitious plan to more than double the number of patients treated between 2003 and 2005. Most of the budget is funded for both 2004 and 2005, mainly because of increased funding from a World Bank

credit and a substantial GFATM grant. There is a funding gap of US\$ 1-2 million in both 2004 and 2005; this money is needed to cover the national prevalence survey, to recruit staff in order to strengthen management at the central level and to carry out additional activities to increase case detection and treatment success rates. The substantially improved funding position means that spending on TB control by the NTP and the major NGOs could almost triple between 2003 and 2005. The larger budgets in 2004 and 2005 will allow for increased spending on first-line drugs, in line with projected increases in the number of patients treated as well as the development of a buffer stock in 2005 (this buffer stock is the reason for the relatively high budget for first-line drugs

in 2005). They will also allow for some investment in infrastructure, and increased spending on initiatives aimed at improving case detection. The NTP budget per patient is projected to increase from US\$ 80 in 2003 to US\$ 116 in 2005; if this happens, the total cost of TB control, including visits to health clinics for observation of treatment and monitoring, and limited hospitalization, is projected to increase from US\$ 10 million in 2003 to US\$ 27 million in 2005 (from US\$ 115 to US\$ 146 per patient treated). It remains to be seen whether the increased funding can be absorbed effectively and whether increased expenditures result in improved case detection.

# **Brazil**

Brazil is one of the largest countries in the WHO Region of the Americas and it has the highest TB burden in the region. Providing TB control and other health services throughout the country poses immense organizational and logistic challenges. However, the data from recent years indicate a steady downward trend in TB incidence in Brazil. Although DOTS is currently available to only some 35% of the population, a concerted effort is being made to include all of the 315 highburden municipalities by 2007. There is increasing awareness of the public health importance of TB by the new Brazilian health authorities, who have recognized the DOTS strategy as the best solution to Brazil's TB control problems. The Brazilian MoH has now prioritized the DOTS strategy in its new programme for TB control. TB and leprosy were declared national priority diseases in 2004 and increased government funds were assigned to control them. In addition, 2004 saw the launch of Brazil's Stop TB Partnership involving numerous technical and do-

nor agencies and other public and private sector partners in TB control.

#### **System of TB control**

Brazil adopted the DOTS strategy in 1998, establishing it in four states as demonstration areas. Brazil has a massive and complex decentralized health-care system. At the state and municipal levels, the TB control programme is represented by local TB coordinators who are responsible to the respective state and municipal health secretaries. Recently, the government created the position of Secretary of Health Surveillance (SVS) within the new structure of the MoH, which has given added priority to TB control. The SVS has also facilitated collaboration of the NTP with the national laboratory and the HIV/AIDS programmes. TB patients are treated in the out patient facilities of the public health service and only a few complicated cases require hospitalization.

TB laboratory services are carried out by the National Public Health Laboratories Network. There is one NRL, 27 central public health laboratories (one per state) and more than 4000 local laboratories.

#### **Surveillance and monitoring**

Among the HBCs, Brazil has a relatively comprehensive TB surveillance system, and the observed downward trend in the case notification rate probably represents a real decline in incidence. The rate of fall is about 3% per year both for smear-positive and for all TB cases, but a faster rate of decline should be achievable by an expanded DOTS programme. DOTS coverage increased to 34% in 2003 and the case detection rate to 18%, giving a detection rate of 55% within DOTS areas. However, an estimated 81% of all new smear-positive TB cases are found nationally (by DOTS and non-DOTS services), suggesting that Brazil could meet and even exceed the target of 70% case detection simply by ensuring that patients already notified are correctly diagnosed and treated by DOTS services.

As DOTS coverage increases, the monitoring of patients on treatment needs to be carried out more rigorously. The treatment success rate under DOTS in 2002 was 75%, with 18% of patients lost through default or transfer to other treatment centres without follow-up. A large proportion of patients (29%) completed treatment without evidence of smear conversion. Among patients registered for retreatment, only 36% were cured. An additional 24% completed treatment, but the demonstration of smear conversion is vital for re-treatment patients, who could be carrying drugresistant bacilli. Treatment success rates were even lower among the subset of patients receiving re-treatment after default (51%) or failure (42%). As control efforts intensify, Brazil's system of routine surveillance should be strengthened as the main instrument for monitoring trends in TB cases and deaths and for evaluating the future impact of control measures.

#### **PROGRESS IN TB CONTROL IN BRAZIL**

#### **Indicators**

DOTS treatment success, 2002 cohort	75%
DOTS case detection rate, 2003	18%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	86%
Government contribution to total TB control costs, including loans, 2004	94%
Government health spending used for TB control, 2004	0.3%

#### **Major achievements**

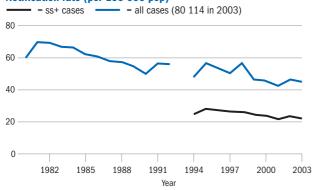
- Approval of TB national plan (2004–2007) by the government
- Launch of the Stop TB Partnership in October 2004
- Organization of 15 regional meetings to discuss the national TB plan and strategies for DOTS expansion, attended by all 27 state TB control coordinators and by the municipal TB control coordinators of all 315 priority municipalities
- Creation of a Task Force Group to monitor and assist the states and priority municipality in DOTS implementation

#### **Major planned activities**

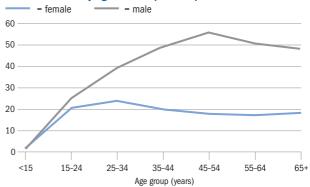
- Establish a TB/HIV coordination body in 2005 to implement strategies to increase provision of VCT to TB patients
- Increase microscopy coverage in all 315 priority municipalities and improve quality control of existing microscopy centres
- Implement a national workplan involving different sectors of civil society and the community for TB control

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	178 470 430	DOTS coverage (%)	7.0	32	25	34
Global rank (by est. number of cases)	15	Notification rate (all cases/100 000 pop)	45	43	46	45
Incidence (all cases/100 000 pop/year)	62	Notification rate (new ss+/100 000 pop)	24	22	23	22
Incidence (new ss+/100 000 pop/year)	28	Detection of all cases (%)	67	65	72	73
Prevalence (all cases/100 000 pop)	92	Case detection rate (new ss+, %)	79	75	82	81
TB mortality (all cases/100 000 pop/year)	8.2	DOTS case detection rate (new ss+, %)	7.5	8.0	9.6	18
TB cases HIV+ (adults aged 15-49, %)	3.8	DOTS case detection rate (new ss+)/coverage (%)	108	25	38	55
New cases multidrug resistant (%)	0.9	DOTS treatment success (new ss+, %)	73	67	75	_

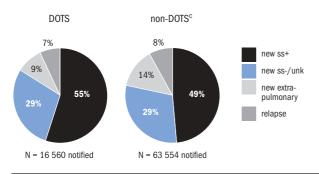
#### Notification rate (per 100 000 pop)



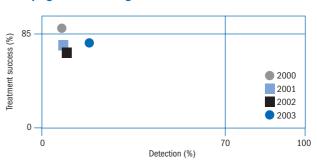
#### Notification rate by age and sex (new ss+)<sup>b</sup>



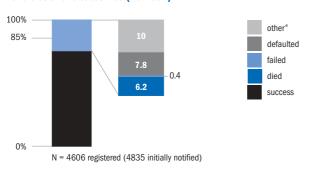
#### **Case types notified**



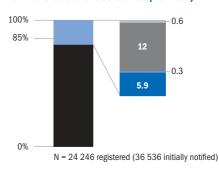
#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



#### Non-DOTS treatment outcomes (new ss+)



ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

The creation of the SVS will strengthen Brazil's TB surveillance system by integrating TB with surveillance and control of other endemic diseases and improving coordination; however, it is also important to optimize Brazil's information system (SINAN) for TB surveillance and DOTS monitoring.

#### **Improving programme** performance

The MoH, together with health authorities at state and municipal levels, is working hard to strengthen TB control and to reorganize primary health-care services for DOTS implementation. It is important to ensure better integration and coordination of activities at the primary health-care level, particularly those included in the Family Health Programme (Programa de Saúde da Família - PSF) and the Community Outreach Programme (Programa de Agentes Comunitários - PAC). Training in DOTS TB control is currently being provided to other public and private health-care professionals. However, appropriate training and continuous good quality supervision and monitoring activities from the state to the municipal and from the

municipal to the local levels are indispensable for effective DOTS implementation. Training for 20 000 Family Health Teams is planned for 2005. Another important area for improving programme performance is the provision of TB control services in high-risk populations such as the indigenous groups and prison populations.

A national TB control plan for 2004-2007 was approved by the government in 2004. It aims to strengthen the NTP and to reach 100% DOTS coverage in the 315 priority municipalities that account for an estimated 70% of the country's TB burden. The plan includes the creation of a training task force to improve HR capacity for TB control, with the goal of offering DOTS services in all basic health-care facilities in all the priority municipalities by the end of 2007. During 2004, five regional meetings were organized to discuss the national TB plan and strategies for DOTS expansion in the first quarter; two more cycles of five regional meetings each were conducted to monitor this plan in the second and third quarter. All 27 state TB control coordinators and the municipal TB control coordinators of

the priority municipalities attended one of these meetings. A Task Force Group was created in 2004 to monitor and assist the states and priority cities in DOTS implementation.

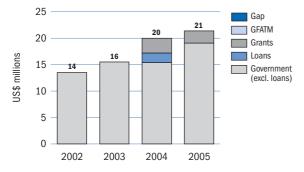
#### Diagnostic and laboratory services

As DOTS services expand to the 315 priority municipalities, laboratory capacity needs to be increased, and quality assurance must be introduced. The TB laboratory manual is under revision and the task force organizing training has begun the strengthening of laboratory services; this will continue in 2005. Laboratory information systems and monitoring and supervision will also be improved. During 2004, three regional managerial courses, with the support of an international consultant, were developed to increase the capacity for sputum smear microscopy and quality assurance. More than 800 laboratory personnel countrywide were trained on those topics.

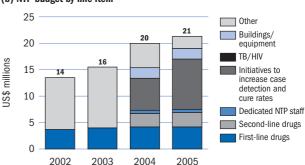
#### TB/HIV coordination

Brazil is a country with a concentrated HIV epidemic. In 2003, the estimated HIV seroprevalence in the general

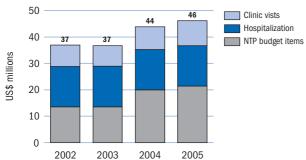
#### (a) NTP budget by source of funding



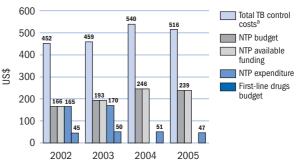
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details

population was 0.65%. The NTP estimates that the prevalence of HIV among new TB patients was 8%. This is substantially higher than the WHO estimate of 3.8%, which may underestimate the effect of shared risk factors for TB and HIV. ART is available to all HIV-infected individuals (including TB patients) through the public health system. The recently created SVS has contributed to the collaboration between the NTP and the National AIDS Programme, and to better coordination between them. A national TB/ HIV plan is now in place and includes the establishment of a TB/HIV coordination body in 2005, plus strategies to increase the provision of VCT to TB patients and to provide DOTS services to HIV-positive individuals suffering from TB.

#### Links with other health-care providers

Private hospitals and clinics are required to refer TB suspects and cases to government TB facilities. A small number of NGOs are involved in DOTS provision, and the NTP is planning to host a meeting of national NGOs in 2005 to formulate a collaborative agreement. Brazil has no PPM-DOTS

taskforce or guidelines, but plans to strengthen ties with the Brazilian Society of Pulmonology and Phthisiology through a collaborative agreement. There are also plans to enhance the involvement of the Brazilian research network in the 2004-2007 national plan for DOTS expansion, particularly in the area of operational research.

#### **Partnerships**

Brazil has established effective international technical partnerships with agencies such as PAHO, WHO, IUATLD and CDC to support adequate DOTS implementation and expansion. Funding partners include USAID (two TBCTA projects), DFB and GLRA. The launch of the Stop TB Partnership in Brazil in October 2004 signifies another important step towards involving different sectors of civil society and the community in TB control, as does the launch of a national advocacy plan to disseminate TB and DOTS informa-

#### **Budgets and expenditures**

The NTP budget has been steadily increasing, from US\$ 14 million in 2002 to US\$ 21 million in 2005 (a 50% increase in four years). As would be expected in an upper-middle income country, the budget is fully funded and most financing is provided by the government, although grant funding was received in 2004 and is expected in 2005. This sound funding situation reflects the commitment of both the government and the international community to TB control. The budget for first-line drugs has been consistently around US\$ 4 million and around US\$ 50 per patient. In 2004 and 2005, there has been an increase in the budget for activities aimed at improving case detection and cure rates, including an extensive training programme and upgrading of the laboratory network. NTP expenditures were US\$ 14 million, equivalent to about US\$ 170 per patient treated, in both 2002 and 2003. When costs not covered by the NTP budget are included (i.e. 2509 dedicated TB hospital beds and visits to clinics for DOT and monitoring during treatment), the cost per patient treated is estimated at US\$ 450-550 during the period 2002-2005. The total cost of TB control is estimated at US\$ 37-46 million.

# Cambodia

Cambodia achieved nationwide DOTS coverage at district level in 1998, at a time when the health services were still relatively centralized. Since then, a policy of progressive decentralization has been followed, designed to improve the access of the population to health care. With the establishment of peripheral health centres, the NTP has gradually introduced its activities in these settings, resulting in substantially improved access to TB control services. By the end of 2004, the remaining health centres will be included. A national TB prevalence survey in 2002 yielded a great deal of valuable information, which continues to be applied in strengthening the programme. Results will be published and will serve as an important basis for the assessment of the burden of TB and the impact of DOTS services on the TB epidemic. The shortage of staff to support the expanding programme is now being addressed, and there are plans to tackle the urgent need for better coordination between the TB and HIV control programmes.

#### System of TB control

Cambodia's NTP operates under the responsibility of the National Center for Tuberculosis and Leprosy Control (CENAT) and within the overall national health system. It comprises TB referral hospitals, provincial TB centres and district TB units. In 1994, TB control was decentralized from provincial hospitals to district hospitals, and in 1999 to health centres. As of 2003, more than 145 TB units and 700 health centres are implementing the DOTS strategy.

There are 180 laboratories in the country including the TB reference laboratory of CENAT, which is responsible for the development of training materials, training of laboratory technicians, and supervision and quality assurance of the provincial laboratories. The reference laboratory carries out culture of mycobacteria and HIV testing but not regular drug susceptibility testing, which will be started in the near future. There are 24 provincial laboratories with responsibility for the supervision and training of health

centre staff in sputum smear microscopy and quarterly reporting to CENAT.

#### **Surveillance and monitoring**

Cambodia's case detection rate under DOTS was 60% in 2003, after the noticeable upturn in case detection since 2001. This assessment of the case detection rate is based on an estimate of incidence that pre-dates the 2002 prevalence survey. Analysis of the results of that survey will allow a reassessment of the burden of TB in the country and of the case detection rate. The proportion of all cases diagnosed as smear-positive in 2003 was 67%, falling from the highest recorded level of 82% in 1999, possibly because of improvements in diagnosis (fewer false-positives).

The treatment success rate reported among new smear-positive cases has exceeded 90% since 1995, which is unusually high given that 13% of TB patients were thought to be coinfected with HIV in 2003. The success rate for re-treatment patients in 2002 was also remarkably high (89%). Despite some uncertainty about case detection and treatment success, Cambodia is in a strong position to evaluate the future impact of the expanding DOTS programme on TB prevalence, incidence and deaths. As found in population-based surveys in other countries, the 2002 survey in Cambodia has yielded much more than an estimate of prevalence, including data that suggest numerous ways improving routine diagnosis and treat-

### **Improving programme** performance

The strong commitment of the Cambodian government to poverty elimination and health infrastructure development will have a positive effect on the control of TB in the future. Capacity building for DOTS expansion in all areas of the NTP continues to be a leading priority for the programme. In response to the low ac-

#### PROGRESS IN TB CONTROL IN CAMBODIA

#### Indicators

DOTS treatment success, 2002 cohort	92%
DOTS case detection rate, 2003	60%
NTP budget available, 2004	81%
Government contribution to NTP budget, including loans, 2004	10%
Government contribution to total TB control costs, including loans, 2004	44%
Government health spending used for TB control, 2004	NA

#### **Major achievements**

- Implementation of DOTS in 320 additional health centres during 2003, for a total of 706 out of 856
- Community-based DOTS introduced in collaboration with NGOs in four operational districts
- Introduction of six-month short-course treatment regimen in three operational districts

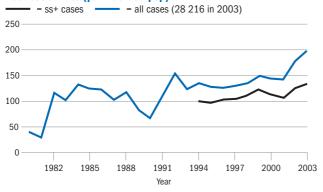
#### **Major planned activities**

- Implement DOTS in an additional 150 health centres to reach 100% coverage by end 2004
- Conduct follow-up study of TB suspects detected during the national TB prevalence survey conducted in 2002
- Continue to train health-care workers on community DOTS and six-month shortcourse treatment regimen

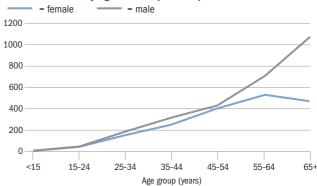
NA indicates not available.

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	14 143 527	DOTS coverage (%)	99	100	100	100
Global rank (by est. number of cases)	23	Notification rate (all cases/100 000 pop)	144	142	178	199
Incidence (all cases/100 000 pop/year)	508	Notification rate (new ss+/100 000 pop)	113	107	125	134
Incidence (new ss+/100 000 pop/year)	225	Detection of all cases (%)	27	27	35	39
Prevalence (all cases/100 000 pop)	762	Case detection rate (new ss+, %)	49	47	55	60
TB mortality (all cases/100 000 pop/year)	95	DOTS case detection rate (new ss+, %)	49	47	55	60
TB cases HIV+ (adults aged 15-49, %)	13	DOTS case detection rate (new ss+)/coverage (%)	49	47	55	60
New cases multidrug resistant (%)	0.0	DOTS treatment success (new ss+, %)	91	92	92	_

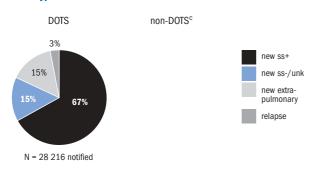
## Notification rate (per 100 000 pop)



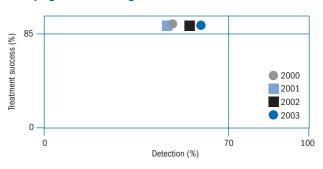
#### Notification rate by age and sex (new ss+)<sup>b</sup>



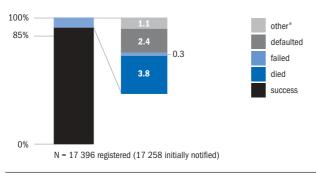
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

## **CAMBODIA**

cess to health services and DOTS in some areas. DOTS services were expanded to 320 additional health centres in 2003. There are plans to implement DOTS in the remaining 150 health centres by the end of 2004. A six-month short-course chemotherapy regimen has been introduced in pilot studies in three operational districts, and training in the new regimen for health-care workers will continue into 2005. A follow-up study of TB suspects detected during the 2002 prevalence survey has started and will be completed during 2004. A drug resistance survey conducted in 2000-2001 found that the prevalence of MDR-TB was negligible among new cases and 3% among re-treatment cases.

The lack of human resource capacity remains a challenge for the NTP. At the request of CENAT, a representative from KIT met with key personnel and staff focus groups to assess human resource development needs in 2003. A workshop was subsequently organized to develop an outline for management training; training activities have been intensified and new staff have been recruited. There is still an urgent need for both in-country and

international training for staff (including managers), and to recruit more staff. The NTP is planning to address these issues through recruitment of staff from outside the NTP with the aid of partners, making use of additional funding from the GFATM and the World Bank.

Other areas where programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers and the community.

#### Diagnostic and laboratory services

Two of 24 provincial laboratories have been upgraded during 2003 to perform culture and drug susceptibility testing. The EQA system, introduced in 2002, is still under development and must be strengthened and expanded. There are too few staff with sufficient training to run the laboratory and diagnostic services in Cambodia. During 2005, training programmes will improve technical knowledge and enhance staff motivation. Drug susceptibility testing is not available in Cambodia, but its introduction is considered a priority.

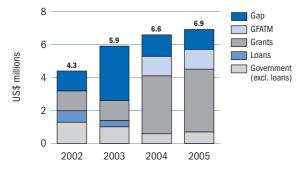
#### TB/HIV coordination

A national TB/HIV prevalence survey in 2003, carried out by the VCT service at CENAT, estimated HIV seroprevalence among TB patients at 12% (similar to the WHO estimate of 13% among adult TB patients). As yet, there are no data on TB incidence or mortality among PLWHA. TB/HIV collaborative pilot studies in four provinces included screening and treatment for TB among PLWHA, isoniazid preventive treatment for PLWHA who are infected with M. tuberculosis, surveillance of HIV in TB patients and ART for HIVinfected TB patients. A workshop to assess the pilot projects concluded that TB/HIV collaboration is hampered by the disease-specific focus of the individual programmes, the quality of TB/HIV counselling and lack of joint IEC material. IEC materials and standardized reporting and recording forms for TB/HIV activities are being developed.

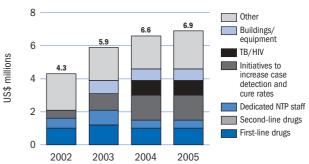
#### Links with other health-care providers

The 2002 prevalence survey showed that among people with TB symptoms who sought any type of health care, 89% went first to the private sector

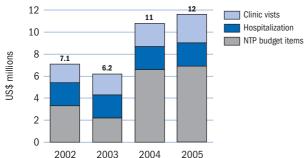
#### (a) NTP budget by source of funding



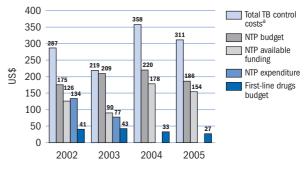
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

### **CAMBODIA**

(pharmacies and doctors). However, the private sector has not yet been formally involved in the NTP in Cambodia. The majority of private providers diagnose and treat TB but the quality of the services provided by them is generally poor, as shown in a study carried out by CENAT in collaboration with the Quality Assurance Project of Cambodia's University Research Corporation. However, many private providers are interested in collaborating with the NTP, and a pilot project involving private practitioners and pharmacies will be launched in 2005.

#### Links with the community

Community-based DOTS has been introduced in four districts, and training started in 2004 for community-based DOT workers.

#### **Partnerships**

Cambodia has a diverse group of technical partners including CDC (TB/HIV pilot programme activities), JICA (training and supervision, laboratory technical support, IEC, procurement, operational research, TB/HIV), KNCV (training and workshops, community DOTS) and WHO (training and supervision, laboratory technical support, IEC, procurement, TB/HIV). The main financial partners are CIDA, GFATM, JICA, USAID, WHO and the World Bank.

#### **Budgets and expenditures**

The NTP budget has increased from about US\$ 4 million in 2002 to almost US\$ 7 million in 2005, in line with planned increases in case detection. Available funding almost doubled between 2002 and 2005, from about US\$ 3 million in 2002 to almost US\$ 6 million in 2005. This improvement is becasue of a large increase in grant funding, including from the GFATM. However, funding gaps have persisted in each year from 2002 to 2005. In both 2004 and 2005, the gap is about US\$ 1 million, equivalent to about 15% of the total budget requirement. The increased budgets in

2004 and 2005 are mainly a result of higher proposed spending on TB/HIV collaborative activities, and initiatives to increase case detection and cure rates (e.g. implementation of community-based care in remote areas and active case-finding).

Reported expenditures were lower than available funding in both 2002 and 2003. On a per patient basis, the NTP budget has varied from between US\$ 175 (in 2002) and US\$ 220 (for 2004), while actual expenditures per patient were US\$ 134 and US\$ 77 in 2002 and 2003, respectively. When costs not covered by the NTP budget are included (i.e. 1200 dedicated TB hospital beds and visits to clinics for DOT and monitoring during treatment), the cost per patient treated is estimated to range from around US\$ 220 to US\$ 360. The total cost of TB control was about US\$ 7 million in 2002. Provided the 2005 budget is fully funded and spent and the projected number of cases are treated, this will rise to almost US\$ 12 million in 2005.

# China

China has seen a radical change in political commitment to TB control during 2003 and 2004. There has been a clear government decision to meet the global targets for diagnosis and treatment of TB by the end of 2005. This decision was endorsed by the State Council at a meeting on TB control in September 2004, and a pledge was secured to make an eightfold increase in central government funding for TB control. Following the accelerated DOTS expansion undertaken in recent years, coverage will reach 95% by the end of 2004 and is expected to reach 100% in 2005. Building on the experience of the severe acute respiratory syndrome (SARS) epidemic, China has further recognized the importance of a public health approach to communicable diseases and has set up a new national Internet-based reporting system, under which all cases of several specified communicable diseases, including TB, must be notified. The recently revised law on infectious diseases also strengthens the mandatory reporting of TB, and this is expected to improve TB case reporting substan-

tially. The main challenge is to ensure the quality of TB services during a phase of rapid expansion and to address the shortages of staff and laboratory services needed to support the expanding programme.

#### **System of TB control**

China introduced DOTS on a wide scale in 1992 by expanding DOTS to 13 of 31 mainland provinces, municipalities and autonomous regions ("provinces" hereafter) using funds from a World Bank loan. By 2000, most counties (1132 of 1208) in these 13 provinces had been using DOTS for at least five years. Further expansion of DOTS activities in other parts of China followed in 2002. By 2003, 91% of the population lived in areas covered by the DOTS strategy. Nationwide coverage is planned for the end of 2005.

The government is increasing its investment in public health substantially, and the MoH has put the control of TB among its top priorities. In 2004, an eight-fold increase in funding for the NTP has been pledged for TB control activities. A recent evaluation of the progress towards the 10year national TB control plan carried out by the MoH, Ministry of Finance, and the National Development and Reform Commission has resulted in further government commitment to TB control at all levels.

The TB laboratory network operates under the guidance of the NTP manager and consists of one national reference laboratory, 31 provincial TB reference laboratories, 336 TB laboratories at the prefecture/city level and 2683 peripheral laboratories. Microscopy is performed by all laboratories, while 16% carry out culture and less than 2% do drug susceptibility testing. Culture is occasionally performed in 5-10% of county laboratories, except in some major cities including Beijing where culture is done routinely for all TB suspects. Drug susceptibility testing is available to diagnose drug resistance at some provincial and prefecture level laboratories.

#### **Surveillance and monitoring**

The estimated incidence rate for China was revised during 20041 and is believed to be falling by 1% per year, as is the measured rate of decline in the annual risk of TB infection over the decade since 1990. However, on the basis of the currently available data, these assessments of trend should be treated as approximate.

China made the second largest contribution to the increase in global case detection between 2002 and 2003, after India. The case detection rate achieved by the DOTS programme was 30% in 2002, and increased sharply to 43% by the end of 2003 as population coverage reached 91%. A rapidly implemented TB control programme faces the challenging task of maintaining quality as the programme

## **PROGRESS IN TB CONTROL IN CHINA**

#### **Indicators** DOTS treatment success, 2002 cohort 93% DOTS case detection rate, 2003 43% NTP budget available, 2004 88% Government contribution to NTP budget, including loans, 2004 74% Government contribution to total TB control costs, including loans, 2004 74% Government health spending used for TB control, 2004 0.5%

#### **Major achievements**

- A State Council TB control meeting on TB control involving all provinces
- Increased political commitment, especially at local levels, and increased funding from government and partners
- Establishment of a nationwide Internet-based system for the compulsory reporting of infectious diseases, including TB
- Monitoring mission to six priority provinces organized by the MoH

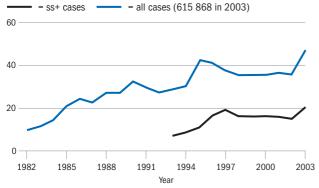
#### **Major planned activities**

- Strengthen system of referral of TB patients from hospitals to local TB dispen-
- Build human resource capacity according to the NTP guidelines
- Expand EQA system to all cities and counties, and drug resistance surveillance in additional provinces

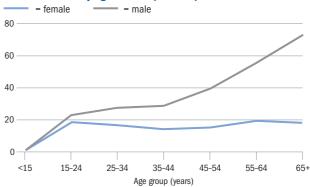
<sup>&</sup>lt;sup>1</sup> Using the annual risk of TB infection (ARTI) measured in 2000, and by applying Stýblo's rule of thumb relating TB incidence to ARTI (smear-positive incidence increases by 50/100 000 population for every 1% increase in ARTI).

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	1 304 196 022	DOTS coverage (%)	68	68	78	91
Global rank (by est. number of cases)	2	Notification rate (all cases/100 000 pop)	36	37	36	47
Incidence (all cases/100 000 pop/year)	102	Notification rate (new ss+/100 000 pop)	16	16	15	21
Incidence (new ss+/100 000 pop/year)	46	Detection of all cases (%)	34	35	35	46
Prevalence (all cases/100 000 pop)	246	Case detection rate (new ss+, %)	34	34	32	45
TB mortality (all cases/100 000 pop/yea	r) 18	DOTS case detection rate (new ss+, %)	31	31	30	43
TB cases HIV+ (adults aged 15-49, %)	0.7	DOTS case detection rate (new ss+)/coverage (%)	45	45	39	47
New cases multidrug resistant (%)	5.3	DOTS treatment success (new ss+, %)	95	96	93	_

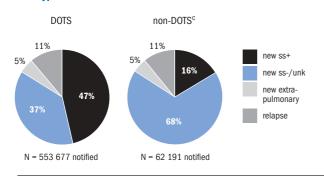
#### Notification rate (per 100 000 pop)



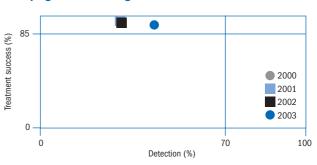
#### Notification rate by age and sex (new ss+)<sup>b</sup>



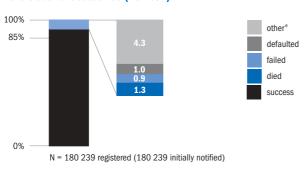
#### **Case types notified**



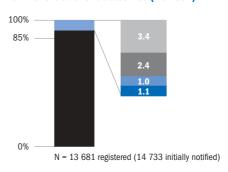
#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



#### Non-DOTS treatment outcomes (new ss+)



ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

expands. At least two aspects of the monitoring data from China need closer scrutiny. One is the steady decrease in the proportion of DOTS patients diagnosed as smear-positive from 1996 to 2003; the other is the exceptionally high treatment success rate, reported to be 93% for the 2002 cohort of new smear-positive patients.

A full analysis of the year 2000 prevalence survey has confirmed that, in the 13 World Bank project provinces implementing DOTS between 1991 and 2000, culture-positive TB prevalence fell by 37% more than in other areas of the country, with a 30% decline directly attributable to DOTS.1 Although the geographical coverage of DOTS increased substantially between 2002 and 2003, the case detection rate within DOTS areas was only 47% in 2003. Case detection will be improved by the new communicable disease surveillance system (implemented by the Chinese Center for Disease Control and Prevention), which has begun to notify TB patients via the Internet from all major hospitals and health centres as well as the TB dispensaries.

#### Improving programme performance

During 2003, 27 of 31 provinces in China began scaling up existing and new TB control projects, and the country's two largest projects - funded by the World Bank/DFID and GFATM - are now fully operational. With anti-TB drugs provided free of charge by grants from the Government of Japan and the central government, 2003 is the first year that all provinces in China have had sufficient resources to implement the complete DOTS technical package. This is the main reason for the increase in case detection for both old and new DOTS areas and accounts for the increase in the case detection rate in DOTS areas from 35% in 2002 to 47% in 2003.

Insufficient human resources both the quantity and expertise of staff - is a major constraint to TB control. China has begun to address this problem by developing a new national TB training plan. With new funding from the GFATM and the ISAC initiative, China plans to recruit and train additional staff at the central and provincial levels.

With the aim of reaching the global targets by 2005, China has developed a national TB health promotion strategy to increase case detection and cure rates, especially among the poor and vulnerable. Special efforts are being made to increase public awareness of TB. The capacity of TB control staff to carry out health promotion activities at national and provincial levels will be enhanced, as will their communication and outreach skills.

Given the size of China, drug resistance surveys are carried out in individual provinces rather than nationally. The first survey began in 1996 in Henan Province, and since then six additional mainland provinces and Hong Kong SAR have reported drugresistance data. China has an organized DRS plan, and many provinces are in various stages of planning and implementation. Another three provinces have completed drug resistance surveys; four more provinces have surveys in progress. A nationwide survey in 2000 estimated that 10% of prevalent bacteriologically confirmed TB cases have MDR-TB disease. MDR-TB patients are treated on an individual basis and have to pay for the services. Second-line drugs are produced in the country and are widely available.

Diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers are three priority areas in which programme performance needs to be improved.

#### Diagnostic and laboratory services

With 500 000 people for every microscopy diagnostic unit, diagnostic services offer a challenge to TB control in China. As activities expand, improving the quality of laboratory services is a priority. Rapid expansion of the new internationally recommended EQA system is also a priority for China, and a new national EQA manual for smear microscopy was developed and issued to TB control institutions at each level. Training courses are being held on EQA implementation. Quality assurance for smear microscopy currently includes a quarterly review, on-site evaluation and panel testing. The national reference laboratories have set an EQA target for 2004 to cover 100% of provincial and prefecture laboratories and 60% of county laboratories. There are, however, no quality assurance systems in place for culture testing.

#### TB/HIV coordination

The Chinese government estimates that there are currently 840 000 people living with HIV in the country, but by the end of 2003, only around 62 000 had been reported to the authorities, of which nearly 9000 were reported with AIDS; reported AIDS deaths have been rapidly increasing. While less than 0.2% of Chinese adults are currently infected with HIV, high rates of HIV infection have been found among intravenous drug users and among people who sold blood plasma to supplement their incomes in provinces such as Anhui, Henan and Shandong.<sup>2</sup> The Government of China is planning to collect data on HIV prevalence among TB patients in provinces known to have a relatively high HIV prevalence, and to use sentinel surveillance or surveys to determine trends in HIV prevalence among TB patients in provinces where the prevalence of HIV is not known. The MoH plans to establish a national TB/HIV coordinating body.

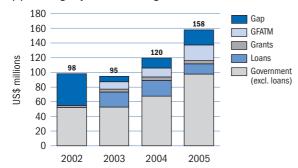
#### Links with other health-care providers

TB suspects and patients seek care from public hospitals at all levels. The focus of PPM DOTS in China is to link hospitals to TB dispensaries, which is potentially the most important way to increase case detection and to improve the quality of patient care. Data from the prevalence survey conducted in 2000 indicate that more than 75% of smear-positive cases are initially managed in either county general hospitals or township hospitals in China. In the past, many cases diagnosed and treated in hospitals were not reported to the TB dispensaries. Patients in hospitals should now also be reported through the new surveillance

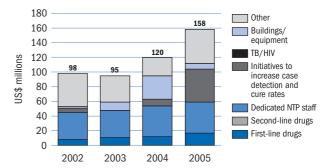
 $<sup>^{\</sup>scriptscriptstyle 1}\,$  China Tuberculosis Control Collaboration. The effect of tuberculosis control in China. Lancet, 2004, 364:417-422.

<sup>&</sup>lt;sup>2</sup> UNAIDS 2004 Report on the global AIDS epidemic. Geneva, Joint United Nations Programme on HIV/AIDS, 2004.

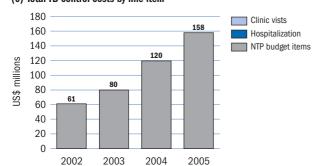
#### (a) NTP budget by source of funding



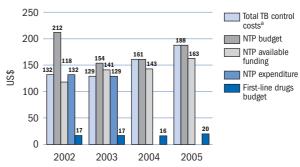
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>







<sup>&</sup>lt;sup>a</sup> Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. See Methods for further details.

system. Pilot initiatives to involve hospitals are in place and are showing encouraging results.

#### **Partnerships**

In addition to the government funding for TB control, funds are provided for TB control projects from outside sources, with technical assistance from WHO and KNCV. The World Bank/ DFID project provides funding for 16 project provinces. The Government of Japan provides funds for anti-TB drugs, microscopes and health promotion materials in 12 provinces. The GFATM has approved US\$ 25.4 million for the first two years of a five-year TB project, with initial funds disbursed to 24 provinces. Another GFATM project was approved in July 2004. The Damien Foundation Belgium supports TB control activities in Tibet (since 1995), Inner-Mongolia (since 2001) and Qinghai (since 2003). CIDA began

funding TB control activities through WHO in 2003, and the project now covers a population of 75 million in its second year. FIDELIS, run by IUATLD, is exploring new approaches to improve quality of DOTS services and increase case detection.

#### **Budgets and expenditures**

In line with plans to reach the global targets for case detection and treatment in 2005, the budget is projected to increase from around US\$ 100 million in 2002 and 2003 to US\$ 160 million in 2005. Due to a large funding gap, actual expenditures were only around US\$ 60 million in 2002.

Budgets from 2003 onwards have been substantially boosted by increased government funding, a new World Bank loan and successful applications to the GFATM. Despite this progress, a funding gap of US\$ 21 million still exists for 2005. This funding gap reflects faster than expected expansion of DOTS to new areas, greater than anticipated increases in case detection and plans to introduce additional initiatives to increase case detection in 2005 (e.g. tracing of patients reported through the general communicable disease reporting system, an increased number of sputum examinations at subcounty level and more IEC activities). Some of the funding gap may be filled by an increase in local government funding, but the extent to which this will occur is currently unclear. An increasing budget for first-line drugs is planned to meet the increase in treated cases, with the cost per patient treated remaining at about US\$ 20 for the period 2002-2005. If expenditures match budgets in 2004 and 2005, the total cost of TB control activities per patient treated will increase from about US\$ 130 in 2002 to US\$ 190 in 2005.

# Democratic Republic of the Congo

The Democratic Republic of the Congo, despite being among the poorest countries in the world, has made substantial progress in TB control in recent years; by 2004, DOTS services were available to approximately 80% of the population. While there is strong government support for the NTP, the provision of adequate TB services throughout the country has been hampered by a combination of difficulties. The country's health infrastructure has suffered in the past from an underdeveloped primary care system, lack of funds and resources as well as from the destructive effects of civil unrest and natural disasters. In spite of these constraints, TB case detection and cure rates have both improved steadily since the early 1990s, and the NTP hopes to reach the global targets by 2005. The likelihood of achieving these objectives has been greatly boosted by an award from the GFATM, as well as increased government funding, which have transformed the financial basis of TB control services and

will allow for the extension and strengthening of activities. Furthermore, the special problems posed by the epidemic of TB in people infected with HIV are being addressed in an expanding programme of collaborative TB/HIV activities.

#### **System of TB control**

The NTP was officially launched in 1980 (Programme National Antituberculeux Integré, PATI 1) and consists of a central unit, 20 provincial coordination centres, 777 TB diagnosis and treatment centres and a network of health posts (consisting of a nurse or health-care worker) distributed in 515 health districts. Better health coverage resulted from a health mapping exercise carried out in 2004, following which the number of health districts was increased from 306 to 515. TB services follow the expanded health network to improve access by providing services closer to where the patients live and to promote healthseeking behaviour.

The TB laboratory network consists of one NRL, which was significantly upgraded in preparation for the application to the GLC, 20 provincial laboratories implementing EQA and 800 district laboratories, giving 1 laboratory per 70 000 inhabitants. There are no microscopy services in any of the peripheral health posts.

#### **Surveillance and monitoring**

As a result of progressive expansion of DOTS services, coverage of approximately 80% was reached in 2004. The TB notification rate for both smearpositive and all forms of TB has increased over the past 20 years, partly as a result of improved case-finding and partly as a result of a rise in TB incidence linked to the spread of HIV. For the Democratic Republic of the Congo, as for some other countries in central Africa, the accuracy of the estimated case detection rate (63% in 2003) is uncertain. The treatment success rate was 78% for the 2002 cohort; 7% of patients died and 13% defaulted or were lost to follow-up after transfer to other treatment centres. Both of the latter indicators were high for patients undergoing re-treatment following relapse, failure or default; the relapse re-treatment success rate was 70%. High HIV prevalence, poor health infrastructure and large numbers of displaced persons contribute to this low treatment success rate. However, preliminary data suggest that the treatment success rate for the first quarter of 2003 was 81%. Improvement of the treatment success rate is a high priority for the NTP.

#### **Improving programme** performance

Revised TB control guidelines have been prepared (Programme National Antituberculeux Integré, PATI 4) and will be published soon. These guidelines, which include the introduction of the 6-month regimen for treatment, have already been used as the basis for training sessions on the progressive introduction of the new regimen.

#### PROGRESS IN TB CONTROL IN THE DEMOCRATIC REPUBLIC OF THE CONGO

#### **Indicators**

DOTS treatment success, 2002 cohort	78%
DOTS case detection rate, 2003	63%
NTP budget available, 2004	84%
Government contribution to NTP budget, including loans, 2004	5%
Government contribution to total TB control costs, including loans, 2004	64%
Government health spending used for TB control, 2004	NA

#### **Major achievements**

- Review of the national TB control guidelines (Programme National Antituberculeux Integré, PATI 4), including introduction of 6-month treatment regimen
- Extensive training at all levels, including initiation of more than 4000 community health workers
- Establishment of a TB/HIV coordinating body to coordinate activities of the National AIDS Control Programme and the NTP
- Improved capacity of laboratories, including provision of 800 microscopes

#### **Major planned activities**

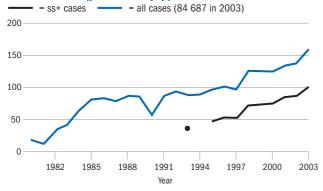
- Host external monitoring mission planned for February 2005
- Prepare five-year strategic plan (2006–2010)
- Implement a national drug resistance survey
- Update and revise NTP technical guidelines
- Expand collaborative TB/HIV activities, following recent award from the President's Emergency Plan for AIDS Relief

NA indicates not available.

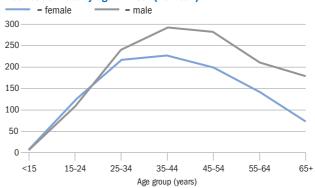
## **DEMOCRATIC REPUBLIC OF THE CONGO**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	52 771 230	DOTS coverage (%)	70	70	70	75
Global rank (by est. number of cases)	11	Notification rate (all cases/100 000 pop)	125	134	138	160
Incidence (all cases/100 000 pop/year)	369	Notification rate (new ss+/100 000 pop)	74	84	87	102
Incidence (new ss+/100 000 pop/year)	160	Detection of all cases (%)	39	40	39	44
Prevalence (all cases/100 000 pop)	564	Case detection rate (new ss+, %)	53	58	57	63
TB mortality (all cases/100 000 pop/year)	81	DOTS case detection rate (new ss+, %)	53	58	57	63
TB cases HIV+ (adults aged 15-49, %)	21	DOTS case detection rate (new ss+)/coverage (%)	76	83	81	84
New cases multidrug resistant (%)	1.5	DOTS treatment success (new ss+, %)	78	77	78	_

#### Notification rate (per 100 000 pop)



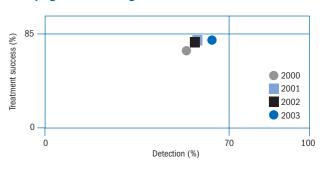
#### Notification rate by age and sex (new ss+)<sup>b</sup>



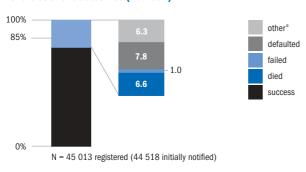
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

### **DEMOCRATIC REPUBLIC OF THE CONGO**

A new strategic plan will be prepared for the period 2006-2010.

The supply of anti-TB drugs is adequate, thanks to a second GDF grant approved in 2004 for another threeyear period. However, the country's drug policy and system of drug management need to be revised to ensure sustainable supply and better drug distribution, with regular reporting on drug stocks at every level.

A national drug resistance survey is planned for 2005. The protocol has been finalized and implementation should start soon. A national policy for the diagnosis and treatment of MDR-TB is being developed.

Extensive training activities were carried out in 2003 at all levels, including initiation of more than 4000 community health workers.

Supervision of laboratory, medical, financial and administrative functions is carried out on a regular basis, but the time devoted to each visit is inadequate and only half of the planned visits were carried out in 2003. Nonetheless, supervision has resulted in improvements in the procedures for recording and analysis of observations made during patient visits. There has also been a striking improvement in the management and flow of funds because of a revision of financial and administrative procedures under the guidance of a newly-recruited finance officer at central level.

Data collection is more reliable than in the past, although delays are experienced and data collection forms remain unnecessarily complicated.

The award of a GFATM grant has greatly increased the funding available for TB control in 2004, making it possible to address the problems of staffing, training, medical supplies and equipment. With improvements in the facilities for diagnosis and patient care, case detection and cure rates should continue to rise in the coming years.

Three areas where programme performance particularly needs to be improved are laboratory services for culture and DST, TB/HIV coordination and links with other health-care providers.

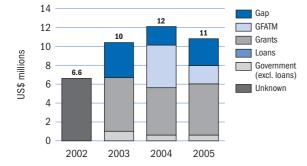
#### Diagnostic and laboratory services

The quality of DOTS implementation relies on an effective laboratory network. New equipment was installed during 2003 and 2004 in most peripheral laboratories, and EQA is implemented in most laboratories. EQA included external laboratory supervision in half of the provinces in 2003. Onthe-spot slide reading is carried out in half of the districts visited during the external visits. The link between the NRL and the NTP needs to be strengthened to ensure effective coordination. The central laboratory is poorly equipped and the quality of slide reading is poor in laboratories where microscopes are old and need to be replaced.

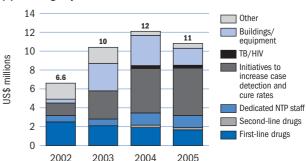
#### TB/HIV coordination

More than 20% of adult TB patients are infected with HIV (WHO estimate). The National AIDS Control Programme and the NTP have established a TB/ HIV coordinating body in 2003, and one NPO has been recruited to support these activities. Since 2002, several DOTS centres have started collaborative TB/HIV activities in Kinshasa, with financial and technical support from WHO, MSF and the World Bank. Following a recent award from the President's Emergency Plan for AIDS Relief, collaborative TB/HIV activities will be expanded in 2005.

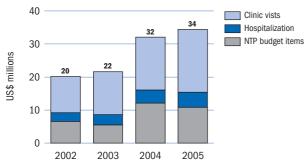
#### (a) NTP budget by source of funding



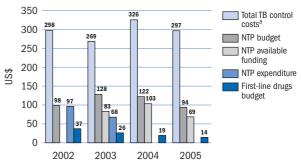
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details

### **DEMOCRATIC REPUBLIC OF THE CONGO**

#### Links with other health-care providers

The NTP is collaborating with general hospitals, medical colleges, and military and police health services; specialist TB hospitals and the prison health-care service do not implement DOTS. Limited formal involvement of the private sector has started with the training of private physicians in DOTS activities. These providers are also represented in some provincial and national task force meetings.

#### **Partnerships**

The Democratic Republic of the Congo benefits from several financial and technical partnerships for TB control. The GFATM is a principal source of funds, and the national budget for TB control has increased in 2004. Additional support from the Government of Belgium was agreed in 2004. UNDP, the principal recipient of the GFATM funds, has established four public and private recipients for the TB control proposal (the NTP, La Ligue Nationale

Antituberculeuse et Antilépreuse du Congo, DFB and the National School of Public Health). Through this arrangement, other partners have been included.

#### **Budgets and expenditures**

The NTP budget has been between US\$ 10 million and US\$ 12 million for the years 2003-2005, compared with about US\$ 7 million in 2002. Almost all of the available funding for 2003-2005 comes from grants, with the government contributing less than 10% of the NTP budget (no breakdown of the budget by funding source in 2002 has been provided to WHO). This makes the Democratic Republic of the Congo largely dependent on external financing. Despite the approval of the GFATM grant in round 2, and sustained commitment from other donors, an important funding gap remains: US\$ 2.9 million (26% of the NTP budget) in 2005.

The budget for first-line drugs has

decreased from US\$ 2.5 million in 2002 to US\$ 1.7 million in 2005, reducing the first-line drug budget per patient treated from US\$ 37 to US\$ 14. In contrast, the budget for initiatives to increase case detection and cure rates has grown from US\$ 1.3 million in 2002 to US\$ 5 million in 2005 and is now the largest single budget item.

The total cost of TB control, which includes the cost of clinic visits and dedicated TB hospital beds, in addition to the NTP budget, is projected to increase from an estimated US\$ 20 million in 2002 to US\$ 34 million in 2005, in line with anticipated increases in the numbers of patients to be treated (the cost per patient treated is around US\$ 300 per patient in both 2002 and 2005). The government contribution to the total cost of TB control is much larger than to the NTP budget, varying from 60% to 76% of total costs.

# **Ethiopia**

Ethiopia has given priority to TB, HIV/ AIDS and malaria prevention and control for more than a decade. The DOTS strategy is being implemented in most districts, and almost all hospitals and health centres provide DOTS services. However, basic health services are not yet accessible to about 40% of the population, and intensive efforts are being made to ensure better access throughout the country. Health facilities suffer from a high turnover of staff to deliver TB services; this constraint is being addressed through a comprehensive HRD plan and training programmes. Available data suggest that the incidence of TB has risen in recent years, partly as a result of the impact of the HIV/AIDS epidemic. Special efforts are being made to address the needs of TB patients coinfected with HIV coinfection. Ethiopia has carried out its first national drug resistance survey and found that the rate of MDR-TB is low. The country has successfully maintained an uninterrupted supply of anti-TB drugs for several years. Approval of a grant from the GFATM opened up additional possibilities to expand and improve TB control services in 2004 and 2005.

#### System of TB control

The health policy in Ethiopia, dating from 1993, gives priority to the control of communicable diseases, including TB, HIV/AIDS and malaria. The health system is being progressively decentralized under the country's primary health-care strategy. Recently, a four-tier health-care delivery structure was established to implement this policy. The primary health-care unit is the basic level of health care for Ethiopia and consists of a health centre with five satellite health posts, each serving 5000 people. In 2005, this network will be extended by the addition of two health extension workers for each subdistrict (kebelle). The health system also includes district (woreda), regional and specialized hospitals, serving 250 000, 1 000 000 and 5 000 000 people, respectively. During the past year, further decentralization to the woredas in major regions of the country has led to an increase in the transfer of health personnel from regions and zones to woredas and a decreasing role of the zone in TB control activities.

In 1994, the NTP (known locally as

the TB and Leprosy Prevention Control Team) was established. Since 2000, it has been part of the Disease Prevention and Control Department of the Federal MoH. In 1996, a Project Development Plan (PDP), designed to support TB control through the NTP for five years, was signed by the Government of Ethiopia, WHO and KNCV. In 2001, this plan was extended for an additional year.

The laboratory services in Ethiopia include one NRL, regional reference laboratories in some regions and peripheral laboratories.

#### **Surveillance and monitoring**

The steady rise in case notifications since 1993 is because of increasing DOTS coverage, improved reporting and the impact of HIV/AIDS. While the relative contributions of these three factors are uncertain, it has been assumed that the national smear-positive case detection rate by the DOTS programme has remained constant at around 36%, while incidence has increased. The case detection rate within DOTS areas was only 38% in 2003, due largely to the important difference between DOTS coverage as defined in this report (95%) and the proportion of the population thought to have access to health services of any kind, including for TB (50%). The proportion of notified cases diagnosed as smear-positive is low in Ethiopia, and has stayed within the range 27-35% during the period 1995-2003.

Despite the moderately high prevalence of HIV infection (4.4% of adults aged 15-49 years in 2003), it remains difficult to explain the extraordinary proportion of cases that are reported as extrapulmonary TB (>34% in 2003, regional variation 29-54%). The vast majority of extrapulmonary cases are reported as lymph node TB; this phenomenon is currently being investigated through a large operational research study in six sites in four

Treatment success among new patients was only 76% in the 2002

#### **PROGRESS IN TB CONTROL IN ETHIOPIA**

#### **Indicators**

DOTS treatment success, 2002 cohort 76% DOTS case detection rate, 2003 36% NTP budget available, 2004 100% Government contribution to NTP budget, including loans, 2004 8% Government contribution to total TB control costs, including loans, 2004 31% Government health spending used for TB control, 2004 10%

#### **Major achievements**

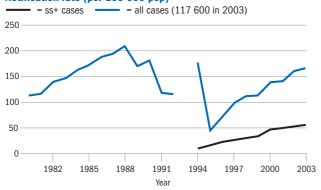
- Provision of DOTS services by 98% of hospitals and health centres
- Uninterrupted drug supply for several years
- Strong HRD plan with up-to-date training material and methodology
- Drug resistance survey completed with relatively low MDR-TB rate reported

#### **Major planned activities**

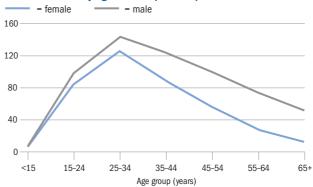
- Commence collaborative TB/HIV activities in pilot sites as well as in hospitals scheduled to provide ART
- Involve communities and private providers in TB control
- Conduct major training activities in all regions and woredas following the recent HRD plan

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	70 678 002	DOTS coverage (%)	85	70	95	95
Global rank (by est. number of cases)	7	Notification rate (all cases/100 000 pop)	139	141	160	166
Incidence (all cases/100 000 pop/year)	356	Notification rate (new ss+/100 000 pop)	47	49	53	56
Incidence (new ss+/100 000 pop/year)	155	Detection of all cases (%)	45	43	47	47
Prevalence (all cases/100 000 pop)	533	Case detection rate (new ss+, %)	35	35	36	36
TB mortality (all cases/100 000 pop/year)	79	DOTS case detection rate (new ss+, %)	35	35	36	36
TB cases HIV+ (adults aged 15-49, %)	21	DOTS case detection rate (new ss+)/coverage (%)	41	50	38	38
New cases multidrug resistant (%)	2.3	DOTS treatment success (new ss+, %)	80	76	76	_

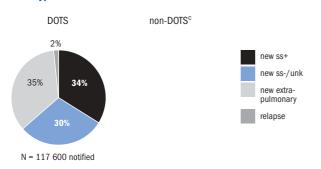
#### Notification rate (per 100 000 pop)



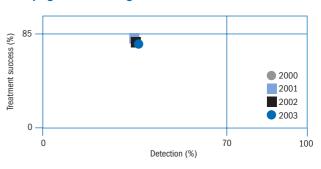
#### Notification rate by age and sex (new ss+)<sup>b</sup>



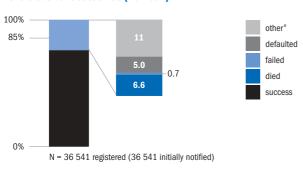
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- <sup>e</sup> "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

cohort, considerably lower than Ethiopia's maximum of 80% reported for 2000. However, this decrease is explained by the fact that the NTP now includes all notified cases in the analysis. Among new patients who were registered for treatment in 2002, the outcome of treatment is not known for 10% following transfer between treatment units; 7% died and 17% completed treatment without evidence of smear conversion.

#### Improving programme performance

In recent years, the high turnover of staff involved in TB control and the effects of decentralization have resulted in a workforce that is not well trained in the principles of TB control. HRD in the NTP therefore received special emphasis in 2003, with the completion of a comprehensive HRD plan, production of a first edition of TB/leprosy training modules and associated materials for all levels of staff involved in TB control, and the creation of a pool of 53 competent TB and leprosy trainers distributed over all regions. Detailed regional training plans for 2005 have been drafted and funding for their implementation secured. The first phase of the plans focuses on in-service training; the second phase will also involve incorporation of TB control principles in the pre-service curricula.

Anti-TB drugs and laboratory supplies are procured by the Pharmaceutical Administration and Supplies Services of the Federal MoH using international competitive bidding, with funding from the GFATM. Despite delays in the procurement process, there has not been any interruption to the availability of drugs, mainly because of the continued maintenance of a one-year buffer stock. NTP training increasingly includes pharmacy staff. Four-drug FDCs have been introduced for the intensive phase of treatment for new patients.

Although almost 95% of the woredas have at least one health facility providing DOTS services, more than half of the smaller health stations/posts do not provide directly observed TB treatment. Covering all

these units is one of the main objectives of the NTP, but implementation has been constrained by a shortage of staff for monitoring and supervision as well as a delay in HRD.

Ethiopia's first drug resistance survey is close to completion, with preliminary results indicating 1.7% MDR among new cases, somewhat lower than the WHO estimate of 2.3%.

Three other areas in which programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers and the community.

#### Diagnostic and laboratory services

All laboratories are supplied with microscopes and reagents by the NTP, and staff are included in TB-related training activities. A system of quality assurance is in place, but implementation is weak. The NTP developed and issued a national laboratory manual for smear microscopy in 2002, which will be revised and re-edited in 2005.

#### TB/HIV coordination

A national TB/HIV coordinating body has been established and specific terms of reference developed. The committee includes the Federal MoH, academia, bilateral donors and the technical partners of the TB and HIV programmes. Nine pilot sites have been selected to pilot collaborative TB/HIV activities under the guidance of the committee. A national TB/HIV surveillance plan is being finalized. TB/HIV activities are managed by a national TB/HIV coordinator (WHO) based at the Federal MoH.

A national TB/HIV orientation workshop and various training courses have been conducted for the staff of pilot sites, in management of TB and other opportunistic infections in PLWHA and in VCT. Guidelines have been developed for the use of isoniazid preventive therapy in PLWHA infected with M. tuberculosis and for the use of co-trimoxazole preventive therapy in HIV-infected TB patients. Isoniazid (through the GDF), cotrimoxazole and HIV test kits (both through the Federal MoH) have been distributed. Ethiopia hosted, facilitated and participated in three major international activities: the meeting of the TB/HIV Global Working Group, TB/ HIV Surveillance International Workshop (CDC/WHO) and two global TB/ HIV managers training courses (WHO/ GLRA).

#### Links with other health-care providers

Observations during monitoring and supervision as well as a small scale study in Addis Ababa have shown that many patients are managed in private clinics. Patient management is generally limited to diagnosis since, officially, anti-TB drugs in Ethiopia are available only in government health facilities. Most patients in whom TB is diagnosed in the private sector are referred to public health centres for registration and treatment. However, anti-TB drugs have been shown to circulate illegally, and treatment of an unknown number of patients is initiated in the private sector, disregarding national treatment guidelines. A pilot project is planned in Addis Ababa so that private providers will be increasingly involved in training activities as well as laboratory quality assurance activities.

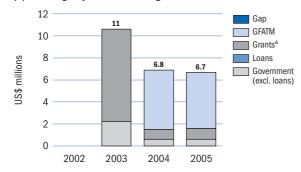
#### Links with the community

Given the sparse distribution of health facilities, and the consequent limited access to DOTS services in Ethiopia, plans are under way to involve the community in TB control. With GFATM funding, pilot projects will start in four districts of four regions. National guidelines for community involvement in DOTS and training modules and materials have been developed and distributed.

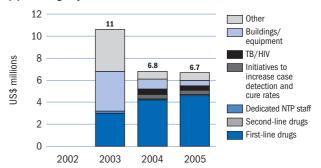
#### **Partnerships**

For many years, the NTP has been consistently supported by the Royal Netherlands Embassy, GLRA and WHO. More recently, support has been received from the GFATM, with a large grant approved in the first round of applications. Other partners are CDC and USAID. MSF Belgium is providing support in the Somali Region, but this will be discontinued in 2005 when the regional health bureau assumes responsibility for the region.

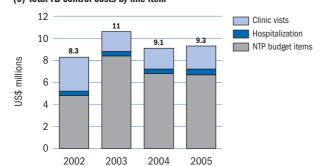
#### (a) NTP budget by source of funding



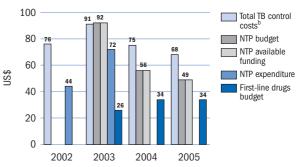
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>b</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



- <sup>a</sup> The 2003 budget data provided to WHO did not separate the GFATM contribution from other grants.
- <sup>b</sup> Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

#### **Budgets and expenditures**

The total NTP budget was US\$ 11 million in 2003, and lower at around US\$ 7 million in both 2004 and 2005 (US\$ 92 and US\$ 49 per patient in 2003 and 2005, respectively). The relatively high total in 2003 was a result of a large budget for capital investments (included in the "buildings and equipment" category) as well as a large budget in the "other" line item category. In practice, only 53% and 34%, respectively, of these budgets were spent. Following approval of a

GFATM grant in round 1 of US\$ 11 million for the first two years, Ethiopia has not reported any funding gaps for 2003–2005. Grants, including from the GFATM, represent more than 90% of the NTP budget in 2004 and 2005, making the NTP highly dependent on external funding. Programme sustainability is a concern, as grants support key areas such as first-line anti-TB drugs. In addition, continuous funding from the GFATM will depend on NTP performance during the first two years of the grant.

As the expected number of TB patients to be treated is increasing, the first-line drug budget is steadily expanding to reach US\$ 4.6 million in 2005, and remains the largest budget item.

The total TB control cost per patient (including the estimated costs of bed-days and clinic visits as well as the costs reflected in NTP budgets and expenditures) has remained relatively low, varying from US\$ 68 to US\$ 91 between 2002 and 2005.

# India

India, the country with the greatest burden of TB, is also the country where the most dramatic advances are being made in DOTS expansion. Thanks to a massive recent scale-up, TB services were available to some 67% of the population by 2003, and full nationwide DOTS coverage is planned for 2005. During 2003 alone, some 250 million additional people were included and treatment provided to more than 900 000 TB patients. At the same time, there has been a considerable improvement in the level of case detection, with India making a greater contribution than any other country to the global increase in casefinding since 2000. Mobilizing all public sector health-care providers, especially medical colleges, as well as many private and other health-care providers outside the government service, has been important to achieving such swift progress, and successful efforts continue to increase their involvement. Maintaining quality during rapid growth is a priority, while addressing the urgent need for addi-

tional staff and laboratory support for the expanded services. The Indian TB control programme is outstanding not only because of the recent progress but also because it has been made at a lower than predicted cost.

#### **System of TB control**

India's Revised National TB Control Programme (locally RNTCP, hereafter NTP) was introduced on a pilot scale in 1993 and, after a period of pilot testing, was formally launched by the government in 1997. By mid-1998, the programme had expanded to serve some 20 million people. There followed a phase of rapid expansion from late 1998 so that, by 2003, the areas covered by the DOTS strategy included 778 million people (around 67% of the population).

The laboratory network currently comprises 3 national reference laboratories (these are the LRS Institute of TB and Respiratory Diseases, Delhi; the National TB Institute, Bangalore and the TB Research Centre, Chennai), 15 state laboratories, 522 district laboratories and nearly 9000 peripheral NTP-designated microscopy centres. The national reference laboratories train state-level laboratory staff, and monitor and oversee the state laboratories. The state laboratories train district laboratory and supervisory staff, and monitor and oversee the peripheral microscopy centres; some of them perform culture and drug susceptibility testing. Sputum smear microscopy services are provided by the district and peripheral level microscopy centres.

#### **Surveillance and monitoring**

Coverage was extended by 250 million people during 2003, with more than 900 000 patients placed on DOTS treatment during that year. Based on this remarkable progress, it is planned to cover a total of 850 million people by the end of 2004 and to reach 100% coverage by October 2005. The estimated smear-positive incidence was revised on the basis of a three-year national tuberculin survey that was completed during 2003. There was a striking improvement in the DOTS case detection rate in 2003, with an estimated 47% of all new smear-positive cases in the country detected by the NTP compared with 31% in 2002, and 69% detected in the areas already covered by the DOTS programme. This increase in case detection represents 39% of the increase in cases detected by DOTS programmes worldwide, and India has made a larger contribution than any other country to the acceleration in global case-finding observed since 2000. The reported treatment success has also increased over the past three years (to 87% for 2002), despite the rapid growth of the national DOTS cohort (to more than 37 000 new smear-positive patients in 2003).

In contrast to the upward trend in case notifications seen in the NTP, the notification rate of all TB cases, from all sources in India, has been falling gradually since 1992. It remains unclear whether this downward trend

#### **PROGRESS IN TB CONTROL IN INDIA**

#### **Indicators**

DOTS treatment success, 2002 cohort	87%
DOTS case detection rate, 2003	47%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	74%
Government contribution to total TB control costs, including loans, 2004	86%
Government health spending used for TB control, 2004	2%

#### **Major achievements**

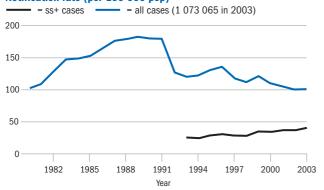
- Expansion of DOTS to cover an additional 250 million population during 2003
- Scaling up of PPM DOTS project in 12 sites
- GFATM round 1 activities started and round 2 agreement signed
- Involvement of medical colleges through national, subnational and state task
- Involvement of health facilities under other ministries
- Publication of new guidelines on EQA and development of a DRS protocol for
- Development of guidelines for management of paediatric TB

#### **Major planned activities**

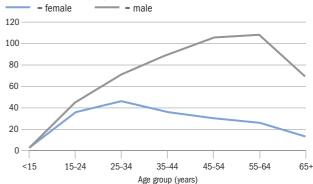
- Prepare for DOTS expansion in remaining states (laboratories, human resource, procurement) - entire country to be covered by October 2005
- Sustain quality of existing DOTS services by implementing a revised supervision and monitoring strategy
- Continue human resource capacity building through revision of all training material

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	1 065 462 272	DOTS coverage (%)	30	45	52	67
Global rank (by est. number of cases)	1	Notification rate (all cases/100 000 pop)	110	105	101	101
Incidence (all cases/100 000 pop/year)	168	Notification rate (new ss+/100 000 pop)	34	37	38	41
Incidence (new ss+/100 000 pop/year)	75	Detection of all cases (%)	65	63	60	60
Prevalence (all cases/100 000 pop)	290	Case detection rate (new ss+, %)	46	50	50	54
TB mortality (all cases/100 000 pop/year	33	DOTS case detection rate (new ss+, %)	12	24	31	47
TB cases HIV+ (adults aged 15-49, %)	5.2	DOTS case detection rate (new ss+)/coverage (%)	42	53	60	69
New cases multidrug resistant (%)	3.4	DOTS treatment success (new ss+, %)	84	85	87	_

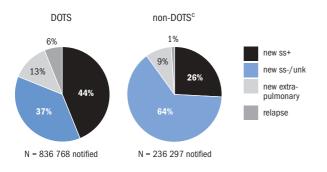
#### Notification rate (per 100 000 pop)



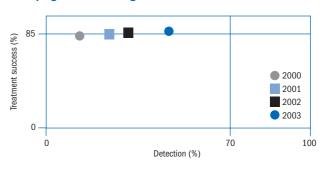
#### Notification rate by age and sex (new ss+)<sup>b</sup>



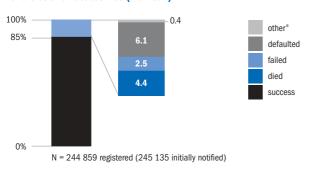
#### **Case types notified**



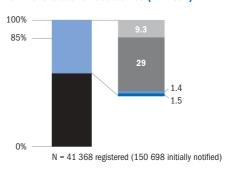
#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)



ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\rm e}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

reflects a real decrease in incidence, or improvements in diagnosis (eliminating false-positives). National data for the years up to 2003 do not yet provide evidence that the NTP has reduced incidence and prevalence, although it is clear that there are significantly fewer deaths among cases notified (18 deaths averted per 100 patients treated are reported at www.tbcindia.org). The epidemiological evidence for impact is most likely to come from areas where the programme has been operating for longest and where the implementation of DOTS has been studied most intensively, notably in the "model DOTS project" being carried out by the Tuberculosis Research Centre in Chennai.

#### Improving programme performance

Maintaining the quality of TB services is crucial as the programme moves towards full coverage, and this will be a major challenge in the coming years. To address it, several activities are under way, including the development of guidelines for the management of paediatric TB and the introduction of a revised supervision and monitoring strategy with detailed indicators for activities at all levels.

The most important constraint to the extension of quality TB services is the shortage of staff to manage the rapidly expanding programme, particularly at central and state levels. To improve this situation, additional technical staff have been recruited to assist the NTP manager, the limits on hiring contractual laboratory technicians have been relaxed and efforts are being made to achieve an adequate distribution of laboratory technicians to states where laboratories are understaffed. Subdistrict contracted laboratory supervisors have greatly contributed to the success of the programme, and efforts will be made to sustain capacity over the next few years. Further political commitment at the state level is needed to ensure that the programme is fully staffed with stable management. The capacity of current staff will be increased through training programmes run in part by expert consultants.

It is estimated that 3.4% of previously untreated TB cases are multidrug resistant. Currently, the NTP does not supply second-line drugs for MDR-TB patients. There are plans to build capacity at the state level for DRS and DOTS-Plus. Although MDR-TB patients are not treated under the NTP, second-line drugs are widely available and used by many practitioners, both public and private.

During 2005, priority will be given to preparing the remaining districts for DOTS implementation. The preparatory activities include the improvement of laboratories and stores, recruitment, relocation and training of staff, and procurement of equipment and supplies. Some of the districts are in areas where operations are difficult to access and where intensive monitoring will be required. Funds secured through the GFATM will be used to expand the programme to cover 56 million population in all 47 districts of the three newly-created states of Chhattisgarh, Jharkhand and Uttaranchal (round 1), and 110 million population in 56 districts of the states of Bihar and Uttar Pradesh (round 2). In addition, GFATM funds will be used to maintain DOTS coverage in 110 million population in the states of Andhra Pradesh and Orissa (round 4).

Diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers and the community are three priority areas to improve programme performance.

#### Diagnostic and laboratory services

The TB laboratory network is being strengthened to meet the needs of the expanding programme, by upgrading existing laboratories, creating new microscopy centres and establishing EQA. Based on new international guidelines, an EQA system for sputum microscopy was adopted at the beginning of 2004 for the NTP smear microscopy laboratory network, and includes a random blinded crosscheck of routine slides each month. Panel testing at the district level is done by the state laboratories once a year. Currently, only NTI Bangalore and TRC Chennai are quality assured for both culture and drug susceptibility testing. The national reference laboratories participate in annual proficiency testing coordinated by the Antwerp and Chennai supranational laboratories. Building capacity for DST at the intermediate laboratory level has started in two state-level laboratories in 2004 and is planned for two others by the end of the 2004; a national plan has also been developed to systematically perform DRS surveys in large states of the country.

#### TB/HIV coordination

An estimated 5.1 million people are infected with HIV in India. HIV is likely to have a significant impact on the TB epidemic in the six states where the prevalence of HIV is greater than 1%, namely Andhra Pradesh, Karnataka, Maharashtra, Manipur, Nagaland and Tamil Nadu. The prevalence of HIV in TB patients has been measured in a number of tertiary care hospital settings, reaching 25% in one such hospital in Pune, Maharashtra, in 2001. However, the results from such studies are not representative of the HIV levels in TB patients in India as a whole. In 2004, HIV surveillance in TB patients has started in four districts in the six high-prevalence states, using a more representative sampling methodology.

Coordination of HIV and TB services has been prioritized in the six states with the highest HIV prevalence. HIV and TB staff have been cross-trained, referral linkages between the district VCT centres of the HIV programme and microscopy centres of the DOTS programme established and a surveillance system to document cross-referrals is currently in the pilot phase. Joint HIV/TB coordination committees will be established at the national and state levels with support from GFATM, and a referral system will be created at the subdistrict level between the existing NTP infrastructure and the VCT centres.

#### Links with other health-care providers

Private and other health-care providers, including NGOs and medical colleges, play an extremely important role in DOTS implementation in India. The Government of India has formulated and published schemes to promote participation of NGOs (2001) and private practitioners (2002) in implementing DOTS. During the past few years, several local initiatives have emerged in both urban and rural settings; the NTP has provided drugs free of charge and has taken responsibility for supervision and monitoring of laboratory and treatment services. The evaluated initiatives have shown an increase in case notification between 3% and 30%. Most projects have also achieved treatment success greater than the programme target of 85%. Encouraged by the success of these early experiments, the NTP, in collaboration with WHO, has embarked on scaling up PPM DOTS in 14 cities across the country. The strategy is to offer technical support to the city TB control programmes to facilitate partnership development through a fulltime PPM consultant assisted by two field supervisors. Future expansion of PPM DOTS will link all public, corporate, voluntary and private individual and institutional providers to the NTP. The programme has adapted the existing recording and reporting system in order to evaluate the PPM-DOTS activities. Monitoring during the two initial quarters showed that PPM-DOTS

providers other than those under the DoH contributed 39% of the cases detected under DOTS in the pilot cities. Public and private medical colleges alone accounted for 18%.

#### Links with the community

Community volunteers are used as DOT providers all over the country. In some parts, there has been effective involvement of the community through patient—provider—community meetings. IEC campaigns also involve the community at large, especially during events such as World TB Day.

#### **Partnerships**

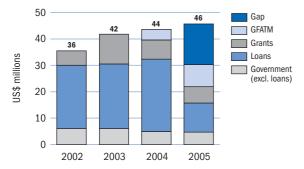
WHO has helped to establish a network of more than 85 field consultants and provides technical support for all aspects of the programme. These field consultants work with the programme managers at state and district level and report directly to the central unit of the NTP. India receives anti-TB drugs for 240 million of its population through the GDF. Financial partners include CIDA, DANIDA, DFID, GFATM, USAID and the World Bank.

#### **Budgets and expenditures**

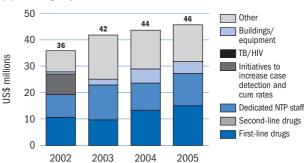
In line with the rapid DOTS expansion taking place in India, the NTP budget has increased from US\$ 36 million in 2002 to a projected US\$ 46 million in 2005. Most funding is provided by the government, through a World Bank credit and domestic government revenue. With an increase in funding from the GFATM, grants will provide about 30% of the budget in 2005. No budget gaps were reported for 2002-2004; although there is currently a funding gap of US\$ 15 million for 2005, it is expected that this will be filled by a combination of additional grants and a new World Bank credit.

The largest budget items are first-line drugs and dedicated staff, which together account for more than 50% of the total budget in each year 2002–2005. The budget per patient treated has remained stable as DOTS has expanded, at about US\$ 35–40. The same is true of total TB control costs (which include visits to health facilities and expenditures on dedicated TB hospital beds in addition to items covered by the NTP budget). The total TB control cost per patient treated has consistently remained at about

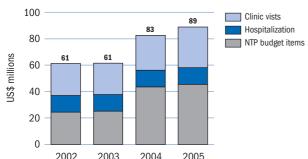
#### (a) NTP budget by source of funding



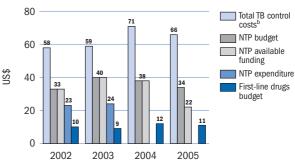
#### (b) NTP budget by line item<sup>a</sup>



#### (c) Total TB control costs by line item<sup>b</sup>



### (d) Per patient costs, budgets, available funding and expenditures



- a TB/HIV collaborative activities and initiatives to increase case detection and cure rates are not budgeted separately, and are thus included under other budget lines
- b Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

### **INDIA**

US\$ 60-70, as total TB control costs have increased from about US\$ 60 million in 2002 to a projected US\$ 89 million in 2005; these figures may be overestimates because they assume that 75% of all DOT is undertaken at health facilities and that about 10 000 dedicated hospital beds are still being used for TB patients. In practice, community workers or volunteers may provide DOT to more than 25% of patients, at no cost to the health system, and increasing numbers of hospital beds previously dedicated to TB patients are being reallocated to other uses. A costing study will be undertaken in 2005 to further refine these estimates. While progress has been made at the planned rate, actual expenditures were lower than budgets in both 2002 and 2003, so the cost of DOTS expansion has been lower than anticipated.

# Indonesia

Indonesia is entering a phase of rapid and comprehensive acceleration of its TB activities thanks to a substantial increase in funding for 2004 and 2005. With a fully-funded budget, many opportunities are being taken to improve surveillance, case detection and laboratory services, to extend the involvement of other health-care providers in DOTS and to improve TB/HIV coordination. A population-based TB prevalence survey was carried out in 2004; the data will provide a more accurate estimate of the national burden of TB, and provide a basis for assessing the future impact of the NTP on the TB epidemic. Although Indonesia has achieved a high level of DOTS coverage (98%), this has not yet been matched by high levels of case detection because of several factors, including a backlog of staff to be trained, suboptimal laboratory support and the lack of effective links with the hospi-

tal sector and private practitioners. Substantial improvements in the weaker areas of the programme should accrue from the greatly increased investment in the NTP.

#### **System of TB control**

In the decentralized primary healthcare system, TB control is offered through the district health services. District populations range from under 10 000 to more than 2 million, with the majority between 50 000 and 150 000.

Indonesia does not yet have a designated NRL for TB. A fully functioning national TB laboratory network is currently being developed. The existing laboratory network, which is not formally linked with the NTP, consists of microscopy health centres and independent health centres where trained laboratory staff carry out smear diagnosis. Provincial health

laboratories provide some assessments of the quality of smear microscopy, and perform culture and drug susceptibility testing on request.

#### **Surveillance and monitoring**

Since the burden of TB has been estimated from old (>20 years) and possibly unreliable data, Indonesia carried out an important national disease prevalence survey during 2004. Analysis of the survey data was still in progress in January 2005, and it is not yet clear whether the best estimate of smear-positive prevalence for 2003 will be significantly different from the WHO estimate of 295 per 100 000 population. Because TB cases have been reported with variable effort and consistency since 1980, the notifications over time give no indication of the underlying trend in incidence. However, the higher notification rates among older men suggest that the epidemic could be in slow decline. The national HIV infection rate remains low (0.1% in adults aged 15-49 years in 2003), but HIV appears to be generating more TB cases among young adults in some parts of Java and Papua.

The very high reported DOTS coverage (98% since 2000) has not been matched by high rates of case detection, although the smear-positive case detection rate has increased markedly between 2000 (19%) and 2003 (33%). Optimizing the functional capacity of health centres plus improved collaboration between the NTP, lung clinics and a limited number of public and private hospitals contributed to this success, and further strengthening of these links is needed. The NTP DOTS programme has been recruiting smearnegative and extrapulmonary cases faster than smear-positive cases since 1995. This may be a result of the heavy reliance on X-ray diagnosis in smear-negative patients, particularly in lung clinics and lung hospitals (involvement since 2002). Among new smear-positive patients, treatment success exceeded the 85% target in

#### **PROGRESS IN TB CONTROL IN INDONESIA**

#### Indicators

DOTS treatment success, 2002 cohort	86%
DOTS case detection rate, 2003	33%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	57%
Government contribution to total TB control costs, including loans, 2004	63%
Government health spending used for TB control, 2004	5%

#### **Major achievements**

- Strengthening of management capacity by placing staff at central and provincial level
- Step-wise and cascade training as part of human resource development
- Detailed planning and budgetary exercises conducted at district level for smooth disbursement of donor funds
- Improved supervision and monitoring from central and provincial level
- Involvement of public chest clinics and limited public and private hospitals in PPM DOTS

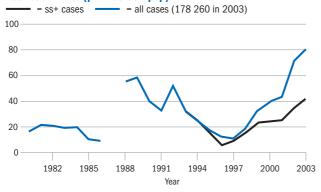
#### **Major planned activities**

- Expand hospital DOTS linkage projects in a phased manner in all provinces and
- Develop and implement an advocacy and communications framework for sustaining political commitment and increasing case detection rate
- Strengthen laboratory network and cooperation between NTP and laboratory directorates at all levels, through joint planning, supervision and monitoring activities at district level
- Accelerate HR development activities by accelerating the training backlog of health-centre staff and by training staff from hospitals and other sectors

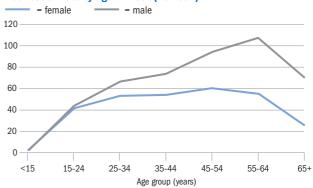
## **INDONESIA**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	219 883 460	DOTS coverage (%)	98	98	98	98
Global rank (by est. number of cases)	3	Notification rate (all cases/100 000 pop)	40	43	71	81
Incidence (all cases/100 000 pop/year)	285	Notification rate (new ss+/100 000 pop)	25	25	35	42
Incidence (new ss+/100 000 pop/year)	128	Detection of all cases (%)	14	15	25	28
Prevalence (all cases/100 000 pop)	675	Case detection rate (new ss+, %)	19	20	27	33
TB mortality (all cases/100 000 pop/year)	65	DOTS case detection rate (new ss+, %)	19	20	27	33
TB cases HIV+ (adults aged 15-49, %)	0.5	DOTS case detection rate (new ss+)/coverage (%)	19	20	28	34
New cases multidrug resistant (%)	0.7	DOTS treatment success (new ss+, %)	87	86	86	_

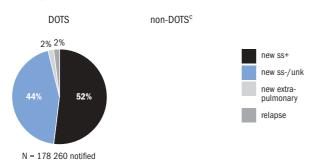
#### Notification rate (per 100 000 pop)



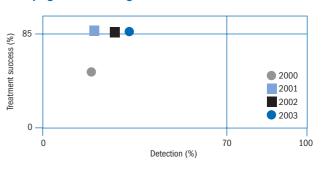
#### Notification rate by age and sex (new ss+)<sup>b</sup>



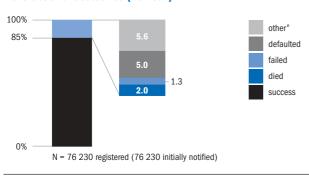
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- e "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

the 2002 cohort. Treatment results have been consistently good since the NTP began to evaluate outcomes comprehensively in 2000, although many patients complete treatment without evidence of cure (15% in 2002). There is no evidence yet that DOTS is reducing the burden of TB in Indonesia, but the 2004 prevalence survey data will give a baseline against which performance can be assessed towards the end of the decade.

#### **Improving programme** performance

The decentralization of health-care delivery has unfortunately had a negative effect on human resource capacity and development. Constraints include a high rotation of staff and hiring restrictions. As of December 2003, only 34% of health centre staff were adequately trained. Steps are being taken to alleviate this situation, and Indonesia was approved for additional funding through ISAC, which will help to reduce the training backlog by intensifying activities through mobile "master trainer's teams". As part of HR development, management capacity has been strengthened at the central and provincial levels during 2004, leading to a considerable improvement in supervision and monitoring by staff at these levels. Closer collaboration between the central, provincial and district health authorities is having a positive impact on TB control activities.

As a result of increased donor support and funding, Indonesia has carried out detailed planning and budgetary exercises at the district level for efficient disbursement of new funds.

Drug resistance surveillance has not yet been instituted in Indonesia. However, laboratory facilities at Surabaya have been upgraded because this laboratory will be used as the reference laboratory for future drug resistance surveys. Limited surveys in Jakarta have found MDR-TB in more than 4% of previously untreated cases; a fully representative survey is needed to determine whether this situation prevails throughout the country (the national WHO estimate is 0.7%). A survey in Central Java is planned for early 2005. There is no national policy

for the management of MDR-TB, and pulmonologists treat MDR-TB cases on an individual basis. Some of the second-line drugs are produced in the country.

#### Diagnostic and laboratory services

The link between TB laboratories and the NTP remains weak but will be made stronger with the establishment of a central laboratory working group and an NRL. A national assessment caried out in November 2004 evaluated the current laboratory services and will be used to guide planning for future improvements. There is also a need to improve and strengthen the EQA system. Priorities should include training of laboratory staff and preparation of a plan and timetable to carry out training and supervision at the provincial level.

#### TB/HIV coordination

Indonesia is classified as a country with low HIV prevalence but with concentrated epidemics, primarily among injecting drug users. A TB/HIV workshop was held in 2002 to consider experiences from central and provincial levels and to develop an action plan for tackling the dual epidemic. A national TB/HIV coordinating body was established and a situation analysis undertaken to assess the linkages between the HIV and TB control programmes in four provinces with high HIV burdens. Guidelines on the management of TB in PLWHA have been published and a pilot project on collaborative TB/HIV activities at the district level is in progress, with funding from WHO.

#### Links with other health-care providers

Indonesia has developed a national strategy for PPM DOTS, focusing primarily on the involvement of public chest clinics and public and private hospitals. Several small-scale pilot projects have been started, and the hospital DOTS linkage project in Yogyakarta has shown a dramatic increase in case detection (>400%) since it began in 2000. Countrywide, very few general hospitals, medical colleges or prison health facilities are involved in DOTS, and there are no treatment providers outside the NTP

that notify cases. However, plans are under way to scale up the successful pilot projects and to start involving private medical practitioners in DOTS.

#### **Partnerships**

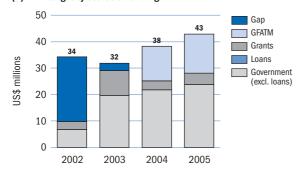
KNCV and WHO are the lead technical partners in Indonesia and support all aspects of DOTS expansion activities. Other technical partners include Kuis/Johns Hopkins, MSH, NLR, TBCTA and World Vision. Major financial partners are the ADB, CIDA, the Dutch Government, GFATM and USAID. A national TB Partners forum meets three to four times a year to share information with partners and donors and to strengthen collaboration between the various participants in TB control.

#### **Budgets and expenditures**

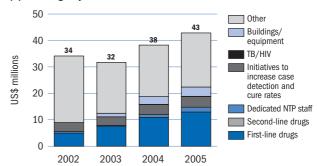
Funding for TB control has improved substantially since 2002, when the NTP reported a funding gap exceeding 50% of the total budget requirement, and expenditures amounting to US\$ 18 million. Available funding more than doubled between 2002 and 2003, although a small funding gap remained. The 2004 budget was fully funded, as is the projected budget of US\$ 43 million for 2005. If the funds are fully disbursed, spending by the NTP in 2005 will more than double that in 2002. This impressive growth in funding is primarily because of a large grant from the GFATM, which will provide 34% of the NTP budget in 2005, in addition to the increase in government funding. The additional funds allow for an increase in the anti-TB drug budget, as well as more spending on initiatives to improve case detection and cure rates. As projected total case detection and total spending increase, the NTP budget and total TB control costs (i.e. the NTP budget plus estimated spending on health clinic visits not covered by the NTP budget) are expected to remain relatively constant per patient, at about US\$ 150-160 and US\$ 180, respectively. It remains to be seen whether the increased funding can be absorbed, and whether increased expenditures result in improved case detection.

### **INDONESIA**

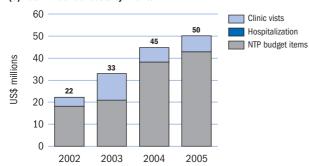
#### (a) NTP budget by source of funding



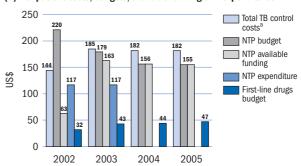
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



<sup>&</sup>lt;sup>a</sup> Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

# Kenya

Kenya adopted the DOTS strategy in the early 1990s and achieved nationwide DOTS coverage in 1996. The diagnosis and treatment of TB are integrated into the Kenyan public health services. TB notification rates have increased five-fold in the past 10 years, the main explanation is probably the impact of the HIV epidemic, although improvements in programme performance may also have contributed; encouraging private physicians to provide DOTS services has increased case-finding in recent years. Furthermore, in collaboration with NGOs, DOTS services are being extended to remote areas, nomadic populations and urban slums. PPM-DOTS initiatives are succeeding but there is still scope to increase the involvement of community workers. An estimated 29% of TB patients are HIVpositive, and TB/HIV coinfection is now a significant problem. Kenya is actively promoting collaboration between the TB and HIV programmes, supported by the "3 by 5" initiative and funding from the President's Emer-

gency Plan for AIDS Relief. It is anticipated that up to 35% of those who start ART in the public sector will be TB patients.

#### **System of TB control**

The NTP (known locally as the National Leprosy and TB Programme) adopted the DOTS strategy in the early 1990s, and TB diagnosis and treatment are integrated into the public health services at all levels. At the central level, the NTP develops TB control policies and offers technical assistance to health-service providers. Other central level responsibilities include surveillance, training, advocacy and resource mobilization activities.

Kenya's laboratory system includes the Central TB Reference Laboratory based at the Kenya Medical Research Institute, which functions as an NRL, four private medical laboratories in Nairobi that are able to perform culture and susceptibility testing for firstline drugs and 619 laboratories that perform smear microscopy. All public and faith-based hospitals, health

centres and some dispensaries carry out sputum microscopy. About three quarters of all registered medical laboratories in Kenya - government, NGO and private - do smear microscopy for the NTP.

#### **Surveillance and monitoring**

The TB notification rate increased fivefold between 1980 and 2003, but the rate of increase is declining. In assessing the case detection rate, it has been assumed that the increase in case notifications reflects a real increase in incidence; it is also possible that case detection has improved in recent years. The spread of HIV infection in Kenya has almost certainly been responsible for much of the increase but also makes it difficult to estimate the true case detection rate without more detailed analysis of subnational data. The proportion of notified cases that are diagnosed as smear-positive has fallen steadily since 1995, possibly because HIVpositive people are more likely than HIV-negative people to present with smear-negative TB. A similar pattern is seen in other countries in eastern and southern Africa where the prevalence of HIV is high.

The treatment success rate in 2002 was 79%, still below the 85% target, largely because 15% of patients defaulted or were transferred to other treatment centres without followup, and 5% died. The loss of patients from the cohort could be associated with HIV infection but may also reflect weaknesses in programme management. For either or both of these reasons, the treatment outcomes for new smear-positive patients have not improved much since 1994. The treatment outcomes among patients registered for re-treatment following relapse were somewhat worse, and the death rate was 10%. Outcomes are not available for patients treated after failure or default.

As long as the incidence of TB remains high because of the HIV epidemic in Kenya, the epidemiological

#### **PROGRESS IN TB CONTROL IN KENYA**

#### **Indicators**

DOTS treatment success, 2002 cohort	
DOTS case detection rate, 2003	46%
NTP budget available, 2004	75%
Government contribution to NTP budget, including loans, 2004	
Government contribution to total TB control costs, including loans, 2004	
Government health spending used for TB control, 2004	7%

#### **Major achievements**

- Significant improvement in human resources capacity in the central unit
- Expansion of PPM DOTS, reinstatement of NRL, establishment of the TB/HIV coordinating body, and development of several guidelines and of the urban TB
- Secured sufficient drugs and funds for DOTS implementation and expansion
- Increased case-finding through decentralization of TB diagnostic services, coupled with improvement of diagnostic procedures
- Development and implementation of the COMBI plan that is aimed at influencing the health-seeking behaviour of the population to improve early case detection

#### **Major planned activities**

- Implement effective TB/HIV collaborative programme: VCT, co-trimoxazole preventive therapy and ART for HIV-infected TB patients
- Improve human resources by recruiting additional staff at central and peripheral levels to boost training and supervision

LATEST ESTIMATES		TRENDS	2000	2001	2002	2003
Population	31 987 119	DOTS coverage (%)	100	100	100	100
Global rank (by est. number of cases)	10	Notification rate (all cases/100 000 pop)	210	235	254	286
Incidence (all cases/100 000 pop/year)	610	Notification rate (new ss+/100 000 pop)	94	101	109	119
Incidence (new ss+/100 000 pop/year)	262	Detection of all cases (%)	46	47	46	47
Prevalence (all cases/100 000 pop)	884	Case detection rate (new ss+, %)	48	47	46	46
TB mortality (all cases/100 000 pop/year)	133	DOTS case detection rate (new ss+, %)	44	47	46	46
TB cases HIV+ (adults aged 15-49, %)	29	DOTS case detection rate (new ss+)/coverage (%)	44	47	46	46
New cases multidrug resistant (%)	0.0	DOTS treatment success (new ss+, %)	80	80	79	_

## Notification rate (per 100 000 pop) = all cases (91 522 in 2003) = ss+ cases 200 100

1991

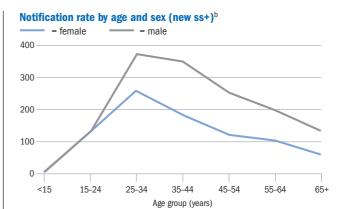
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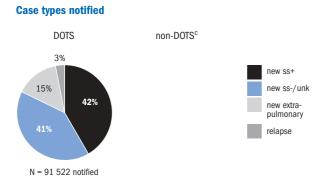
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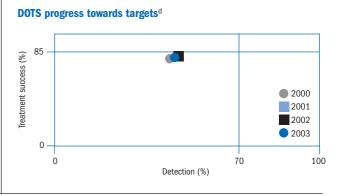
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2000

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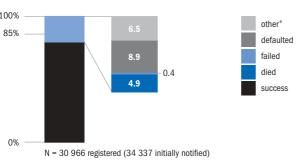




1982

1985

1988



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- e "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

impact of the DOTS programme will be hard to evaluate from routine surveillance data alone. A populationbased survey of disease prevalence would give a better estimate of the current burden of TB in Kenya and provide a baseline against which to assess the future impact of control programmes.

#### Improving programme performance

The Government of Kenya is committed to providing anti-TB drugs for all new patients. The NTP also receives support for drugs from the GDF. The last official data from a drug resistance survey were reported in 1995. In December 2004, the GLC approved a DOTS-Plus pilot project, with funding from the GFATM.

A national TB control plan is being developed for 2005-2010. Tuberculin surveys were carried out in 1958-1959, 1986–1990 and 1990–1995. A fourth tuberculin survey has already started, in collaboration with KNCV, and should be completed in 2006.

An urban TB control project is planned, with a focus on expanding TB services to slum populations in cities; funds from the GFATM will allow several new activities to start. Diagnostic and treatment services are expanding, and NTP activities in collaboration with the national AIDS programme, private sector representatives (physicians and pharmacies), prison authorities and selected NGOs are continuing in several districts.

The COMBI communication plan was launched in April 2004, when materials developed by several agencies were presented. A system to monitor and evaluate the implementation, distribution and impact of these materials is being established.

Three other areas in which programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers and the community.

#### Diagnostic and laboratory services

Expansion of the diagnostic services continues, with the number of laboratories that perform smear microscopy increasing from 542 in 2003 to 619 in 2004. EQA for smear microscopy, in accordance with international guidelines, is being adopted for regional and district laboratories and its implementation started in May 2004 in some districts. Discussions on the establishment of a network of public and private laboratories and the inclusion of these laboratories in the existing EQA system are in progress. Major constraints for the laboratory services include inadequate human resources and an insufficient budget allocation for supervisory activities. Until 2003, the Central TB Reference Laboratory did not have a clear mandate to function as an NRL. At present, one of the main priorities for the NTP is to continue to improve the technical capacity of this laboratory. The NRL is currently equipped to carry out rapid liquid culture techniques and DST on all re-treatment cases from Nairobi as well as re-treatment and failure cases from other provinces. New laboratory guidelines for sputum examination by AFB microscopy have been developed and are to be published in 2005.

#### TB/HIV coordination

Kenya, like many other countries in sub-Saharan Africa, is severely affected by the HIV/AIDS epidemic. An estimated 29% of adult TB patients in Kenya are HIV-positive; a new survey of HIV in TB patients is planned for 2005. In 2003, a national TB/HIV coordinating body was set up, including representatives from the TB and HIV programmes, research institutions, technical agencies, donors and representatives of PLWHA. A national TB/HIV coordinator was appointed and is the secretary of the steering committee. TB/HIV activities have started in Nakuru District, and by the end of 2005, should have started in about 30 other districts.

Kenya is one of the pilot sites for the "3 by 5" initiative and is receiving funding from the President's Emergency Plan for AIDS Relief. In 2005, about 45 000 TB patients should be offered HIV testing and a package of prevention and care, including ART. It is estimated that about 35% of patients who are eligible for ART will be

identified through the TB control services. A monitoring and evaluation system for TB/HIV activities is now being developed and tested in selected districts.

#### Links with other health-care providers

An initiative to encourage private physicians to provide DOTS services in Nairobi was started in 2001 and is now being implemented in several other towns and settings. This has led to an increase in case notification rates, and treatment results have been satisfactory. Guidelines for PPM-DOTS have been developed and staff trained. Collaboration between the NTP, NGOs and a variety of public sector health providers and related institutions, including general hospitals, medical colleges and health services in refugee camps, prisons, military and the police, is still in progress.

#### Links with the community

Community-based DOTS was successfully pilot tested in Machakos District between 1998 and 2000, and 11 other districts have recently started training community volunteers. District teams, comprising nurses, social workers, health educators and public health workers, train community volunteers in increasing awareness of TB, early referral of suspects and treatment support.

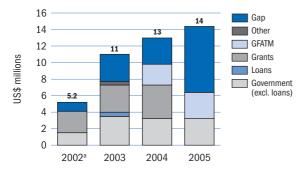
#### **Partnerships**

Financial support to the NTP in Kenya is mostly provided by CIDA, CDC and USAID. The NTP has a three-year agreement with the GDF for anti-TB drugs, which expired at the end of 2004. The World Bank supported the NTP through a loan for the purchase of anti-TB drugs. A GFATM grant agreement was signed in round 2 and will provide significant funding for DOTS expansion activities. KNCV and WHO are the main technical partners.

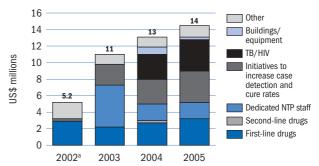
#### **Budgets and expenditures**

The NTP budget has increased steadily from US\$ 5.2 million in 2002 to US\$ 14 million in 2005; the budget per patient has increased from about US\$ 67 per patient in 2002 (for about 60 000 patients) to US\$ 142 per

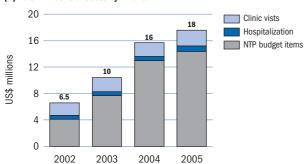
#### (a) NTP budget by source of funding



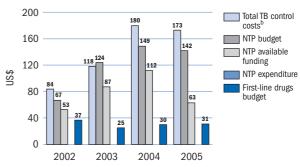
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>b</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



- <sup>a</sup> Does not include budget for buildings/equipment and dedicated NTP staff.
- b Total TB control costs for 2002 and 2003 are based on available funding, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

patient in 2005 (based on a projection by the NTP that about 100 000 patients will be treated in 2005). The government contribution to the NTP budget has been fairly constant at about US\$ 3.5 million per year (although this is underestimated to some extent because funds budgeted for investment in buildings and equipment are not reflected in the NTP budget). All grants, except the GFATM grant, end in June 2005 and need to be renegotiated. This explains why a funding gap of US\$ 8 million, equiva-

lent to about 50% of the NTP budget. is reported for 2005. It is anticipated that external funding will be secured to reduce this gap.

The increased budgets in 2004 and 2005 are to allow increased spending on collaborative TB/HIV activities and initiatives aimed at improving case detection and cure rates. Implementation of these activities will depend on closing the funding gap. No expenditure data are available for the years 2002 or 2003.

If the NTP budget is fully funded

and the money is spent, total TB control costs (which include visits to health clinics and expenditures related to hospitalization in addition to NTP budget items) will be about US\$ 16–18 million in 2004 and 2005, or about US\$ 180 per patient treated (compared with about US\$ 120 in 2003). The increase in total costs per patient is almost entirely due to changes in costs included in the NTP budget.

# Mozambique

Mozambique has a longstanding commitment to TB control, and the DOTS strategy was introduced in all districts by 2000. Nevertheless, the NTP still faces substantial difficulties in providing adequate TB services throughout the country. Efforts are being made to strengthen the country's health infrastructure, which has suffered in the past from inadequate resources as well as the destructive effects of civil unrest and natural disasters. Despite these difficulties, TB case detection and cure rates have improved in recent years. The award of a GFATM grant will greatly increase the amount of funds available for TB control in 2005 and make it possible to address problems related to staffing, training and medical supplies and equipment. With improvements in facilities for diagnosis and patient care, case detection and cure rates should continue to improve in the next few years. Progress will be gradual because it will take time to build up the necessary cadre of well-trained staff to carry out the full programme of DOTS activities. With the help of in-

ternational partners, Mozambique is beginning to tackle the special challenges associated with high rates of Tb and HIV coinfection. The extent of MDR-TB is being investigated in a new national survey.

#### **System of TB control**

The NTP was officially established in 1977 and consists of a central unit, 3 regional coordinators, 12 provincial coordinators and 149 health area district coordinators. There are 3 central hospitals (one in each region), 7 provincial hospitals, 27 rural hospitals and 162 health centres, all involved in DOTS implementation. There are also approximately 800 health posts, managed by rural health workers, which are not part of the DOTS programme.

The TB laboratory network has an NRL in Maputo City that performs culture and DST. There are 45 intermediate laboratories, 11 of which are located in the capital cities of the provinces, and 163 peripheral laboratories located mostly in health centres of the district capital cities. There are no microscopy services in any of the health posts.

#### **Surveillance and monitoring**

The total number of TB cases notified, both smear-positive and all forms, continued to increase between 2002 and 2003. The proportion of new pulmonary cases diagnosed as smearpositive was 67% in 2003. This is towards the lower end of the expected range of 65-80%, as often seen in countries with high rates of HIV infection. For Mozambique, as for some other countries in southern Africa, the accuracy of the estimated case detection rate (45% in 2003) is uncertain. The treatment success rate was 78% for the 2002 cohort and has improved each year since 1995. In 2002, 11% of patients died and 10% defaulted or were lost to follow-up after transfer to other treatment centres. Both of the latter indicators were high for patients undergoing re-treatment following relapse, failure or default; the overall retreatment success rate was 67%. Considering progress towards the MDGs, the priority for Mozambique is to bring treatment success and case detection rates closer to target levels.

## Improving programme

performance Implementation of the NTP's five-year national strategic plan began in 2003. Financial constraints have hindered almost all aspects of programme performance in Mozambique, partly because of a change in financial transfer mechanisms at the central level. Future improvements in the programme rely heavily on the GFATM grant, which was signed in April 2004. Disbursement of funds started in September 2004 but has not yet reached the sub-recipients. However, it may prove difficult for the central unit to meet the staffing requirements needed to begin implementation of the overall GFATM plan. A number of key staff need to be recruited, including TB coordinators for the central, southern and northern regions as well as

## PROGRESS IN TB CONTROL IN MOZAMBIQUE

#### **Indicators**

DOTS treatment success, 2002 cohort	78%
DOTS detection rate, 2003	45%
NTP budget available, 2004	44%
Government contribution to NTP budget, including loans, 2004	30%
Government contribution to total TB control costs, including loans, 2004	46%
Government health spending used for TR control 2004	6%

#### **Major achievements**

- Implementation of a five-year strategic national plan for the NTP (2003–2008)
- Approval of GFATM funding for overall NTP strengthening, and GDF funding for FDC anti-TB drugs
- Development of a national TB/HIV collaborative project

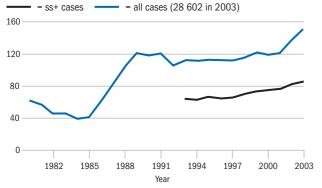
#### **Major planned activities**

- Develop a drug management system to estimate the number of drugs and supplies needed at central and provincial level, and a plan to manage drug short-
- Establish laboratory quality control in collaboration with provincial laboratory supervisors
- Commence collaborative TB/HIV activities in demonstration project sites, including surveillance of HIV in TB patients in 2005 and the introduction of isoniazid preventive therapy and co-trimoxazole preventive therapy

## **MOZAMBIQUE**

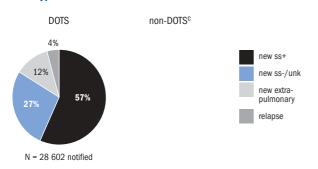
LATEST ESTIMATES®		TRENDS	2000	2001	2002	2003
Population	18 863 291	DOTS coverage (%)	100	100	100	100
Global rank (by est. number of cases)	18	Notification rate (all cases/100 000 pop)	118	121	138	152
Incidence (all cases/100 000 pop/year)	457	Notification rate (new ss+/100 000 pop)	74	77	82	86
Incidence (new ss+/100 000 pop/year)	190	Detection of all cases (%)	30	29	32	33
Prevalence (all cases/100 000 pop)	636	Case detection rate (new ss+, %)	45	44	45	45
TB mortality (all cases/100 000 pop/year)	129	DOTS case detection rate (new ss+, %)	45	44	45	45
TB cases HIV+ (adults aged 15-49, %)	49	DOTS case detection rate (new ss+)/coverage (%)	45	44	45	45
New cases multidrug resistant (%)	3.5	DOTS treatment success (new ss+, %)	75	77	78	_

#### Notification rate (per 100 000 pop)

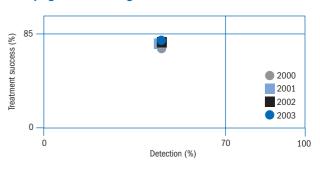


#### Notification rate by age and sex (new ss+)<sup>b</sup>

#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- e "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

## **MOZAMBIQUE**

technical and administrative support staff and laboratory technicians. In preparation for the GFATM grant, six TB managers were trained at the WHO Collaborating Centre for TB and Lung Disease in Sondalo, Italy, in July 2003.

In 2003 and 2004, a serious shortage of anti-TB drugs occurred because of lack of funds. In May 2004, there were no stocks of ethambutol or pyrazinamide at the central level and a severe shortage of drugs was reported in some provinces. Fortunately, a three-year grant was approved by the GDF at the time of these shortages. FDC anti-TB drugs will be procured in place of loose formulations from 2004. A drug management system will be developed with GFATM funds; this will include a simple computer spreadsheet to estimate the amount of drugs and supplies needed at the central and provincial levels, a mandatory oneyear national buffer stock of drugs and laboratory reagents to be maintained at the central level, and a mechanism for responding to unforeseen shortages by mobilizing additional support.

A new national drug resistance survey is planned. The protocol has been finalized and implementation should

start in January 2005. DST is performed on isolates from patients failing re-treatment. However, no proper treatment is available for MDR-TB patients. A proposal will be submitted to the GLC in 2005.

Three other areas where programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

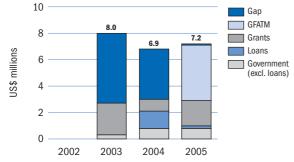
The quality of DOTS implementation is limited by the poor laboratory network. Existing laboratories are inadequately distributed throughout the country, and more than half of the population is served by health centres that are 10 km or further from a diagnostic facility. The protocol for quality assurance of smear microscopy has been finalized but was not put into practice as scheduled in January 2004 because of lack of funds. Regular laboratory supervision has also stopped due to a lack of funds. There are plans to establish laboratory quality control (LQC) in collaboration with provincial laboratory supervisors and to identify and train staff in LQC. Provision of culture testing and quality control call for the upgrading of laboratories in Beira (central region) and Nampula (northern region), but infrastructure improvements have been postponed until GFATM funding becomes available. The NRL will be strengthened only if funds can be obtained from sources other than the GFATM. During 2003, a number of health facilities that had been out of service were rehabilitated and several health posts upgraded to health centres with smear microscopy services.

The lack of technicians throughout the laboratory services in Mozambique will be addressed through intensified training of existing technicians and, with forthcoming funding, re-hiring of qualified technicians.

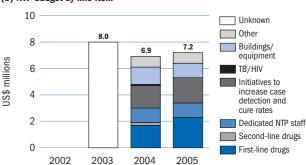
#### TB/HIV coordination

A collaborative TB/HIV project (2004-2005) is being funded by WHO and USAID and implemented with support from KNCV. Good progress is being made at the central level. A TB/HIV coordinator has been recruited and a body to coordinate collaborative TB/ HIV activities at all levels established. The team will oversee the develop-

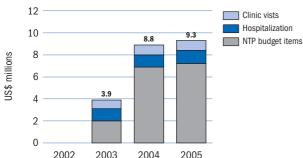
## (a) NTP budget by source of funding



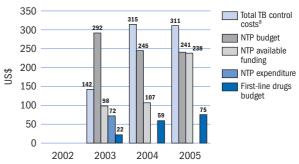
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

## **MOZAMBIQUE**

ment of a national policy for collaborative TB/HIV activities and its implementation, including planned demonstration projects. Planned project activities include surveillance of HIV in TB patients in 2005 and the introduction of isoniazid and cotrimoxazole preventive therapy. Training materials for TB/HIV are being developed and a workshop is planned. An important step has been the inclusion of the TB/HIV monitoring and evaluation indicators in the integrated health network system for HIV/AIDS; good progress is being made.

#### Links with other health-care providers

Private sector involvement is restricted to a few private hospitals that diagnose and treat TB under NTP guidance. The NTP is beginning to involve medical colleges, specialist TB clinics, and prison, army and police health facilities in DOTS activities.

#### **Partnerships**

External funding for TB control comes from the governments of Italy, Norway and the USA (USAID) and from the

GFATM. WHO and KNCV are the main technical partners; CARE International, CDC and MSF-Belgium/Luxembourg are additional partners in the TB/HIV project.

#### **Budgets and expenditures**

The NTP budget decreased from an estimated US\$ 8.0 million in 2003 to US\$ 6.9 million in 2004. However, available funding has increased from around US\$ 3 million in 2003 to almost US\$ 7 million in 2005. This means that the funding gap has fallen from 66% of the budget in 2003 to a projected 1% of the budget in 2005. This improved funding situation is mainly a result of an increase in loans and grants, including approval of a GFATM grant in round 2. The 2005 budget will be highly dependent on external financing, with around 85% of funds provided by grants, even though the government's contribution to the NTP budget has remained constant in absolute terms.

One of the largest budget line items is first-line anti-TB drugs. This budget has been increasing in recent years

to allow creation and consolidation of a buffer stock (this is why the budget per patient for first-line drugs has increased since 2003). There has also been a large increase in the budget for initiatives to increase case detection and cure rates between 2003 and 2005. These initiatives will be scaled up once GFATM funds become available. For the first time, there is a budget for collaborative TB/HIV activities in 2004 and 2005; however, this represents only around 1% of the NTP budget.

The total TB cost per patient based on budget data is about US\$ 300 for the years 2004 and 2005. However, the total cost per patient based on actual expenditure was substantially lower in 2003 at US\$ 142 because of the large funding gap. The total annual cost of TB control (including visits to health clinics and hospitalization as well as NTP budget items) is projected to increase from US\$ 3.9 million in 2003 to US\$ 9.3 million in 2005.

# Myanmar

After a period of rapid DOTS expansion, Myanmar achieved nationwide DOTS coverage by the end of 2003. That year, more than 75 000 TB cases were reported, corresponding to a case detection rate of 73%; more than 80% of the 2002 cohort were treated successfully. With a strong national health infrastructure and government recognition of TB as a top priority, the country is now within sight of becoming the second of the current group of HBCs1 to reach the global targets for DOTS implementation (after Viet Nam). Myanmar has made these commendable achievements with little external donor support. This situation will change radically with a massive

increase in funding, mainly from the GFATM. When they become accessible, these funds will provide major opportunities for capital investment in infrastructure as well as important improvements in staffing at all levels and in the quality of laboratory services. They will also enable sustainability and strengthening of all aspects of the NTP including further boosting of treatment outcomes. Several NGOs now participate in the provision of TB control services, and the NTP is promoting the involvement of other health-care providers, particularly private physicians and clinicians from large hospitals. A national TB prevalence survey would provide a more

accurate estimation of incidence and a baseline for assessing the impact of DOTS services on the TB epidemic.

### **System of TB control**

The NTP functions through a central level office and 12 state or divisional TB centres. There is one central drug store and two subnational stores in upper and lower Myanmar. Township hospitals serve as the DOTS treatment units, and TB registers are maintained at this level for the population in each township.

The NRL was established in 2001, and there are two subnational laboratories. Since 2003, all state and divisional laboratories participate in a quality assurance network. Sputum smear microscopy is done in 309 of 324 townships. The NRL carries out drug susceptibility testing and, together with the subnational laboratory in Mandalay, also performs culture.

#### **Surveillance and monitoring**

The total number of reported TB cases increased from less than 15 000 in 1998 to more than 75 000 in 2003, with DOTS coverage reported to be 95% of the population during 2003 (rising to 100% towards the end of the year). During the same period, the smear-positive case detection rate increased from 29% to an estimated 73%, exceeding the 70% target. During this period of rapid DOTS expansion, the proportion of all new cases diagnosed as smear-positive fell from 68% (1998) to 36% (2003), which raises questions about the accuracy of the microscopic diagnosis. Treatment success has exceeded 80% since 1997, but moderately high default rates (9% in 2002) have limited the rate of success. As expected, the treatment success rates are somewhat lower for patients undergoing retreatment (76% among relapses, 75% among all re-treatment cases combined). Nonetheless, on current evi-

## Technical Working Group on TB established by WHO to facilitate coordination and collaboration between agencies

#### **Major planned activities**

- Strengthen the national laboratory network by expanding the network of smear microscopy centres, improve quality control and upgrade the subnational laboratory in Mandalay
- Increase programme management capacity through GFATM funding (training, case management, supervision and monitoring, drug management, human resource development)
- Implement pilot project on ARV for HIV-infected TB patients in Mandalay
- Scale up PPM-DOTS projects
- Prepare 10-year plan for NTP (2006–2015), with emphasis on achievement of
- Apply to GDF for a second three-year grant
- Apply to ISAC for funding of technical assistance in GFATM project implementa-

## **PROGRESS IN TB CONTROL IN MYANMAR**

#### **Indicators**

DOTS treatment success, 2002 cohort	81%
DOTS case detection rate, 2003	73%
NTP budget available, 2004	34%
Government contribution to NTP budget, including loans, 2004	6%
Government contribution to total TB control costs, including loans, 2004	18%
Government health spending used for TB control, 2004	0.4%

#### **Major achievements**

- DOTS expanded to all townships and case detection of new smear-positive patients above global target
- Operational guidelines on the involvement of private practitioners in DOTS pub-
- Treatment guidelines for HIV-infected TB patients published
- Nationwide introduction of FDC anti-TB drugs
- GFATM grant agreement signed in August 2004 and first funds distributed to principle recipient (UNDP)
- First nationwide drug resistance survey completed

<sup>&</sup>lt;sup>1</sup> Peru was excluded from the original group of HBCs, having met the targets and successfully reduced incidence.

## **MYANMAR**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	49 485 491	DOTS coverage (%)	77	84	88	95
Global rank (by est. number of cases)	20	Notification rate (all cases/100 000 pop)	65	89	117	153
Incidence (all cases/100 000 pop/year)	171	Notification rate (new ss+/100 000 pop)	36	44	49	55
Incidence (new ss+/100 000 pop/year)	76	Detection of all cases (%)	38	52	68	90
Prevalence (all cases/100 000 pop)	187	Case detection rate (new ss+, %)	48	58	65	73
TB mortality (all cases/100 000 pop/year)	25	DOTS case detection rate (new ss+, %)	48	56	65	73
TB cases HIV+ (adults aged 15-49, %)	6.8	DOTS case detection rate (new ss+)/coverage (%)	62	67	74	77
New cases multidrug resistant (%)	4.0	DOTS treatment success (new ss+, %)	82	81	81	_

## Notification rate (per 100 000 pop) = all cases (75 744 in 2003) = ss+ cases 120 80 40

1991

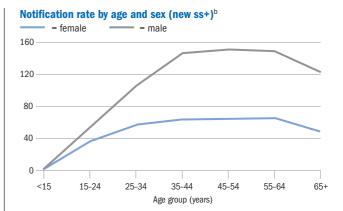
Year

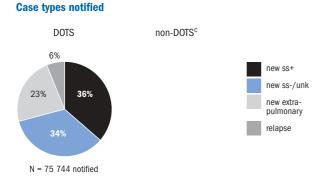
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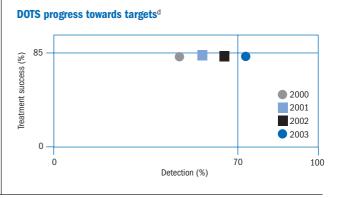
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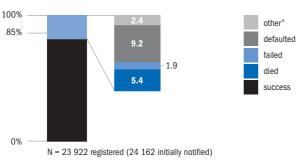




1985

1982

1988



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

dence, Myanmar is close to reaching the targets for case detection and treatment success. One important caveat is that the denominator of the case detection rate - the estimated smear-positive incidence rate - is based on disease prevalence surveys carried out up to 1994. A decade later, a new national prevalence survey would provide a valuable reassessment of the burden of TB in Myanmar, give a baseline against which to evaluate DOTS impact and yield important information about the quality of diagnosis and treatment.

## Improving programme performance

Development of human resource capacity has been strengthened, but many challenges still remain. An HR database is in place and shows that approximately a quarter of all sanctioned posts in the NTP are vacant. An HR development plan was prepared with WHO in 2003 and has already resulted in intensified training activities and the appointment of new staff. Future plans include cascade training of staff involved in TB control at all levels, including community volunteers, national NGOs and private physicians. The GFATM will support training activities and HR development; the NTP will apply for additional support through ISAC's second round of funding.

GFATM funding will also be used to increase monitoring, supervision and evaluation of programme activities and to strengthen DOTS by involving community and national NGOs and private providers, and by improving tracing of defaulters. The role of WHO will be to provide technical support to the principal recipient and subrecipients for planning, implementation and monitoring and evaluation of the TB component of the GFATM grant.

Myanmar currently receives anti-TB drugs through the GDF, which will provide a third year's supply of drugs for 2005, including a buffer stock. The GDF is considering a second term of three years beyond 2005. Following a successful pilot project to introduce FDC anti-TB drugs in the divisions of Mandalay and Yangon, all State/Divisional TB Officers were trained in FDC anti-TB drug management, and FDCs

have been introduced nationwide through cascade training of the Township Medical Officers. A nationwide drug resistance survey was completed in 2003, with the prevalence of MDR-TB among new cases estimated at 4.0%. There is no national policy on MDR-TB management; patients are treated on an individual basis.

Three other areas where programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

Although the TB laboratory infrastructure is improving, strengthening of the national laboratory network is needed, especially the expansion of sputum smear microscopy centres, reinforcement of quality assurance at the township level and upgrading of the subnational laboratory in Mandalay. Another constraint is the shortage of qualified staff, especially junior laboratory technicians, which will be addressed as a priority for 2005. In addition, strengthening techniques for culture and drug susceptibility testing at the NRL in Yangon and the subnational laboratory in Mandalay are planned. It is also planned to introduce culture in four state/divisional laboratories in Bago, Mawlamyine, Pathein and Taunggyi.

#### TB/HIV coordination

The HIV prevalence among TB patients was estimated to be 4.5%, based on surveillance carried out in 20 sentinel sites from 1995-1997. This is lower than the current WHO estimate of 6.8%. No recent estimates from surveys are available, but a new HIV prevalence study among TB patients is planned once funding from the GFATM becomes accessible. Funding from GFATM will also support TB/HIV training for NTP staff at all levels. Political commitment has been demonstrated by the establishment by the MoH of a high-level coordinating body on TB/HIV. TB/HIV prevention and control activities were implemented in five pilot townships in 2000, including VCT for TB patients and provision of HIV education and prevention for HIVinfected TB patients. These activities were discontinued because of lack of funding. Currently, there are limited collaborative TB/HIV activities in the country. Treatment guidelines for TB/ HIV have been developed. VCT is available at a small number of VCT centres, at some drug treatment centres and at hospitals offering prevention of mother-to-child transmission programmes.

Although a supply of drugs for ART has arrived in Myanmar, ART is not yet available through the public sector. Some international NGOs such as MSF Holland and MSF Switzerland are providing ART to TB patients on a small scale. A WHO/National AIDS Programme/NTP/IUATLD/Total Exploration and Production Myanmar project will put 200 TB patients on ART in five townships in Mandalay Division in 2005. Partners within the government, and national and international NGOs have expressed interest in working with the NTP to strengthen existing TB/HIV activities and to actively engage in extending TB/HIV activities.

#### Links with other health-care providers

Involvement of general hospitals has increased rapidly during the past two years. In some areas, TB cases notified from general hospitals represent a substantial proportion of all cases registered under DOTS. However, a high proportion of the cases notified from hospitals are extrapulmonary or sputum smear-negative pulmonary TB, which raises some concern about the diagnostic quality in these hospitals. Involvement of army, police and prison health services has started but is still limited.

The Myanmar NTP has developed national guidelines for involvement of private practitioners in TB control. So far, two initiatives to involve private providers have been launched. In Mandalay Division, the NTP, together with the Department of Medical Research, started a project in 2002 to involve private physicians in diagnosis and treatment under the NTP. In Yangon, an international NGO (Population Services International) started implementing DOTS in 2004 as part of its existing franchising scheme, under which private physicians deliver diagnostic services at low cost and provide treatment with drugs free of charge from the NTP. Both initiatives

## **MYANMAR**

have contributed substantially to increased case detection in targeted townships. Several additional initiatives are planned, including a training programme for private physicians, which will be coordinated by the Myanmar Medical Association, and a joint initiative by JICA and NTP to involve private practitioners in selected townships in Mandalay and Yangon Divisions.

#### **Partnerships**

Many national and a limited number of international NGOs work together on TB control in Myanmar. IUATLD supports operational research, TB/HIV activities including ART, and procurement of cars and laboratory equipment and supplies. WHO provides technical support and assists with HR development and procurement of drugs and laboratory supplies. JICA offers laboratory training, and anti-TB drugs are supplied by the GDF. The GFATM will soon be the main funding partner. The Country Coordination Mechanism has merged its working group with the Technical Working Group on TB hosted by WHO. The role of this working group is to support the United Nations Theme Group on Health (UNTGH) and the GFATM principal

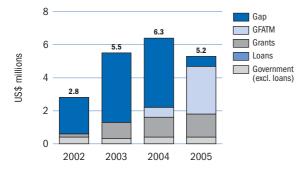
recipient, the UNDP, by providing technical advice on all aspects of the implementation of the TB control programme funded by the GFATM and by other external sources, and by coordinating all operational and technical aspects between implementing agencies.

## **Budgets and expenditures**

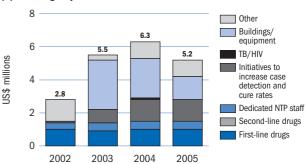
The NTP budget was around US\$ 3 million in 2002, but a large funding gap meant that actual expenditures were only around US\$ 1 million, primarily for staff and first-line drugs. The establishment of the GFATM has created new funding opportunities for the NTP, and following a successful GFATM application in 2003 the budget for 2004 was US\$ 6.3 million. It was anticipated that the GFATM would provide about US\$ 4 million of the reguired funds. However, because of delays in signing the initial two-year grant agreement, the first disbursement was only received by the GFATM principal recipient in September 2004, and thus a substantial funding gap remained. Provided that GFATM funds can be transferred to subrecipients and further disbursements are made according to the grant agreement, funding for the NTP will dramatically

improve in 2005, and the GFATM will be by far the most important source of financing. If this happens, funding per patient treated is likely to rise as well, from expenditures of US\$ 20 per patient in 2002 to a budget of US\$ 70 per patient in 2005. Since case detection is already high, increased funding will mainly provide for improvements in the quality of the existing infrastructure, which are needed to sustain the achievements that have already been made, and to support further improvements in case detection. Much of the increased budget in 2005 is for capital investments that will benefit patients for many years. Thus budgets for subsequent years will be lower (some of these investments were originally planned for 2004, but needed to be deferred to 2005 because of lack of funds; this explains why the budget developed for 2004 was higher than that for 2005). Costs beyond those reflected in the NTP budget are limited in Myanmar, at about US\$ 1 million per year. If the NTP budget is fully funded in 2005, total TB control costs will rise from about US\$ 2 million in 2002 and 2003 (about US\$ 30 per patient treated) to US\$ 6.1 million in 2005 (about US\$ 80 per patient treated).

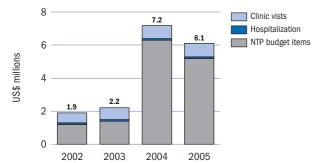
#### (a) NTP budget by source of funding



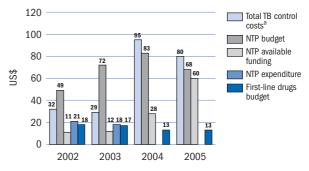
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details

# **Nigeria**

Among African countries, Nigeria has the highest estimated number of new TB cases annually. Following the Abuja Declaration<sup>1</sup> in 2001, the DOTS strategy was adopted nationally and is now being applied in all states. There has been rapid progress in DOTS expansion in 2003, with relatively high treatment success. Unfortunately, political commitment has not yet been translated into strong support for the health system, and much of the approved government funding for health care has not been released for use in health programmes. This situation has discouraged a number of external donors, including the GFATM, who are reluctant to provide additional funds while government funding is very limited. Although Nigeria has an extensive national health infrastructure, it lacks the resources needed to function effectively. Nigeria is now decentralizing its health system and clarifying the responsibilities and services at each level, which should result in better management and

coordination. The spread of HIV infection is adding to the burden of TB; more than a quarter of adults with TB are coinfected with HIV. Notwithstanding the hesitation of some external donors, Nigeria now has an excellent opportunity to develop a programme of collaborative TB/HIV activities with the help of an award from the President's Emergency Plan for AIDS Relief.

#### System of TB control

Although the NTP was launched in 1991, the nationwide adoption and expansion of the DOTS strategy began only recently, following the Abuja Declaration to Stop TB in October 2001. Previously, only half of the states in Nigeria were supported by international NGOs (mainly dealing with leprosy) that were able to provide TB diagnosis and treatment; these did not include Lagos or the Federal Capital Territory (FCT), Abuja. All 37 states have at least one local government area (LGA) that is implementing DOTS.

The public health sector accounts for less than half of the health services provided in Nigeria, the rest being met by NGOs and the private sector, including hospitals, clinics and pharmacies. Health sector reform is under way in order to clearly establish the roles and responsibilities for health service provision at each level, and a Health Act will define the decentralization of functions. Tertiary care is provided and health regulations and technical guidelines developed at the federal level. States are responsible for secondary care and specialized services, while the LGAs are responsible for providing primary health care. The basic unit of health care is the ward. Each ward has 10 000-20 000 people and there are an average of 10 wards per LGA. Ward staff utilize community resources to help deliver the minimum package of care. Public health services will be decentralized from the LGA to the ward, and TB and leprosy control will be included in the minimum package of health services.

The NTP is organized at the federal, state and LGA levels. There is a central unit at the federal level led by a national coordinator. Each of the 37 state programmes is run by a state TB and leprosy control officer. The LGA is the main operational level of the programme, and most LGAs have a TB/leprosy control supervisor. The LGA TB/leprosy control supervisor is, in most cases, a community health officer or nurse who oversees activities in the health facilities.

The NRL in the National Institute for Medical Research in Lagos is responsible for overall supervision and quality assurance of the laboratory network. Six zonal reference laboratories supervise peripheral laboratories. The peripheral laboratories in PHC facilities, NGOs and private facilities all do direct smear microscopy.

#### **PROGRESS IN TB CONTROL IN NIGERIA**

#### **Indicators**

DOTS treatment success, 2002 cohort	79%
DOTS case detection rate, 2003	18%
NTP budget available, 2004	73%
Government contribution to NTP budget, including loans, 2004	37%
Government contribution to total TB control costs, including loans, 2004	66%
Government health spending used for TB control, 2004	4%

## **Major achievements**

- Establishment of DOTS services in three more LGAs in each of 17 states, including training of general health workers and laboratory technicians, and purchase and distribution of laboratory materials
- Approval of a second year of funding from the GDF for anti-TB drugs and distribution of current anti-TB drugs to DOTS facilities throughout the country
- Appointment of a focal point for collaborative TB/HIV activities by the National AIDS and STD Control Programme
- Referral of TB patients with HIV/AIDS for HIV care and support, including ART in 25 pilot sites

#### **Major planned activities**

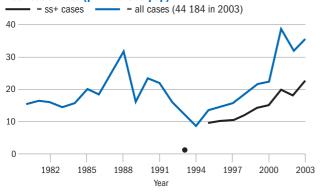
- Assess HR needs and strengthen capacity of general PHC and hospital staff in integrated TB control activities
- Establish at least one microscopy centre in each of the remaining LGAs, and strengthen collaboration between the microscopy centres and the NRL
- Establish a standardized quality assurance system for the entire country in 2005

<sup>&</sup>lt;sup>1</sup> Abuja declaration on HIV/AIDS, tuberculosis and other related infectious diseases. Addis Ababa, The Economic Commission for Africa, 2001.

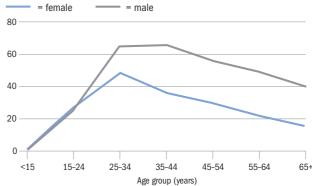
## **NIGERIA**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	124 009 171	DOTS coverage (%)	47	55	55	60
Global rank (by est. number of cases)	4	Notification rate (all cases/100 000 pop)	23	39	32	36
Incidence (all cases/100 000 pop/year)	293	Notification rate (new ss+/100 000 pop)	15	20	18	23
Incidence (new ss+/100 000 pop/year)	126	Detection of all cases (%)	8.9	15	11	12
Prevalence (all cases/100 000 pop)	546	Case detection rate (new ss+, %)	14	17	15	18
TB mortality (all cases/100 000 pop/year)	85	DOTS case detection rate (new ss+, %)	14	14	13	18
TB cases HIV+ (adults aged 15-49, %)	27	DOTS case detection rate (new ss+)/coverage (%)	30	25	24	30
New cases multidrug resistant (%)	1.7	DOTS treatment success (new ss+, %)	79	79	79	_

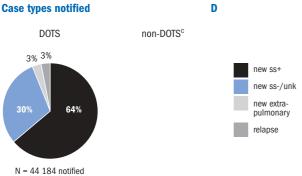
## Notification rate (per 100 000 pop)



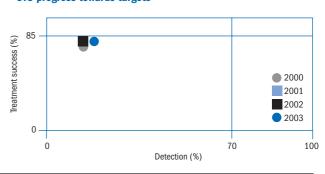
### Notification rate by age and sex (new ss+)<sup>b</sup>



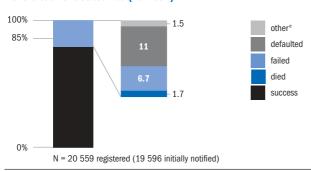
## **Case types notified**



## **OTS** progress towards targets<sup>d</sup>



## **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

 $ss+\ indicates\ smear-positive;\ ss-,\ smear-negative;\ pop,\ population;\ unk,\ unknown.$ 

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- e "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

#### **Surveillance and monitoring**

Among African countries, Nigeria has the highest estimated number of new TB cases each year. An estimated 6% of all adults, and 27% of adult TB patients, are infected with HIV. The increase in case notifications since 1994 is almost certainly due to a rise in TB incidence associated with the spread of HIV, rather than to improvements in case detection. DOTS coverage has changed little over the nine years for which data have been submitted to WHO (1995-2003), although there was a small increase between 2002 (55%) and 2003 (60%). The proportion of cases that were smear-positive fell between 1995 and 2003. While this could be due in part to increases in TB among HIV-infected people, the reasons for the observed trend need to be investigated further. Although the DOTS case detection rate has increased, the estimate for 2003 remains low at 18%. The treatment success rate was 79% in the 2002 DOTS cohort, with a high default rate (11%). Treatment success has increased only slightly since 1997. Nigeria has not yet taken steps to evaluate the impact of DOTS in reducing transmission, incidence, prevalence or deaths.

## Improving programme performance

A major constraint for PHC and the TB control programme is the failure of the government to release funds that have been budgeted and allocated for health and TB control services at all levels. This reflects a low level of political commitment and results in reliance on external funding for TB control operations, mostly from CIDA, DFB, GLRA, NLR and USAID. CIDA funding from mid-2002 to the end of 2003 has made it possible to expand DOTS to the remaining 16 non-DOTS states and FCT Abuja (thus expanding DOTS to all the 36 states of the Federation, including FCT Abuja), to strengthen the central unit's infrastructure and coordination and to establish three zonal TB coordination and control offices. A TBCTA/USAID grant has provided funds for some TB control activities in 2004, including the expansion of the TB laboratory network, providing

supervision and monitoring activities at the central and zonal levels, and training staff and developing human resources for collaborative TB/HIV activities.

Another challenge facing the NTP in Nigeria is the lack of professional health staff in the LGAs. The PHC facilities are staffed mainly by nurses and community health workers, and the physician to population ratio is between 1:160 000 and 1:400 000. Although the TB programme trains supervisors and other senior staff, very few general PHC and hospital staff have been trained in integrated TB control activities. HR needs are being assessed with a view to revising the HRD plan. Currently, states are responsible for training their own staff, while the federal government supports training programmes for TB control in collaboration with research institutions and universities. The National TB and Leprosy Training Centre in Zaria, established in 1991, is responsible for providing the necessary staff training at the LGA and health facility levels. The centre provides a three-month course for LGA TB control supervisors and a two-week course for laboratory technicians.

The supply of anti-TB drugs is adequate, and an application to the GDF for a second year of support has been approved. However, the country's drug policy dates from 1990 and is currently under revision. The federal government is responsible for legislation concerning drugs, while the management and procurement of drugs are decentralized to individual facilities. There is no system of drug control at national or provincial levels once drugs have been approved. There are no drug resistance data for the country.

#### Diagnostic and laboratory services

The number of TB laboratories is increasing, and smear microscopy is now available in 504 out of 774 LGAs. However, few of these laboratories are covered by a quality assurance system. Nigeria plans to establish a standardized quality assurance system for the whole country in 2005. Stocks of laboratory reagents are low because of the lack of government funding. Most laboratories receive reagents from the NTP (funded by WHO) or from NGOs, and some are charging patients. In 2005, the NTP plans to establish at least one microscopy centre in each of the remaining LGAs and to strengthen collaboration between the microscopy centres and the NRL.

#### TB/HIV coordination

The National AIDS and STD Control Programme has appointed a staff member to act as the focal point for collaborative TB/HIV activities. An NPO will be recruited to support these activities using funds provided by the Norwegian government. Many DOTS and ART centres are now starting collaborative TB/HIV activities. In 25 sites, TB patients with HIV/AIDS will have access to comprehensive HIV/ AIDS care and support, including the provision of ART.

At the central level, a proposal to develop a strategy document for collaborative TB/HIV activities has been finalized and preparations for a highlevel mission are being made in relation to the "3 by 5" initiative. Collaborative TB/HIV activities are constrained by the shortage and high cost of HIV test kits and the shortage of antiretrovirals and drugs for opportunistic infections at both HIV and TB treatment centres. Following a recent award from the President's Emergency Plan for AIDS Relief, collaborative TB/HIV activities will be expanded in 2005.

#### Links with other health-care providers

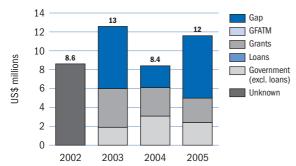
The NTP has successfully pilot tested the involvement of private clinics in the delivery of DOTS services; this initiative is being expanded to six states with financial support from FIDELIS. Several NGOs are already involved, with efforts being made to strengthen collaboration with general hospitals, specialist TB clinics, medical colleges and prison health serv-

#### **Partnerships**

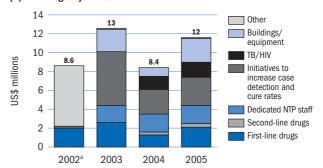
Major technical partners include DFB, DFID, GLRA, IUATLD, Netherlands Leprosy Relief and WHO. CIDA and USAID (TBCTA) are the main funding partners. The GDF provides anti-TB drugs and will start to provide laboratory test kits

## **NIGERIA**

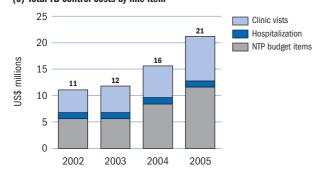
#### (a) NTP budget by source of funding



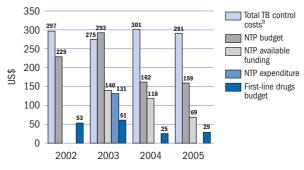
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>b</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



- a In the 2002 budget, the costs of dedicated staff and of building and equipment were not evaluated
- b Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

in 2005. International leprosy organizations have provided technical assistance for TB control for more than a decade.

## **Budgets and expenditures**

The NTP budget increased from US\$ 8.6 million in 2002 to US\$ 12 million in 2005. However, funding from both the government and donors has been declining since 2003, and in 2005 the funding gap is expected to be around US\$ 7 million, equivalent to 57% of the budget. There are two main reasons for persistent funding gaps. One is that, while a GFATM grant was approved in January 2003, this

was subsequently revoked because of lack of counterpart funds from the government. In 2003, a second reason was that funding from the government was planned at US\$ 3.9 million but reached only US\$ 1.9 million. The largest budget line item each year between 2003 and 2005 is for expansion of DOTS to new LGAs (included in the line item "initiatives to increase case detection and cure rates"). Dedicated TB staff, first-line drugs and buildings and equipment are also relatively large budget items. The budget per patient treated has ranged from US\$ 160 to US\$ 300. Actual expenditures in 2003 were US\$ 5.6 million (equivalent to US\$ 131 per patient treated), slightly lower than the available funding of US\$ 6.0 million.

Total TB control costs, including visits to health clinics and spending on dedicated TB hospital beds as well as items covered by the NTP budget, are estimated at US\$ 12 million in 2003 (about US\$ 300 per patient treated). If the budget gap for 2005 is filled and the number of patients treated increased to nearly 73 000 as projected, then total TB control costs would reach about US\$ 20 million in 2005 (also about US\$ 300 per patient treated).

# **Pakistan**

DOTS coverage has increased rapidly in Pakistan since 2000, reaching 63% in 2003. With plans to include the remaining districts, nationwide DOTS coverage should be achieved in 2005. Pakistan has been highly successful in mobilizing financial support for TB control from the international community, and this has given impetus to the programme. The NTP is well structured and has created a strong TB control network during the past five years, with an effective mechanism for coordinating a range of activities and partnerships. Both case detection and treatment outcomes are improving, but remain below the global targets at 17% and 77% respectively. Recent health sector reforms give increased responsibility to the districts for setting priorities for health programmes and to the NTP for ensuring that TB control is a priority at district level. As the programme advances towards nationwide DOTS coverage, the NTP will have to respond to the increasing demand for anti-TB drugs, equipment and reagents, and to ensure that the quality of the services continues to improve.

#### System of TB control

The NTP is responsible, under the MoH, for the overall coordination of TB control in the country. The specific responsibilities of the NTP include formulation of policy, strategic planning, technical support and supervision, monitoring and evaluation, coordination and communication with partners and research. The provincial and regional TB control managers are responsible for planning, implementing, monitoring and evaluating TB control activities in each province and region. However, districts serve as the main administrative units for the programme; the district authorities are primarily responsible for activities at that level. District hospitals and rural health centres provide diagnostic and treatment services; the basic health units and dispensaries provide treatment. In rural areas, "lady health workers" play an important role in referring TB suspects from communities and in providing DOT. In some big cities, treatment is not yet provided in all health

Pakistan has one national, four provincial and two regional reference laboratories. The national laboratory and three of the provincial laboratories have facilities for culture and drug susceptibility testing. In the districts, 619 diagnostic centres do microscopy.

#### **Surveillance and monitoring**

No national survey of TB infection or disease has been carried out in Pakistan, and case notifications were erratic until the introduction of DOTS in the early 1990s. The incidence of TB and its trend are uncertain. DOTS coverage increased rapidly from 9% in 2000 to 63% by 2003. During the same period, the smear-positive case detection rate increased from 3% to 17%. While these two indicators have increased, their ratio has not changed, suggesting that the case detection rate within DOTS areas has stayed in the range 20-30% since 2000. A possible reason for the low rate of case detection is that only 30% of all notified TB cases were diagnosed as smear-positive in 2003. Since it is expected that about 45% of incident cases would be smear-positive, the low proportion of reported smearpositives suggests that some smearpositive cases may have been notified as smear-negative.

The treatment success rate in the 2002 cohort was 77%, similar to that in 2001; the default rate remained high at 14%. Furthermore, 13% of treated patients, who were counted as successfully treated, completed treatment without evidence of smear conversion. Information on treatment success outside the DOTS programme is not available. Among relapse cases treated under DOTS, the treatment success was high (81%), but the proportion of patients whose cure was not laboratory confirmed was even higher than among new patients (53%). Among patients who had defaulted on previous treatment, treatment success was only 58%, mostly as a result of patients defaulting again (22% of the cohort).

## **PROGRESS IN TB CONTROL IN PAKISTAN**

#### **Indicators**

DOTS treatment success, 2002 cohort 77% DOTS detection rate, 2003 17% NTP budget available, 2004 27% Government contribution to NTP budget, including loans, 2004 7% Government contribution to total TB control costs, including loans, 2004 26% Government health spending used for TB control, 2004 5%

#### **Major achievements**

- Rapid DOTS expansion to cover a total of 94 out of 121 districts
- Provincial and district capacity-building to improve monitoring and supervision
- Establishment of the National Pakistan Stop TB Partnership to increase TB awareness and political commitment of local authorities

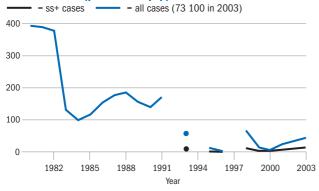
#### **Major planned activities**

- Expand DOTS to cover all districts by 2005
- Develop an EQA system for smear microscopy
- Implement PPM-DOTS through FIDELIS and GFATM funding
- Launch communication strategies to improve TB awareness among health-care providers and the public

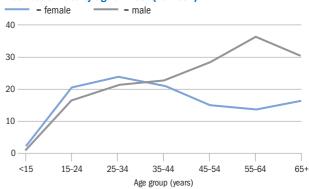
## **PAKISTAN**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	153 577 848	DOTS coverage (%)	9.0	24	45	63
Global rank (by est. number of cases)	6	Notification rate (all cases/100 000 pop)	7.7	23	35	48
Incidence (all cases/100 000 pop/year)	181	Notification rate (new ss+/100 000 pop)	2.3	7.5	11	14
Incidence (new ss+/100 000 pop/year)	82	Detection of all cases (%)	4.3	13	19	26
Prevalence (all cases/100 000 pop)	359	Case detection rate (new ss+, %)	2.8	9.2	13	17
TB mortality (all cases/100 000 pop/year)	43	DOTS case detection rate (new ss+, %)	2.8	5.2	13	17
TB cases HIV+ (adults aged 15-49, %)	0.6	DOTS case detection rate (new ss+)/coverage (%)	31	22	28	27
New cases multidrug resistant (%)	9.6	DOTS treatment success (new ss+, %)	74	77	77	_

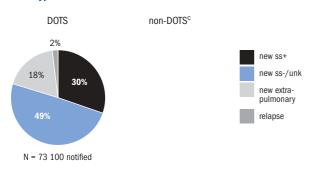
#### Notification rate (per 100 000 pop)



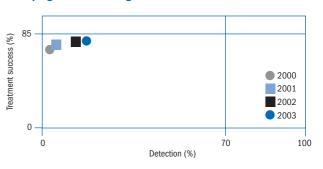
#### Notification rate by age and sex (new ss+)<sup>b</sup>



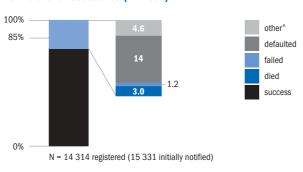
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

### **Improving programme** performance

Under the recent health reforms, district governments were authorized to prioritize their district health needs. The NTP needs to ensure that districts take ownership of their local TB control effort and make it a priority. The new National TB Control Programme Plan for 2006–2010 and Provincial TB Control Programme Strategic Plans for each of the four provinces for the same period have been drafted. These will be used to advocate for TB control at the national, provincial and district levels. The induction of national programme officers through USAID funding has helped to develop provincial and district capacity for monitoring and supervision.

The Government of Pakistan is committed to TB control under the DOTS strategy, and the programme is receiving adequate attention from policymakers, as evidenced by the rapid expansion of DOTS since 2000. DOTS coverage will be expanded to the remaining 20 districts in 2005. With the rapid expansion of DOTS, the NTP faces constraints including inadequate public sector resources. As coverage is increased, the new national plan will progressively focus on the quality of care, enhanced case detection, monitoring and supervision and activities to de-stigmatize the disease. The Pakistan Stop TB Partnership is being launched and has appointed a Stop TB Ambassador. This is the first initiative in Pakistan to include non-traditional partners in TB control activities.

In 2002, the NTP received a twoyear grant from the GDF, and this was extended to cover 2005. However, the current level of drug procurement will not be sufficient to meet the increasing needs arising from rapid DOTS expansion. As the use of FDCs is being advocated in the four provinces, the NTP has revised the treatment guidelines and has drafted training materials on their use. There are no drug resistance data available for Pakistan, although WHO estimates a prevalence of MDR in new TB patients of 10%. Patients in whom MDR-TB is diagnosed are not treated under the NTP.

The NTP has recognized the importance of behaviour change, communication and community mobilization in achieving countrywide implementation of DOTS, and support from various donors has been sought to develop effective strategies. Television spots, posters, leaflets, videos and other materials have been developed to raise public awareness. These strategies will be launched in 2005, with the aim of spreading public awareness among both health-care providers and the general public. Innovative approaches, coupled with operational research, are being explored to involve non-traditional partners such as politicians, industrialists, local district governments and religious leaders in TB control activities.

Three areas in which programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

The NTP plans to establish an intermediate level laboratory network consisting of one reference laboratory in each district in 2004-2005 and to expand the number of hospitals and rural health centres that serve as diagnostic centres. The national and provincial reference laboratories have been strengthened with the procurement of laboratory equipment, materials and vehicles. Supervision and overall support for the provincial laboratories need further strengthening; guidelines need to be developed as well as tools for supervision, appropriate for the country setting. There are currently no systems in place for quality assurance of microscopy services at the district level, and this is a priority in the 2006–2010 national TB control plan. Other needs include training of laboratory staff and improvement in laboratory operating procedures.

#### TB/HIV coordination

The prevalence of HIV in the general population appears to be low, but the lack of adequate epidemiological data precludes an accurate assessment of the HIV situation in Pakistan. A TB/ HIV plan and a national TB/HIV coordinating body are both being developed. TB/HIV awareness activities have been undertaken in conjunction with the South Asian Association for Regional Cooperation (SAARC) TB/HIV Awareness Year (2004).

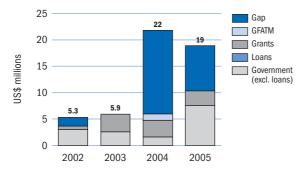
#### Links with other health-care providers

Pakistan has developed a national strategy for PPM DOTS. Few initiatives have been launched so far, but there is a strong commitment to encourage the active involvement of more healthcare providers, including governmental, semigovernmental and the private sector, in DOTS expansion. Funds from the GFATM are being used to expand PPM and BCC (behaviour change and communication) activities. Several FIDELIS projects linking the NTP to other health-care providers are planned, including improving TB case detection by encouraging intersectoral collaboration in three urban areas and strengthening DOTS implementation in four districts of Punjab. NGOs are involved in some districts of each province and territory of the country; these include the Abasseen Foundation, Aga Khan Foundation, Asia Foundation, Association for Social Development, Marie Adelaide Leprosy Center, Mercy Corps International, Pakistan Anti-TB Association and many other local NGOs

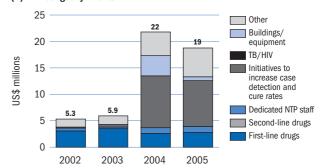
#### **Partnerships**

Partnerships for TB control in Pakistan have been strengthened, and technical and financial support has increased significantly. The NTP has launched the National Pakistan Stop TB Partnership to increase TB awareness and the political commitment of local authorities. Major technical partners include GLRA, GTZ, IUATLD, JICA and WHO. DFID has offered assistance to develop PPM partnerships; USAID has provided support to strengthen the capacity for DOTS implementation in the districts. The governments of Canada, Germany and Japan are the main financial partners for TB control activities. Pakistan will receive ISAC initiative funding (through CIDA) for DOTS expansion and sustainability through the involvement of district governments.

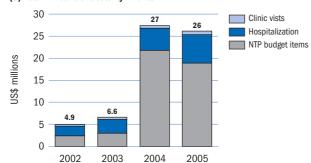
#### (a) NTP budget by source of funding



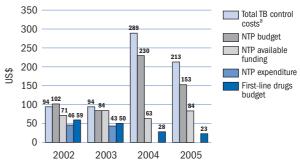
#### (b) NTP budget by line item











Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

#### **Budgets and expenditures**

The budgets for both 2004 and 2005 are substantially higher than in previous years, at about US\$ 20 million compared with US\$ 5-6 million in 2002 and 2003 (US\$ 153 per patient in 2005 compared with around US\$ 100 in both 2002 and 2003). These budgetary increases reflect the development of more ambitious plans to accelerate DOTS expansion (to achieve 100% coverage by 2005) and to increase case detection and cure rates throughout the country, and the associated revision of existing national and provincial strategic plans and budgets in 2004 (most of the budgetary increase between 2003 and 2004 is in the category "initiatives to increase case detection and cure rates"). The revised plans include PPM-DOTS strategies and community mobilization activities.

While the funding available for 2004 and 2005 is similar to that in 2002 and 2003, the considerably increased budgets for 2004 and 2005 mean that large funding gaps currently exist: US\$ 16 million in the fiscal year 2004 and US\$ 8.6 million in 2005 (the fiscal year starts in July). The NTP is already engaged in efforts to mobilize funds to fill these gaps. For example, it is expected that PPM-DOTS strategies will be funded through the national health and population facility (this is supported by the Pakistani government and DFID), and a further application to the GFATM is planned. In addition, provinces are revising their budgetary allocations in the context of the revised strategic plans. Some positive results are already apparent: in December 2004, the Punjab government approved a revised three-year budget allocation of US\$ 8.6 million

for TB control activities in the province, including US\$ 2.4 million for the first year (of which US\$ 1.3 million is for drugs).

If the revised NTP budget is fully funded, the total cost of TB control (including health clinic visits for observation of treatment and monitoring and limited hospitalization as well as NTP budget items) will increase from around US\$ 5 million in 2002 to US\$ 26 million in 2005 (and from around US\$ 100 to US\$ 213 per patient treated). It remains to be seen whether increased funding can be absorbed effectively and whether increased expenditures result in improved case detection and cure rates.

# **Philippines**

The Philippines achieved full DOTS coverage in 2003, has met the global target for treatment success in each of the past four years and is coming close to the target for case detection. TB control has progressed thanks to strong government commitment and a relatively well-staffed programme, while innovative partnership arrangements are making important contributions to TB control activities and resource mobilization. The financial position is favourable, with the budget for TB control activities fully funded for 2004 and 2005. The use of barangay (small local district) health workers to treat and follow patients has been a very beneficial national policy that has helped to achieve high treatment success rates. Involving medical schools and private physicians in DOTS services is now a government priority because this will increase case detection and ensure that standard methods for diagnosis and treatment are used in the private sector. Surveillance for TB drug resistance is in progress. The Philippines is one of the few high-burden countries that has started to implement DOTS-Plus treatment for MDR-TB

cases. Providing TB control and other health services to population groups in remote mountainous areas and small islands, and accessing insecure areas, present continuing challenges.

#### **System of TB control**

The NTP has recently been reorganized as part of the national health sector reform process. Following restructuring and considerable decentralization of the Department of Health, the number of staff at central level was substantially reduced. Although additional staff have subsequently been employed at central level, there are still too few to carry out regular monitoring of programme activities in the regions. This means that regional coordinators are now responsible for most coordination and technical assistance, even though they may be responsible for more than one health programme and thus have limited time for TB control activities. Fortunately, the number of staff in the provinces and in rural health units is sufficient, and most staff have adequate training in all aspects of TB control. Each of the country's 16 regions has a centre for health development that provides technical support to the provincial health offices. Provincial TB coordinators supervise staff in the rural health units, which are the main focus of TB control in the Philippines.

The TB laboratory network is structured as follows: the NRL is responsible for developing policy, management, training of microscopists, supervision of intermediate laboratories and DRS. Regional and provincial laboratories implement the policies developed by the NRL and provide EQA to the peripheral laboratories. The primary role of the peripheral laboratories at the rural and city health units is sputum smear microscopy. Culture and drug susceptibility testing are carried out by the NRL, one private laboratory and one NGO-affiliated laboratory. Seven regional laboratories have the capacity to perform culture.

#### **Surveillance and monitoring**

The TB case notification rate was decreasing before 2001 but has increased slightly since then. In 2003, as in previous years, the highest notification rates were among adults aged 45 years and older. These observations suggest that the TB incidence rate is probably in decline in the Philippines, with this reduction obscured since 2000 by DOTS expansion and the greater effort given to case-finding. The DOTS case detection rate increased rapidly to 48% in 2000 and then more slowly to 68% in 2003. Treatment success was reported as 88% in the 2000, 2001 and 2002 DOTS cohorts, and 91% of new smearpositive cases notified in 2002 were registered for treatment in that year.

With the public sector DOTS programme nearing full implementation, greater efforts are being made to diagnose and treat patients in collaboration with the private sector. The NTP must now also consider how to evaluate the epidemiological impact of the DOTS programme. Two prevalence surveys were done in the Philippines before the implementation of DOTS

## PROGRESS IN TB CONTROL IN THE PHILIPPINES

#### Indicators

DOTS treatment success, 2002 cohort	88%
DOTS case detection rate, 2003	68%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	36%
Government contribution to total TB control costs, including loans, 2004	82%
Government health spending used for TB control, 2004	3%

#### **Major achievements**

- Scaling up of PPM DOTS in two thirds of medical schools and more than 2000 private providers to increase the case detection rate
- Nationwide implementation of FDC anti-TB drugs, with increased health-worker
- TB control in children was piloted in urban and rural areas, and TB control in high-risk populations started

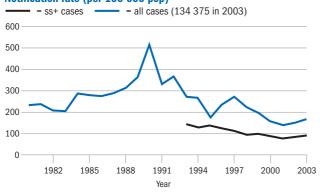
#### **Major planned activities**

- Establish additional PPM-DOTS sites to cover the entire country
- Implement EQA for smear microscopy nationwide, including hospitals and private laboratories
- Strengthen the national reference laboratory function in laboratory networking

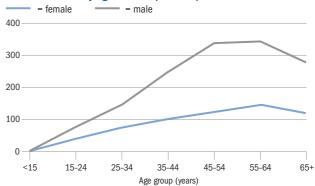
## **PHILIPPINES**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	79 999 016	DOTS coverage (%)	90	95	98	100
Global rank (by est. number of cases)	9	Notification rate (all cases/100 000 pop)	158	139	151	168
Incidence (all cases/100 000 pop/year)	296	Notification rate (new ss+/100 000 pop)	89	77	83	91
Incidence (new ss+/100 000 pop/year)	133	Detection of all cases (%)	52	46	50	57
Prevalence (all cases/100 000 pop)	458	Case detection rate (new ss+, %)	65	57	62	68
TB mortality (all cases/100 000 pop/year)	49	DOTS case detection rate (new ss+, %)	48	57	62	68
TB cases HIV+ (adults aged 15-49, %)	0.1	DOTS case detection rate (new ss+)/coverage (%)	54	60	63	68
New cases multidrug resistant (%)	3.2	DOTS treatment success (new ss+, %)	88	88	88	_

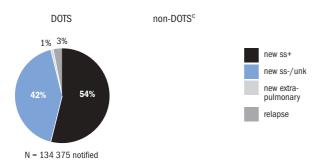
## Notification rate (per 100 000 pop)



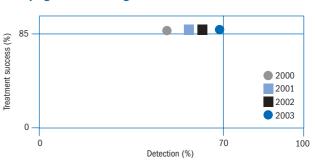
#### Notification rate by age and sex (new ss+)<sup>b</sup>



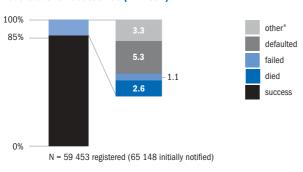
#### **Case types notified**



## **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

that showed little reduction in culturepositive or smear-positive disease between 1981-1983 and 1997. A new national TB prevalence survey is scheduled for 2007. This will show whether or not the Philippines can meet, or has already met, the Millennium Development Goal of halving prevalence between 1990 and 2015.

### Improving programme performance

The Philippines reached 100% DOTS coverage in 2003 as a result of strengthened DOTS expansion efforts, backed by government commitment and funding for TB control as a priority public health programme. As part of the health sector reform process, management capacity and programme infrastructure were upgraded, and TB control activities became the responsibility of the Infectious Diseases Office under the National Centre for Disease Control and Prevention. Following the reorganization of the Department of Health, the procedural manual for the NTP and the Comprehensive and Unified Policy for TB Control in the Philippines will be revised. This policy provides a framework for collaboration with other government agencies and with the private sector, which in turn will help to harmonize and unify TB control efforts in the Philippines.

Nationwide implementation of FDC anti-TB drugs started after successful training for health-care workers. To improve case detection, TB control initiatives focused on children were pilot tested in urban and rural areas. and TB control activities in high-risk populations begun. A TB outpatient benefit package, PhilHealth, was introduced to improve treatment success rates.

The first nationwide DRS survey started in June 2003. This will provide the first reliable estimate of the magnitude of MDR-TB in the country. In 2000, the GLC approved a DOTS-Plus project at Makati Medical Center in Manila (a private medical centre collaborating with the NTP), with an initial cohort of 200 patients. With support from the GFATM, this cohort has been expanded to 750 MDR-TB patients in 2004. As yet, no MDR-TB

patients are treated in the public sector.

In addition to management of MDR-TB, diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers could also be improved.

#### Diagnostic and laboratory services

The laboratory service at intermediate and peripheral levels is good and staff are well-trained; however, EQA for sputum smear microscopy is not in place in every laboratory and needs to be strengthened where it already exists. An updated national manual for EQA for direct sputum smear microscopy was developed and distributed in late 2004. A priority for 2005 is to establish models for EQA, to monitor and evaluate the model EQA implementation and to expand to other laboratories. Eventually, hospitals and private laboratory facilities will be included in EQA activities. Laboratory networking is to be developed at all levels of the health service; successful networking will require reinforcement of the NRL.

#### TB/HIV coordination

There are no existing data on  $\ensuremath{\mathsf{TB}}/\ensuremath{\mathsf{HIV}}$ coinfection in the Philippines. However, HIV prevalence in the general population, and among TB patients, remains low (<1%). Given the worsening HIV/AIDS epidemics in neighbouring countries, it is important to monitor HIV prevalence in the general population as well as among high-risk groups including TB patients.

## Links with other health-care providers

With the aim of consolidating and scaling up initiatives to involve private health-care providers in DOTS, the Philippines issued guidelines on PPM DOTS in 2004, and a national committee for PPM DOTS has been established. Operational guidelines for PPM DOTS in the Philippines were published, endorsed by the Secretary of Health and distributed. PPM-DOTS units, whose role is to coordinate private sector involvement in provision of DOTS services, have been set up in over 50 sites nationwide. More than 2000 private providers have been trained, and six professional societies have introduced the DOTS strategy in

their training curricula. Two thirds of medical schools have become or are in the process of becoming involved in DOTS activities. The positive impact of these initiatives on case detection has been demonstrated in a few sites, but there is a need to incorporate a careful and more comprehensive strategy for monitoring and evaluation of the current scale-up of PPM DOTS in the Philippines.

#### **Partnerships**

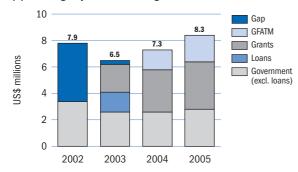
The Philippines benefits from several partnerships that strengthen the programme and support DOTS expansion. Overall external technical collaboration is coordinated by WHO. Other external technical support is provided by CDC, JICA, KNCV, Medicos del Mundo (Spain), USAID and World Vision, which has helped to maintain technical quality during the expansion phase. An important innovation led by the Department of Health is the organization of the Philippines Coalition Against TB (PhilCAT). This includes a substantial group of NGO and private sector entities that collaborate to help private sector TB control activities and to mobilize local resources. The major funding partners are CIDA, GFATM, JICA and USAID.

### **Budgets and expenditures**

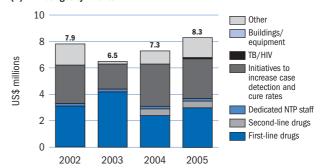
The budget specifically for TB control activities has been similar in the four years 2002-2005, at about US\$ 7-8 million. However, funding gaps existed in 2002 and 2003, whereas no funding gap has been reported for 2004 or 2005. This improved funding situation is linked to an increasing level of grant funding, much of which is related to initiatives to increase case detection - in particular, USAID funds for PPM DOTS. Funding from the government has fallen, related to austerity measures that affect public spending as a whole. In contrast to most other HBCs, there is also a budget for second-line drugs in 2004 and 2005, linked to implementation of DOTS-Plus in Manila. On a per patient basis, the overall NTP budget has fallen from US\$ 66 in 2002 to US\$ 48 in 2005. This is mainly explained by a reduction in the cost of first-line drugs, which has fallen from US\$ 26 per pa-

## **PHILIPPINES**

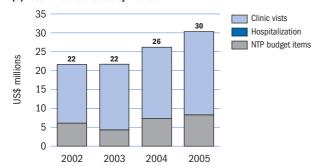
#### (a) NTP budget by source of funding



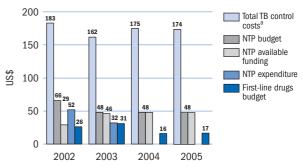
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

tient treated in 2002 to US\$ 17 per patient treated in 2005. When costs beyond those reflected in the budget specifically for TB control are also included, i.e. health clinic visits for DOT and monitoring during treatment, the cost per patient has been about US\$ 160-180 for the past four years, and total TB control costs have been around US\$ 22-30 million per year.

# **Russian Federation**

Following a considerable rise in the TB burden in the Russian Federation during the 1990s, a peak was reached in 2000 when some 132 000 cases of TB were notified. Since then, a progressive reduction in the number of reported cases has occurred, mainly because of a decline in the number of cases registered in the prison sector. However, the number of cases in the general population has increased, particularly among children. The DOTS strategy is not widely used in the Russian Federation. In oblasts where it is being applied, both case detection and treatment outcomes are still low. However, government commitment to TB control is strong, and a recent World Bank loan will allow accelerated expansion of both TB and HIV/AIDS programmes. In addition, the Russian Federation made a successful application to the GFATM in 2004, opening up additional opportunities to extend and improve these programmes. The

increasing public health importance of TB/HIV coinfection is being addressed through a national TB/HIV coordinating body, which has developed a national strategy for TB/HIV control. Also receiving special attention is the growing MDR-TB epidemic in the Russian Federation; links between the MDR-TB and TB/HIV epidemics are being investigated. A major challenge is to improve the laboratory network to meet international standards and provide reliable diagnostic services for the TB control programme.

#### **System of TB control**

The Russian Federal Target Programme "Prevention and Control of Social Diseases (2002-2006)", with the subprogramme "Urgent Measures of TB Control in Russia" was approved in 2001. The Programme aims to stabilize the epidemiological situation of social diseases through improvement of current organizations and newly

established services. The plan covers strengthening the capacities of health facilities, research institutes and centres that carry out prevention, timely detection, diagnosis and treat-

Several federal laws and regulations were developed to strengthen the foundation of the TB control programme. The national five-year plan, "Provision of guaranteed diagnostic and treatment procedures for TB patients and the development of TB services in Russia (2003-2006)", was developed as the main framework for activities and cooperation with international partners. Reduction of TB incidence, disability and mortality is currently one of the priorities of state policy in the Russian Federation.

Within the federal TB control programme, five research institutes are responsible for organizing and supervising research, training and implementation of TB control in a wide network of more than 500 TB control facilities in 88 regions of the Russian Federation. These are the Research Institute of Phthisiopulmonology of Sechenov Moscow Medical Academy (RIPP MMA), the Central TB Research Institute of the Russian Academy of Medical Sciences (CTRI RAMS), St Petersburg Institute of Phthisiopulmonology, Ural Research Institute of Phthisiopulmonology and the Novosibirsk TB Research Institute. The TB dispensaries in turn supervise and monitor regional TB hospitals, sanatoria and TB units at district polyclinics. Under the Ministry of Justice, 37 hospitals and 57 treatment facilities provide treatment for TB patients within the penitentiary system.

The five federal TB research institute laboratories and 377 TB dispensary laboratories perform culture and drug susceptibility testing. In the territories, 348 centres with hospitals and sanatoria perform culture, and more than 11 000 centres perform smear microscopy.

### PROGRESS IN TB CONTROL IN THE RUSSIAN FEDERATION

#### **Indicators**

DOTS treatment success, 2002 cohort	67%
DOTS case detection rate, 2003	8.8%
NTP budget available, 2004	84%
Government contribution to NTP budget, including loans, 2004	83%
Government contribution to total TB control costs, including loans, 2004	87%
Government health spending used for TB control, 2004	4%

#### **Major achievements**

- Beginning of implementation of the AIDS and TB control project funded by a World Bank loan
- Approval by the MoH of the new recording and reporting system, including cohort analysis, introduced in 37 regions in 2004 and countrywide in 2005
- Successful application to the GFATM round 4 for TB control
- Development of a strategy on TB/HIV control and countrywide training of regional TB/HIV coordinators
- Training of trainers in the revised TB control strategy
- Substantial progress in the Thematic Working Group on MDR-TB control

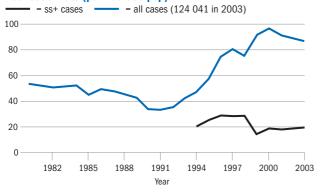
#### **Major planned activities**

- Expand the revised TB control strategy through the World Bank loan project and prepare for GFATM project implementation
- Develop national guidelines and a framework for the management of MDR-TB
- Implement TB/HIV control strategy
- Strengthen laboratory system: capacity development for smear, culture and drug susceptibility testing; establish the national reference laboratories network and quality assurance system; implement drug resistance surveillance in 10 oblasts
- Improve anti-TB drug supply system

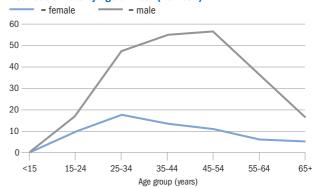
## **RUSSIAN FEDERATION**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	143 246 223	DOTS coverage (%)	12	16	25	25
Global rank (by est. number of cases)	12	Notification rate (all cases/100 000 pop)	97	91	89	87
Incidence (all cases/100 000 pop/year)	112	Notification rate (new ss+/100 000 pop)	19	18	19	20
Incidence (new ss+/100 000 pop/year)	50	Detection of all cases (%)	79	76	77	77
Prevalence (all cases/100 000 pop)	160	Case detection rate (new ss+, %)	35	34	37	40
TB mortality (all cases/100 000 pop/year)	20	DOTS case detection rate (new ss+, %)	4.6	5.2	6.9	8.8
TB cases HIV+ (adults aged 15-49, %)	6.2	DOTS case detection rate (new ss+)/coverage (%)	39	33	28	35
New cases multidrug resistant (%)	6.0	DOTS treatment success (new ss+, %)	68	67	67	_

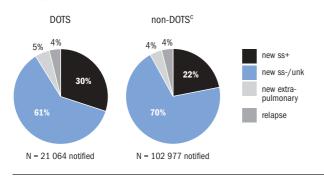
#### Notification rate (per 100 000 pop)



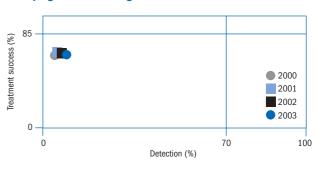
### Notification rate by age and sex (new ss+)b



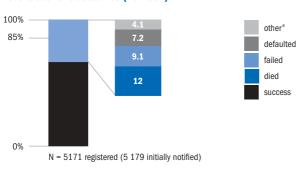
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

#### **Surveillance and monitoring**

The increase in the number of annual TB case notifications after 1990 reached a peak of around 141 000 cases in 2000, and the number of reported cases has fallen each successive year since then. Similar trends have also been observed in other countries of the former Soviet Union that have also reported fewer cases, or at least a slowing in the rate of increase in the number of patients. This stabilization in notification rates in the Russian Federation could be the result of improved TB control or of general improvements in peoples' health, but is most likely because of the decline in notified cases over the past few years observed in the prison sector. Since 2001, a decline of notified cases in the prison sector has occurred from more than 24 000 cases to around 16 000 in 2003. TB mortality rates remain increasingly high at around 20 per 100 000 population.

In 2003, DOTS coverage was low, with only 8.8% of cases detected under DOTS. It is therefore unlikely that the DOTS strategy had a major impact on incidence. In 2004, DOTS was implemented in 37 regions of the Russian Federation, with increased coverage to around 45% of the population; however, detection and treatment outcomes remain suboptimal. Moreover, treatment success in the 2002 cohort was low even in DOTS areas (67%) because many patients died (13%), failed treatment (9%) or were lost to follow-up (11%). The treatment outcomes for new smear-positive patients under DOTS have not improved in eight successive cohorts (1995-2002). Among the 962 DOTS relapse cases in 2002, fewer than half were successfully treated (46%), mainly because 26% failed re-treatment.

Treatment outcomes are not available for re-treatment after default or failure. Given the high prevalence of MDR-TB in the Russian Federation, it is important that these data be collated and analysed in future. Although sputum smear microscopy is increasingly used for diagnosis, the proportion of new pulmonary TB patients with a positive sputum smear was still only 33% in 2003 in DOTS regions and did not exceed 24% in non-DOTS regions,

with an overall average for the Russian Federation of 25.3%. Nevertheless, the Russian Federation is different from many other HBCs in having a fairly comprehensive system for recording and reporting the total numbers of TB cases and deaths. This system of routine surveillance (rather than population-based surveys) should, with some refinements, be adequate for monitoring epidemiological trends and the future impact of TB control

### Improving programme performance

The current state policy aims to stabilize and improve the epidemiological situation, which is evidenced by an increase in federal budget allocations for TB control. The commitment of the federal government to TB control continues to grow, with sustained activities of the high-level working group (HLWG), one of the mechanisms of international cooperation in the field of medicine. The HLWG comprises representatives of the Ministry of Health and Social Development of the Russian Federation, the Ministry of Justice, RIPP MMA, CTRI RAMS, WHO and the Council of Europe. The federal government has adopted a number of regulations for TB control including: the Executive Order No. 109 of 21 March 2003 "On Improvement of TB Control in the Russian Federation" that focuses on laboratory diagnosis, chemotherapy standards, organization of treatment, prevention of TB transmission, system of centralized control and management of main TB interventions at the level of TB facilities in regions of the Russian Federation, and the introduction of the new reporting and dispensary follow-up system; the Executive Order No. 50 of 13 February 2004 "On Implementation of Registration and Reporting Documentation for Tuberculosis Monitoring" that includes cohort analysis and assessment of detection and treatment effectiveness in line with international standards; a recording form "Individual Card of TB/HIV Patient"; and recommendations on decreasing TB burden among high HIV prevalence populations.

Registers for recording and report-

ing TB based on cohort analysis were introduced by the Russian MoH in February 2004. From April 2004, new reporting forms were being introduced in 37 territories in both the civil and penitentiary sectors, and will be used country-wide from January 2005.

The activities of the federal TB control programme and expansion of the revised strategy are constrained by the shortage of staff and age of the existing medical staff working in TB services, many of whom are retiring. A detailed assessment of HR needs is under way and several activities are in progress to address HR capacity, including a staff development plan, as part of the overall TB plan for 2003-2007, and further training of TB service personnel supported by the World Bank-funded TB/AIDS project. New national guidelines and recommendations have been developed, published and distributed on case detection, TB treatment, laboratory services and TB/HIV control.

MDR-TB is a major challenge for TB control in the Russian Federation. MDR-TB patients outside the DOTS-Plus projects are treated on an individual basis and according to the availability of second-line drugs. Data on the prevalence of drug resistance are reported routinely from Ivanovo, Orel and Tomsk oblasts where the prevalence of MDR-TB among new cases ranges from 2.6% in Orel to 13.7% in Tomsk. Data from a few additional oblasts will be available shortly, and a plan to survey oblasts systematically is being developed. GLC-approved DOTS-Plus projects are being implemented in Archangelsk, Ivanovo, Orel and Tomsk. The GLC has approved the treatment of 2830 MDR-TB patients. The project in Tomsk, which was the first of these projects to start, has been successful in treating MDR-TB patients and has recently been expanded with financial support from the GFATM.

#### Diagnostic and laboratory services

The physical infrastructure of many diagnostic facilities in the Russian Federation does not meet Russian and international standards for laboratory design and safety. In addition, in many instances, laboratory equipment is

## **RUSSIAN FEDERATION**

outdated. Updating infrastructure of existing facilities and ensuring availability of quality equipment and supplies is an enormous challenge facing the national TB control programme.

Quality assurance is being addressed by the introduction of a federal system of EQA for smear microscopy approved by the MoH. However, given the financial constraints, the system has not yet been introduced in all diagnostic centres, nor have internal quality control procedures. The TB laboratory network faces a serious shortage of staff; HR capacity building through training and development of a model for effective laboratory services at the central level should lead to improvements in laboratory diagnosis at all levels.

#### TB/HIV coordination

HIV/AIDS is becoming a significant public health problem in the Russian Federation. A thematic working group, "TB in HIV-infected people", has been established within the HLWG on TB and has developed recommendations on decreasing the TB burden among PLWHA. The group comprises leading national TB and HIV experts from re-

search institutes, health facilities, WHO and international partners. Its basic objective is to develop a framework for establishment of the national system of TB care among HIV-infected people. The first stage resulted in the preparation of the "Recommendations on decreasing TB burden in high HIV prevalence populations" based on national and international practices.

Regional TB/HIV coordinators have been appointed in many regions of the Russian Federation. A number of federal-level seminars were held in 2004 where these coordinators were trained in principles of the newly developed TB/HIV strategy.

#### Links with other health-care providers

Collaboration between all relevant public sector health-care providers and related institutions is being strengthened, including general hospitals, TB hospitals, medical colleges, prison health services and the health services of the armed forces and of the police. The Ministry of Railway Communication, the Federal Security Service and a number of other ministries and departments have their own TB control services, and links with them

need to be strengthened. The private sector plays a minor role in TB diagnosis and treatment.

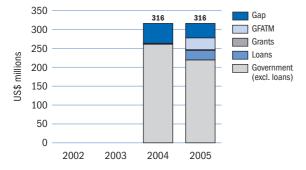
#### Links with the community

Several community groups contribute to the provision of TB control activities in the Russian Federation, including an NGO of TB patients (NABAT), the Russian Red Cross, Russian TB Society and other regional foundations and societies. These groups participate in annual World TB Day campaigns and provide health education and social support for TB patients.

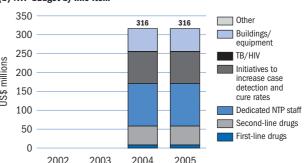
#### **Partnerships**

The HLWG, established in 1999, continues to play an important role in the development of TB control. It is responsible for coordinating TB activities between the national and international partners, and it works on recommendations for executive policy documents (prikaz) that regulate implementation of national TB control. Many national and international NGOs and technical agencies are partners in TB control within the 88 territories of the Russian Federation. Major donors include USAID, the Swedish

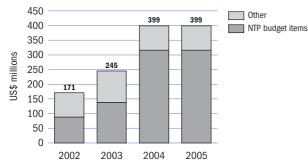
#### (a) NTP budget by source of funding



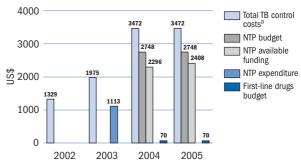
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



a Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. "Other" includes costs for hospitalization and fluorography not reflected in the budget estimates submitted to WHO

## **RUSSIAN FEDERATION**

International Development Agency, the EU, the Government of Finland and DFID. The Russian Federation successfully applied to the GFATM in round 4; funds for TB control activities should be available in 2005. A loan agreement between the Russian Federation and the World Bank to fund the project on "AIDS and TB Control" was signed in September 2003 and became effective in December 2003.

#### **Budgets and expenditures**

Financial data were prepared by WHO staff (Moscow office) using data available in the public domain, and are therefore estimates rather than official figures. Sources of data included the Ministry of Health and Social Development and the Federal Agency for Health Care and Social Development of the Russian Federation.

The total budget for TB control in both 2004 and 2005 is estimated at US\$ 316 million (almost US\$ 3000

per new TB patient). About US\$ 250-260 million is available from the government in both 2004 and 2005 (including funds from a World Bank loan), a substantial increase compared with 2003. In 2005, the GFATM is expected to provide a further US\$ 30 million, but grants from other sources are limited. While the available funding of about US\$ 270 million in 2004 and 2005 is substantial by the standards of other HBCs, a funding gap of about US\$ 40-50 million has been estimated for both years, primarily for the purchase of secondline drugs (US\$ 18 million) and for investment in buildings and equipment (US\$ 24 million). The Russian Federation accounts for almost one third of the total funding gap reported by the 22 HBCs.

The largest budget line items are for staff working exclusively on TB control (US\$ 113 million in both years), initiatives to increase case

detection and cure rates (US\$ 84 million in both years), investment in buildings and equipment (US\$ 60 million in both years) and second-line drugs (about US\$ 45 million in both years). The budgets for staff, investment in buildings and equipment and secondline drugs are relatively large compared with those in other HBCs, and reflect the country's extensive network of dedicated TB control facilities and the large number of patients with MDR-TB. When costs beyond those reflected in the reported budgets are included (i.e. the operating costs of a network of 81 425 dedicated TB beds and the cost of mass screening using fluorography), the total cost of TB control is estimated to be about US\$ 400 million in both 2004 and 2005 (about US\$ 3500 per patient treated), up from an estimated US\$ 245 million in

# South Africa

In 1996, South Africa established an NTP and adopted DOTS as its TB control strategy. Despite government commitment to making TB control a priority, and the implementation of the DOTS strategy in all provinces and almost all districts, it is not known with confidence how much TB there is in the country. Inadequate case reporting systems, a shortage of trained staff at the provincial level and problems associated with the laboratory network hinder effective TB surveillance. Recognizing these shortcomings, the NTP has recently taken a number of steps to remedy the situation, and better data can be expected in the future. However, treatment success rates remain low and many patients are lost to follow-up. A concerted effort will be needed if South Africa is to reach the target cure rates. TB/HIV coinfection is a significant public health problem and is being addressed through a national programme of collaborative TB/HIV activities. MDR-TB prevalence is estimated to be about 2% in new TB patients and 7% in re-treatment cases. Second-line

drug treatment is available in provincial MDR-TB units, though at high cost. A number of NGOs are involved in providing TB services and are also mobilizing support in the communities, but more needs to be done to encourage broader private sector participation.

#### **System of TB control**

South Africa's health system is decentralized. The National Department of Health provides general guidelines, but the implementation and delivery of services is the responsibility of the provincial authorities. The management structure and the implementation of TB control services vary considerably among provinces. The development of administrative districts, with health management structures in each province, is in progress and not yet complete. The basic unit for TB control and management is the individual primary care institution. Community health workers play an important role in patient care, but their involvement needs to be better organized and recorded.

The National Health Laboratory Service (NHLS) is the main provider of TB laboratory services in eight of the nine provinces in South Africa (all except KwaZulu-Natal) and is divided into central, coastal and northern regions. The laboratories of the NHLS are centralized, work under contract, and include primary health-care, regional, academic and referral laboratories. Communication between them is through a laboratory information system. Smear microscopy is performed in all laboratories; culture, identification and DST are performed in 11 referral laboratories throughout the country. In KwaZulu-Natal, 73 laboratories do smear microscopy, two have culture facilities and one referral laboratory carries out DST.

## **Surveillance and monitoring**

The incidence of TB in South Africa is uncertain because of weaknesses in the reporting system. Furthermore, the rise in TB incidence caused by the spread of HIV cannot easily be distinguished from improvement in case detection. It is likely, however, that the actual incidence of TB is higher than the current WHO estimate because case detection in 2003 was reported to be 118%.1

The treatment success rate in the 2002 cohort was 68% and has been consistently low since recording began in 1996. In 2002-2003, 22% of new smear-positive patients were lost to follow-up, either through default or transfer, and 9% died. A further 14% completed treatment but without evidence of smear conversion. The outcome among re-treatment cases was substantially worse, with a treatment success rate of 53% and with 34% lost to follow-up. As noted in the 2004

### **PROGRESS IN TB CONTROL IN SOUTH AFRICA**

#### Indicators

DOTS treatment success, 2002 cohort	68%
DOTS case detection rate, 2003	118% 1
NTP budget available, 2004	NA
Government contribution to NTP budget, including loans, 2004	NA
Government contribution to total TB control costs, including loans, 2004	NA
Government health spending used for TB control, 2004	7%

## **Major achievements**

- Implementation of the advocacy and social mobilization plan in five provinces (Eastern Cape, Western Cape, Gauteng, Limpopo and Free State)
- Implementation of a uniform, cohort-based reporting and recording system in all provinces
- Development of guidelines for care of HIV-infected TB patients, including access to ART

#### **Major planned activities**

- Strengthen DOT in the provinces and improve quality of data collected
- Implement and strengthen collaborative TB/HIV activities in subdistricts
- Shorten delays in diagnosis by sputum smear microscopy
- Improve laboratory infrastructure and coverage of services in remote areas

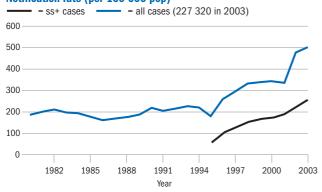
NA indicates not available.

Note that the "case detection rate" can exceed 100% because this is calculated as the ratio of cases reported in a given year to the estimated incidence in that year. Because the numerator is derived from the pool of prevalent cases. a proportion of which has arisen in previous years, the ratio can exceed 100%.

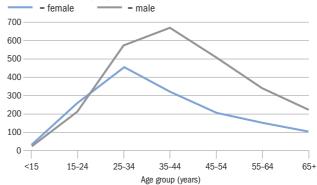
## **SOUTH AFRICA**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	45 026 470	DOTS coverage (%)	77	77	98	99.5
Global rank (by est. number of cases)	8	Notification rate (all cases/100 000 pop)	344	334	481	505
Incidence (all cases/100 000 pop/year)	536	Notification rate (new ss+/100 000 pop)	173	189	221	258
Incidence (new ss+/100 000 pop/year)	218	Detection of all cases (%)	74	68	94	94
Prevalence (all cases/100 000 pop)	458	Case detection rate (new ss+, %)	91	95	106	118
TB mortality (all cases/100 000 pop/year)	73	DOTS case detection rate (new ss+, %)	75	81	105	118
TB cases HIV+ (adults aged 15-49, %)	61	DOTS case detection rate (new ss+)/coverage (%)	97	105	107	119
New cases multidrug resistant (%)	1.6	DOTS treatment success (new ss+, %)	66	65	68	_

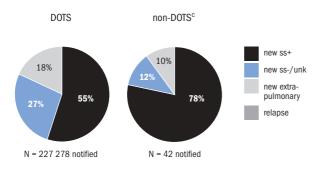
#### Notification rate (per 100 000 pop)



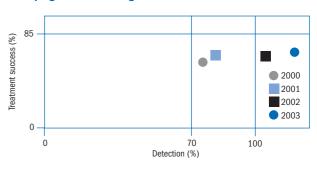
#### Notification rate by age and sex (new ss+)<sup>b</sup>



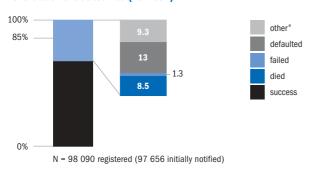
#### **Case types notified**



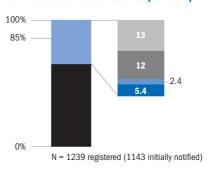
#### **DOTS** progress towards targets<sup>d</sup>



## **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)



ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

## **SOUTH AFRICA**

WHO report, it remains unclear why so many patients are lost to followup, and efforts need to be made to promote better adherence and to achieve better treatment outcomes.

Because the surveillance and monitoring data are still weak, and the electronic TB register was introduced only at the end of 2003, it is difficult to assess the TB burden and trends and to evaluate the impact of the DOTS programme. A national disease prevalence survey would help to determine how much TB there is in South Africa and would provide a baseline against which to measure the future impact of DOTS and related control methods for HIV and AIDS.

## **Improving programme** performance

The current TB control plan, the "Medium Term Development Plan" (2002-2005), was developed and endorsed by the national government and by eight of the country's nine provinces. A 10-year review of the programme is scheduled in 2005, and a new five-year plan will be developed in line with the strategies developed by the Department of Health.

South Africa has overcome some of the important constraints to achieving the global targets identified in the last report. A uniform, cohort-based recording and reporting system has been set up in all provinces, and the establishment of the electronic TB register will allow tracking of patients between health facilities. While staffing shortages still pose a problem at the provincial level, there has been an increase in staff at the national level. To address the lack of capacity, a training manual has been developed for medical practitioners and training workshops are being held in all provinces. The WHO training manual for trainers of facility health-care workers is being adapted and training of trainers will be conducted in all provinces. A database of trained staff has been established. A national TB manual is being developed.

A national advocacy and social mobilization plan entitled "Stop TB because you can" is being used to improve community awareness about TB through sustained and highly visible

campaigns. The plan has been used to advocate the need for more resources for TB control at all levels of the government and to bring together all partners involved in TB control. It is now implemented in five provinces (Eastern Cape, Western Cape, Gauteng, Limpopo and Free State).

New drug combinations, following the WHO-recommended treatment guidelines, were phased in during 2003 but this led to problems with drug supplies and to a shortage of first-line drugs. Furthermore, the sole supplier of streptomycin has stopped manufacturing the drug. While FDCs are now available in most districts, some districts have yet to train health staff in treatment regimens using FDCs.

Data collected from the most recent prevalence survey (2000–2002) estimated 7500 prevalent MDR-TB cases and about 450 new MDR-TB cases per year, corresponding to MDR-TB levels of 1.7% (new cases) and 6.6% (re-treatment cases). Treatment facilities for MDR-TB have been established in eight provinces. The Medical Research Council is currently developing a national policy on MDR-TB management. A standardized treatment regimen is provided to MDR-TB patients. The country is not planning to submit an application to the GLC as most second-line drugs are available in the country and many are locally produced.

#### Diagnostic and laboratory services

Nearly all laboratories participate in a quarterly EQA programme run by the NHLS, but the current programme does not yet completely satisfy international guidelines. The delays in sputum smear diagnosis are still too long and reporting mechanisms are inadequate in some laboratories. The NHLS plans to establish a national TB reference laboratory and to introduce a pilot EQA study for sputum smear microscopy that will comply with international guidelines. Other priorities for the NHLS are to improve the laboratory infrastructure and the coverage of services in remote rural areas, as well as training and monitoring.

#### TB/HIV coordination

South Africa had an estimated HIV prevalence of 22% among all adults at the end of 2003. A recent national survey estimated the HIV prevalence among TB patients to be 55% in 2002, close to the WHO estimate of 61% in 2003. There is a national TB/HIV coordinating body for collaborative activities, which have been implemented in 44 out of 174 subdistricts; it is planned to cover the entire country by 2007. TB/HIV provincial coordinators and national staff have been recruited and national guidelines for care of HIV-infected TB patients, including access to ART, have been developed. VCT is offered routinely to TB patients, but the acceptance rate remains low.

#### Links with other health-care providers and the community

A few public and private hospitals as well as prison health services implement DOTS. Several large private corporations, in particular in the mining industry, provide DOTS through their corporate health facilities and contribute about 20% of all reported cases. Several NGOs are involved in the delivery of TB control services and many have recruited community health workers and volunteers as DOTS providers. As noted in last year's report, a PPM-DOTS plan is still needed and more private sector participation should be encouraged.

#### **Partnerships**

South Africa has a country TB coordinating group that meets four times a year. Many partners and technical agencies support DOTS implementation and expansion, including CDC (surveillance and TB/HIV activities), DFID (district management and inpatient care of TB patients), IUATLD (laboratory support and programme management), KNCV (training and research) and WHO (training and TB/HIV activities). USAID is one of the main sources of funds and the GFATM has approved one grant to fund TB/HIV activities.

#### **Budgets and expenditures**

As in previous years, South Africa did not submit financial information to

## **SOUTH AFRICA**

WHO because the NTP does not have access to district and provincial financial data. South Africa was awarded one TB/HIV grant from the GFATM in round 2 for US\$ 8.4 million over two years; to date no funds have been disbursed. The Government of Belgium is also funding TB/HIV activities to the amount of US\$ 8.3 million over five years, of which US\$ 1.2 million has been disbursed. Estimates made in previous WHO reports suggest that the total annual cost of TB control in South Africa is about US\$ 300 million.

# **Thailand**

Thailand has had nationwide DOTS coverage since 2002 and reached the global target for case detection in 2003. Recent data suggest that the incidence of TB is declining slowly in Thailand. Considerable efforts are being made to extend TB control services to marginalized and deprived population groups, and this has boosted the case detection rate. However, treatment success is still well below the DOTS target and too many patients die, fail to complete their treatment or are lost to follow-up. In contrast to most HBCs, diagnostic laboratories in Thailand are relatively well equipped and maintained, but the shortage of adequately trained staff is still a problem. The estimated prevalence of HIV in Thailand is higher than in any other country in the WHO South-East Asia Region. A national TB/HIV coordinating body has been set up and is planning joint TB/HIV activities. The recent reform and decentralization of the country's health sector is changing the responsibilities and funding arrangements for TB control; the full implications of this for TB control are still unclear.

#### System of TB control

The central office of the NTP has become a cluster within the Bureau of AIDS, TB and STIs, following the recent reorganization of the Department of Disease Control (DDC) of the MoPH. The TB cluster is responsible for the development of technical policies, planning and monitoring of TB control in the country. The procurement and distribution of anti-TB drugs have been decentralized to the provincial and district levels as part of the healthsector reform process. Twelve regional TB centres and the TB cluster in Bangkok are responsible for monitoring, training and supervising of provincial and district-level staff. Health inspectors monitor the provincial hospitals and health offices and have a strong influence on provincial and district health-care programmes. Certain programmes are now given priority and

efforts have been made to include TB control among these priority programmes.

Under the health-sector reform project, a number of managerial tasks for the TB control programme, including planning and budgeting for activities such as training and supervision, have been decentralized to the provincial and district levels. District TB Coordinators (DTCs) are responsible for coordinating TB control activities, and work in close collaboration with the TB clinics in the hospitals. One effect of the health-sector reform policies will be to weaken the role of provincial and district health offices, as planning and budgeting authority will now rest with the provincial and district hospitals. In many districts, clinic staff in TB hospitals have assumed some of the responsibilities of the DTCs.

Laboratory diagnostic services in Thailand are provided by one NRL, 167 provincial and 678 district laboratories. All laboratories do smear microscopy, about 85 do mycobacterial culture and eight have facilities for DST. Regional and university laboratories perform culture on request for sputum smear-negative cases.

#### **Surveillance and monitoring**

Annual case notifications from 1980 to 1995 suggest that the underlying trend in incidence is downwards, masked since 1998 by improvements in case detection. Notification rates are highest in elderly men and women, which is consistent with a long-term downward trend in TB incidence. However, the recent impact of HIV on TB incidence cannot be determined from the nationally aggregated data. The prevalence of HIV among adult TB cases was estimated to be 8.7% in 2003, but HIV prevalence has been falling for several years, and TB incidence may also still be falling.

According to the most recent estimate, Thailand has exceeded the target for case detection, reaching 72% in 2003, following the rapid increase in DOTS population coverage between 1995 and 2002. In contrast, treatment success was well below target at 74% in the 2002 cohort, mainly

## **PROGRESS IN TB CONTROL IN THAILAND**

#### Indicators

DOTS treatment success, 2002 cohort	74%
DOTS case detection rate, 2003	72%
NTP budget available, 2004	100%
Government contribution to NTP budget, including loans, 2004	NA
Government contribution to total TB control costs, including loans, 2004	NA
Government health spending used for TB control, 2004	0.5%

#### **Major achievements**

- A meeting of TB coordinators from the regions, Bangkok and the prison service that addressed the referral and transfer system and overall strengthening of the TB network
- Recent DOTS expansion to marginalized population groups including people in border areas, migrants, prisoners and the urban poor leading to increased case

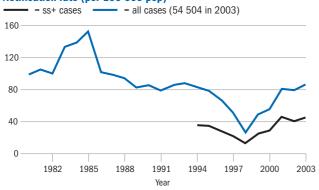
#### **Major planned activities**

- Implement TB/HIV collaborative activities including routine VCT for all TB patients, according to national guidelines
- Develop a comprehensive human resource plan for all levels of the NTP
- Build capacity for mycobacterial culture in provincial hospitals and strengthening the existing culture facilities in regional TB reference laboratories

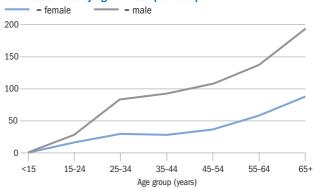
NA indicates not available.

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	62 833 330	DOTS coverage (%)	70	82	100	100
Global rank (by est. number of cases)	17	Notification rate (all cases/100 000 pop)	56	81	80	87
Incidence (all cases/100 000 pop/year)	142	Notification rate (new ss+/100 000 pop)	29	46	41	45
Incidence (new ss+/100 000 pop/year)	63	Detection of all cases (%)	39	57	56	61
Prevalence (all cases/100 000 pop)	208	Case detection rate (new ss+, %)	46	73	65	72
TB mortality (all cases/100 000 pop/year)	19	DOTS case detection rate (new ss+, %)	46	73	65	72
TB cases HIV+ (adults aged 15-49, %)	8.7	DOTS case detection rate (new ss+)/coverage (%)	66	89	65	72
New cases multidrug resistant (%)	0.9	DOTS treatment success (new ss+, %)	69	75	74	_

## Notification rate (per 100 000 pop)



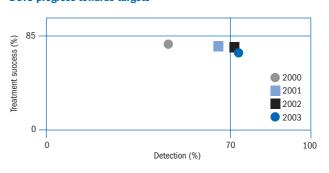
#### Notification rate by age and sex (new ss+)<sup>b</sup>



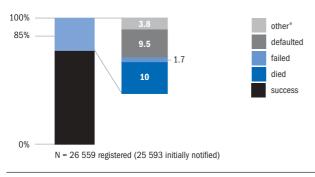
#### **Case types notified**



## **DOTS** progress towards targets<sup>d</sup>



## **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

because 11% of patients died while on treatment and 13% defaulted or were transferred between treatment centres without subsequent follow-up of treatment outcome. Among patients registered for re-treatment, the success rate was only 62%; 17% of patients died while on treatment. There has been no systematic improvement in treatment success in Thailand since data were first submitted to WHO in 1995. Given the rapid recent increase in DOTS coverage, and the apparently high rate of case detection, Thailand should consider assessing the quality of treatment observation and ensure that reporting and recording are accurate. A recent programme review showed that parts of the database were incomplete and inconsistent.

### Improving programme performance

In 2003, Thailand introduced a countrywide health insurance scheme for all clinical services known as the universal coverage (UC) scheme. The budget for this programme covers drugs and supplies for an "essential package of care" delivered at MoPH facilities and other health-care facilities under contract with the MoPH. Since 2002, following health-sector reforms, anti-TB drugs have been financed through the UC scheme. It is intended that training, supervision and monitoring activities for specific disease control programmes be financed through a non-UC budget available through the DDC at the MoPH. The TB cluster at the central level must use non-UC funds for the organization of national training courses, supervision in the regions and the organization of monitoring meetings for regional staff. During 2004, many training and monitoring activities required by NTP policy could not be carried out because of lack of funding at the peripheral level. Although funds are available for supervisory activities in 2004, the funding of training activities and monitoring meetings will require further negotiation with the DDC. The administrative process for the provision of non-UC funds at provincial and district levels is still being developed.

During the expansion of DOTS activities from 1996-2001, initial training was carried out for health-care workers at all levels. However, because of the high turnover of staff and the lack of systematic refresher courses, there is now a shortage of adequately trained TB control staff in Thailand. As noted last year, many of the regional TB offices have been weakened because staff posts have been cut and additional duties have been assigned to existing staff. A comprehensive HR development plan has been prepared; the immediate challenge is to ensure that sufficient funding is available to implement it.

Referral and transfer systems between treatment units and between prisons and MoPH facilities are weak. Timely information is often not communicated and there is no specific budget for communication between provinces. A TB network meeting was held in 2003 for the TB coordinators of the 12 regions, Bangkok and the prison service to address the referral and transfer system and overall strengthening of the TB network. Efforts are being made to improve data collection, and there are plans to introduce an electronic data management system.

The recent DOTS expansion to marginalized population groups including people living in border areas, migrants, prisoners and the urban poor have contributed to the high case detection rate in Thailand. NGOs and other organizations outside the MoPH system have been particularly active in expanding DOTS services to these groups.

Funding for anti-TB drugs has been adequate in the past but may be threatened if the purchase of drugs must be financed from fixed province and district budgets. Currently, most anti-TB drugs used by the NTP are manufactured in Thailand and are more costly than internationally procured drugs. Renegotiating prices with the government pharmaceutical organization or exploring additional procurement channels may help to release local funds for other TB-related activities such as training and supervision. The prevalence of MDR-TB among new cases decreased from 2.1% in 1997 to 0.9% in 2001. A nationwide drug resistance survey is

planned for 2005. At present, the NTP does not diagnose and treat MDR-TB patients. However, policy guidelines on MDR-TB management are being developed.

Three areas where programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

Compared with most other HBCs, Thailand has relatively well-equipped laboratories with few supply or maintenance problems. Thailand is planning to broaden the range of diagnostic services for TB by developing further capacity for doing TB culture in provincial hospitals and by strengthening the existing culture facilities of regional TB reference laboratories. The rapid detection of drug resistance is a priority for the NRL. EQA activities cover all TB laboratories in MoPH facilities, and efforts are to being made to include the private sector in the quality assurance scheme. Laboratory training activities are being expanded to include training for all TB control staff and targeted training for laboratory staff in technical areas where laboratory performance needs to be improved.

#### TB/HIV coordination

The estimated prevalence of HIV in Thailand (1.5% of adults aged 15-49 at the end of 2003) is the highest in the WHO South-East Asia Region. The prevalence of HIV among TB patients in sentinel surveys was between 10% and 15% in the country as a whole, but up to 30% in some regions (higher than the WHO estimate of 9% of adult TB patients). A national TB/HIV coordinating body was first established in 2001 and a national TB/HIV strategy has been in place since 2004. National TB/HIV guidelines were prepared in 2004 and will be implemented in January 2005.

VCT is offered to all TB patients in four pilot provinces: Chiang Rai, Ubon Ratchathani, Phuket and two districts in Bangkok. Data on specific indicators such as the proportion offered counselling, the proportion tested and the proportion found to be HIV posi-

## **THAILAND**

tive will be collected and analysed. It is planned to train all TB clinic staff in the country in VCT by 2005.

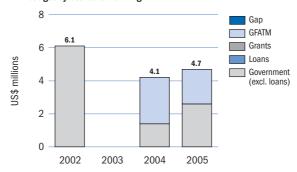
#### Links with other health-care providers

The NTP has established a task force for PPM DOTS and has begun to collaborate with some private hospitals, private physicians and NGOs. There is a need to strengthen ties with all public sector providers and institutions involved in TB treatment and diagnosis, especially since the healthsector reform process has led to a more diversified network for delivering TB care. Public hospitals throughout the country are involved, but the participation of medical colleges and of prison and military health services remains limited.

#### **Partnerships**

CDC, RIT and WHO are the main technical partners in Thailand, assisting with DOTS expansion and TB/HIV activities. CDC (USAID) and GFATM are the major funding partners for surveillance, laboratory services, training and collaborative TB/HIV activities.

#### NTP budget by source of funding



#### **Budgets and expenditures**

Comprehensive data on NTP budgets and expenditures are not available for the period 2003-2005. This is because, under the new health insurance scheme introduced in 2003, provincial and district hospitals receive budgets (calculated on the basis of fixed per capita rates) to provide a package of clinical care. It is not clear how much funding for the TB control programme is provided from these budgets. Meanwhile, programme support functions such as training and supervision are covered through a separate budget. Budget figures reported to WHO for 2004 and 2005 therefore reflect only

the budget managed by the TB cluster in Bangkok. As a result, the reported budget has fallen from US\$ 6.1 million in 2002 to US\$ 4.7 million in 2005, despite an increase in funding from the GFATM. The development of national budgets in future will depend on the NTP's ability to implement a comprehensive financial monitoring system that allows budgets and available funding to be reported by all provinces and districts. Estimates made in previous WHO reports indicate that the total cost of TB control is about US\$ 10 million per year, and around US\$ 170 per patient treated.

# Uganda

Uganda is pursuing universal access to DOTS as its TB control strategy. All districts were implementing DOTS as far back as 1997, but the low coverage of general health services means that more than 50% of the population are still without access to TB services, although this proportion is decreasing. TB is a component of Uganda's minimum health-care package, and TB services are integrated in primary health care. However, many health facilities do not yet provide TB diagnosis and care. In 2000, Uganda adopted the community-based TB care model as the best strategy to control TB in the country. Community involvement is recognized as crucial for the success of TB control in Uganda. The NTP is committed to expanding community-based DOTS (CB-DOTS) to all 56 districts; 51 districts have already been covered. Although GFATM funds were approved for Uganda in round 2, a substantial funding gap still remains, meaning that some of the planned activities for 2005 may not be carried out. In December 2004, the Uganda Stop TB Partnership (USTP) was

launched as a major initiative to raise awareness of TB as an important public health problem and to mobilize additional resources for TB control.

#### **System of TB control**

Uganda has a decentralized system of governance, with central ministries responsible for policies, standards, quality control, resource mobilization and training. Districts are responsible for management of services at peripheral level. Health service delivery, including TB control, is the responsibility of the health subdistricts (HSDs), which are the functional units for TB control. A total of 214 HSDs each serve about 100 000 people. TB service delivery is fully integrated in the primary health-care system.

TB control in Uganda is organized with a central unit at the MoH run by the NTP manager and one administrative officer. Hitherto, the NTP manager has been assisted by six zonal tuberculosis and leprosy supervisors (ZTLSs) based at the periphery, who oversee TB control in their zones. At district level, a district health team (DHT) oversees TB control. The District Tuberculosis and Leprosy Supervisor, a member of the DHT, is responsible for TB control including data collection, analysis and reporting. Below this level, general health workers handle TB control activities as part of their general duties.

The MoH has recently developed the second five-year "Health Sector Strategic Plan" (HSSP II) covering 2005/2006-2009/2010. HSSP II envisages continued implementation of the minimum health-care package of which TB is one component and foresees continued use of TB performance indicators for monitoring progress of HSSP II implementation. As part of continuing health sector reform, the NTP plans to recall the ZTLSs to the centre in order to form, under the guidance of the NTP manager, a strong central team with improved capacity in policy formulation and technical guidance to districts and partners on TB management. The ZTLSs will provide a strong technical link between government and partners, and support to the DHTs. Each district has three or more health facilities providing TB diagnostic and treatment services. Through CB-DOTS, treatment is provided at the community level. CB-DOTS is an important service delivery mechanism that is patient-centred and based on participation by civil society, providing accessible, cost-effective TB care; this mechanism is vital to the success of TB control in Uganda.

The NRL in Kampala is responsible for training, DST and EQA. The national coordinator of the laboratory network is responsible for the NRL. Ten regional laboratories based at regional hospitals also provide training and EQA, in addition to smear microscopy. The district laboratories' main responsibility is the supervision of peripheral laboratories, which serve as the main diagnostic units.

## **PROGRESS IN TB CONTROL IN UGANDA**

## **Indicators**

DOTS treatment success, 2002 cohort	60%	
DOTS detection rate, 2003	44%	
NTP budget available, 2004	83%	
Government contribution to NTP budget, including loans, 2004		
Government contribution to total TB control costs, including loans, 2004	38%	
Government health spending used for TB control, 2004		

#### **Major achievements**

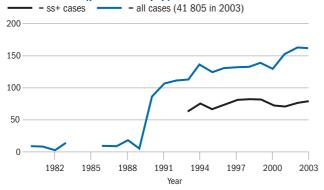
- Expansion of community-based DOTS (CB DOTS) to an additional 11 districts, corresponding to an additional 20% of the country's population
- Formation and launch of the Uganda Stop TB Partnership (USTP) to better harness efforts of all partners on TB control
- Secured additional staff to build NTP capacity; secured additional resources through ISAC

#### **Major planned activities**

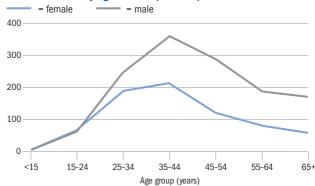
- Complete expansion and consolidation of CB DOTS, ensuring district-wide coverage and high quality of services
- Institute EQA of all laboratories in the country, and strengthen it where it exists
- Update NTP strategic TB control plan to include PPM DOTS as part of the DOTS expansion plan
- Operationalize the USTP

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	25 826 968	DOTS coverage (%)	100	100	100	100
Global rank (by est. number of cases)	16	Notification rate (all cases/100 000 pop)	129	152	163	162
Incidence (all cases/100 000 pop/year)	411	Notification rate (new ss+/100 000 pop)	73	71	76	79
Incidence (new ss+/100 000 pop/year)	179	Detection of all cases (%)	38	42	42	39
Prevalence (all cases/100 000 pop)	652	Case detection rate (new ss+, %)	50	46	46	44
TB mortality (all cases/100 000 pop/year)	96	DOTS case detection rate (new ss+, %)	50	46	46	44
TB cases HIV+ (adults aged 15-49, %)	21	DOTS case detection rate (new ss+)/coverage (%)	50	46	46	44
New cases multidrug resistant (%)	0.5	DOTS treatment success (new ss+, %)	63	56	60	_

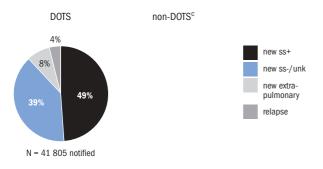
#### Notification rate (per 100 000 pop)



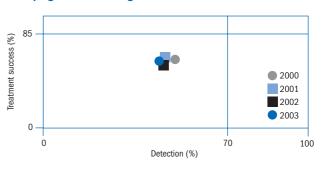
#### Notification rate by age and sex (new ss+)<sup>b</sup>



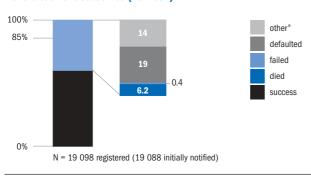
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

#### **Surveillance and monitoring**

While strong evidence exists that HIV prevalence has been falling in Uganda since the early 1990s, the notification rate of TB cases (all forms) has increased since 2000. The notification rate of new smear-positive cases has remained steady for the past decade. In order to estimate the case detection rate, it has been assumed that the increase in the notification rates of all forms of TB reflects an increase in incidence - and that the case detection rate of new smear-positive cases has fallen since 2000 from 50% to 44%. A plausible alternative explanation is that the case detection rate has improved as the coverage of CB DOTS has been extended throughout the country - and that TB incidence has stabilized or begun to fall. However, given the low access to health services (less than 50%), it is unlikely that the case detection rate is much higher than estimated here.

Notwithstanding the uncertainty surrounding the assessment of the case detection rate, it is clear that the NTP must work to improve treatment outcomes, which have been consistently low. Only 60% of new smearpositive patients were successfully treated in the 2002 cohort; 33% defaulted, were transferred without follow-up or were not evaluated. Cure was bacteriologically confirmed in only half the patients successfully treated; the final smear examination was not done for the other patients. The pattern is similar among patients registered for re-treatment.

A small disease prevalence survey was carried out in Kampala in 2001-2002,1 finding a prevalence of smearpositive TB of 440 cases per 100 000 population in the periurban community sampled. However, a larger national survey is needed to assess the total burden of TB in Uganda and to set a baseline against which to measure the impact of DOTS. Alternatively, or in addition, a systematic evaluation of the process of diagnosis and reporting in Uganda would allow a reassess-

ment of the case detection rate. The NTP acknowledges that progress has been hindered by non-prioritization of sputum smear examination (15% of new cases were put on treatment without sputum smear results), poor recording and the absence of strategies to recover interrupters and to capture the true treatment outcome of patients who transfer between treatment units. These factors are being addressed systematically.

### Improving programme performance

Since early 2004, the NTP has benefited from the ISAC initiative, which greatly contributed to the increased capacity of the central team. An international WHO staff member supports the NTP central unit. The central unit has deployed three recently recruited professional officers at regional level to support the DHTs, with redistribution of regional supervisors to weak areas. However, major HR deficiencies still exist at the central level. CB DOTS is being expanded and, by the end of 2004, was being implemented in 51 out of 56 districts. The full impact of this expansion has not yet been seen.

CB DOTS is being implemented in a phased manner in the HSDs of some districts. Uganda has so far engaged two international NGOs (International Medical Corps and the Malaria Consortium) to implement CB DOTS in remote areas; the NTP plans to expand CB DOTS to the remaining districts early in 2005.

Supervision and monitoring activities have been expanded to all levels of TB control, including the community. A new DOTS expansion plan for the next five years is being developed by the NTP.

In 1996-1997, a DRS was conducted in areas of the country supported by GLRA, giving an estimate of 0.5% MDR in new pulmonary cases and 4.4% MDR in re-treatment cases. Resources are being sought to carry out a new DRS. Uganda plans to apply to the GLC in the context of the new DOTS expansion plan.

Three other areas in which programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers and the community.

#### Diagnostic and laboratory services

The two main challenges facing the diagnostic and laboratory services are the shortage of qualified laboratory personnel in the general health service and the lack of a countrywide EQA system for sputum smear microscopy. In 2003, only 12 out of 56 districts had implemented EQA for smear microscopy. In 2005, the NTP plans to establish routine EQA in the remaining districts and to strengthen it in those districts where it exists. The NTP will advocate for recruitment of qualified personnel in peripheral laboratories and will train existing personnel as microscopists in the interim.

#### TB/HIV coordination

An interim national TB/HIV coordinating body comprising the managers of the NTP and of the National AIDS Control Programme and partners (including the AIDS Information Centre, the AIDS Support Organization, GLRA, USAID-funded organizations and WHO) was formed in 2004 to formulate a policy as well as to prepare a proposal for collaborative TB/HIV activities. The committee will ensure phased implementation of collaborative TB/HIV activities in pilot districts and, based on the experiences gained, will frame the policy and strategy for rapid nationwide expansion.

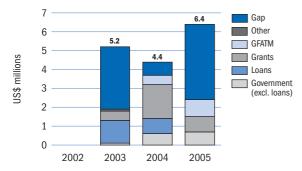
In 2004, WHO appointed an NPO to coordinate TB/HIV activities and oversee the establishment of the committee; its first meeting was planned for mid-January 2005.

#### Links with other health-care providers

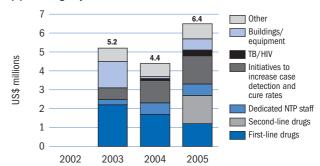
A situation analysis has shown that many patients in urban areas are treated in the private sector. The NTP has initiated a small-scale collaborative project with private hospitals, with plans to expand this initiative to involve individual private medical practitioners. NGOs play an important role in DOTS implementation, and the NTP has involved many general hospitals, a few medical colleges, and prison, army and police health facilities in TB control.

<sup>&</sup>lt;sup>1</sup> Guwatudde D et al. Burden of tuberculosis in Kampala, Uganda, Bulletin of the World Health Organization, 2003, 81:799-805.

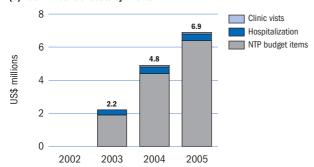
#### (a) NTP budget by source of funding



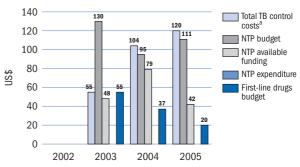
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>







a Total TB control costs for 2003 are based on available funding, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

#### Links with the community

The success and sustainability of CB DOTS is largely dependent on community involvement and ownership of the programme. In the CB-DOTS strategy, communities participate in selecting lay community members to support patients and ensure treatment compliance. The selected community members work on a voluntary basis, observing and recording the ingestion of each day's medication. In addition, they encourage patients to go for follow-up sputum smears and suspects to go for examination in health units. Community volunteers are responsible to the community and also to the formal health system through a public health worker who supervises and replenishes drug supplies.

#### **Partnerships**

The MoH is committed to attaining the targets for DOTS implementation, and has indicated an interest in forming a strong partnership to help the country accelerate towards the 2005 targets. The formation of the USTP was spearheaded by the MoH and supported by WHO and the Global Stop TB Partnership, as well as other part-

ners, in the context of the ISAC initiative. The USTP was launched in December 2004, with the aims of harnessing the contributions of all stakeholders to TB control and raising the profile of TB as a major public health problem. WHO Uganda has offered to host the USTP Secretariat; a Memorandum of Understanding is being prepared to guide their operations.

#### **Budgets and expenditures**

The NTP budget has been between US\$ 4.4 million and US\$ 6.4 million during the period 2003-2005 (equivalent to US\$ 95-130 per patient). Despite the approval of a GFATM grant in round 2, Uganda suffers from a persistent funding gap, which is expected to reach US\$ 4 million in 2005, representing 62% of the NTP budget.

The budget for first-line anti-TB drugs has decreased from US\$ 2.2 million in 2003 to US\$ 1.2 million in 2005, whereas the expected number of patients to be treated is rising. The drug budget per patient treated has thus been reduced from US\$ 55 to US\$ 20. This budget for first-line drugs is fully funded. In contrast, while the

budgets for initiatives to increase case detection and cure rates and for collaborative TB/HIV activities have been increasing between 2003 and 2005, implementation of all of the planned activities will depend on the availability of additional funds. A need for second-line drugs has also been identified; a budget of US\$ 1.5 million has been included in 2005, which is for a stock sufficient to treat 1000 MDR-TB patients. However, funding has not yet been secured.

The total cost of TB control, including the costs of clinic visits and hospital stays as well as the NTP budget, will increase from an estimated US\$ 2.2 million in 2003 to US\$ 6.9 million in 2005 (US\$ 55-120 per patient), provided the existing budget gap for 2005 is filled. If no additional funds are secured, total costs will reach only about US\$ 3 million in 2005; the cost per patient will be US\$ 51. The costs of clinic visits, at around US\$ 0.1 million, is relatively low given the small number of visits required to health facilities following the nationwide introduction of CB DOTS

# United Republic of Tanzania

The United Republic of Tanzania was among the first countries to adopt the DOTS strategy. Nationwide DOTS coverage was attained in 2002, largely through the successful integration of TB control in the general health services. After reaching a peak in 2001, the number of reported TB cases has remained steady, which may perhaps indicate an end to the rise in TB incidence previously associated with the HIV epidemic. As the HIV prevalence has been constant in the country since 1996, the DOTS programme should be able to achieve a progressive reduction in TB incidence from now on. Improvements have been made in the treatment of patients, but a relatively high death rate is still an obstacle to reaching the global target for successful treatment. While progress has been made in control of both TB and HIV and in ART scale-up, the TB and HIV control programmes have not worked together in the past, and the particular needs arising from the epidemic of TB/HIV coinfection have not received special attention until recently. The government has now developed comprehensive plans for collaborative TB/HIV activities and, thanks to an award from the GFATM.

it should be possible to implement them all. Building on the well-managed TB control programme, the collaborative TB/HIV activities will give additional impetus to TB case-finding and treatment. The MoH is also preparing to establish DOTS-Plus within the regular DOTS programme. Further strengthening of human resources, particularly at central level, is essential to meet the needs of these rapidly expanding programme activities.

#### **System of TB control**

The NTP is well organized and managed. Under the direction of a small central unit, the regional and district TB coordinators supervise the activities of hospitals and other health centres and monitor programme performance, using formal quality assurance practices. The district health committees are responsible for developing district health plans that include both TB and HIV. Recognizing the importance of the dual epidemic of TB and HIV, the NTP has decided to implement the full package of collaborative TB/HIV activities as part of a comprehensive TB and HIV/AIDS control strategy.

The NRL oversees 2 zonal, 18 re-

gional and 701 district laboratories. Culture is done at the NRL and zonal laboratories, while DST is carried out only at the NRL.

#### **Surveillance and monitoring**

The total annual TB notification rate has increased three-fold between 1980 and 2001, and has fallen slightly since then. The notification rate of smear-positive cases has fallen slightly since 1998. Assuming that this is a consequence of the earlier levelling off of the HIV epidemic rather than a decline in case detection rates, the DOTS programme should now begin to reduce the incidence of TB, provided the programme performance is maintained or improved. The estimated rate of smear-positive case detection in 2003 (43%) was low, but the reliability of this estimate is not easily verified using the available tuberculin testing data because the usual methods of analysis based on the Stýblo ratio may not apply when the prevalence of HIV is high (see Methods). For this reason, a systematic and quantitative assessment of the completeness of surveillance data or a survey of the prevalence of disease would be very informative. Given the high rate of HIV infection in the country, the treatment success rates are good: 80% for new cases, 79% for relapse cases, 65% for re-treatment after failure and 71% for re-treatment after default. Treatment outcomes for new smear-positive patients have improved steadily since 1995, but the high death rates (11% for the 2002 cohort) are the main obstacle to reaching the 85% target for treatment suc-

#### Improving programme performance

To improve case detection, the number of diagnostic centres has been increased in the districts, and a start has been made on integrating the delivery of TB control into the general health services and into the private sector. In 2003, 1250 general health-

#### PROGRESS IN TB CONTROL IN THE UNITED REPUBLIC OF TANZANIA

#### **Indicators**

DOTS treatment success, 2002 cohort	80%
DOTS case detection rate, 2003	43%
NTP budget available, 2004	76%
Government contribution to NTP budget, including loans, 2004	14%
Government contribution to total TB control costs, including loans, 2004	64%
Government health spending used for TB control, 2004	11%

#### **Major achievements**

- Increased number of diagnostic centres at district level
- Training of 1250 general health-care workers in case detection and treatment
- Maintained high treatment success despite high prevalence of HIV among TB
- Strengthening of MDR-TB services and infrastructure in preparation for application to the GLC

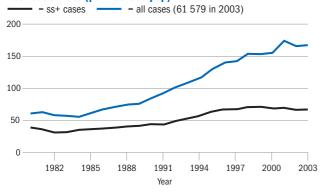
#### **Major planned activities**

- Expand DOTS by involving communities, the private sector, specialist TB clinics, medical colleges and prison health services
- Introduce DOTS-Plus activities
- Expand collaborative TB/HIV activities to all districts by 2007

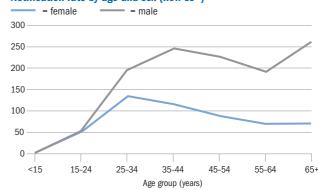
## **UNITED REPUBLIC OF TANZANIA**

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	36 976 622	DOTS coverage (%)	100	100	100	100
Global rank (by est. number of cases)	14	Notification rate (all cases/100 000 pop)	156	173	166	167
Incidence (all cases/100 000 pop/year)	371	Notification rate (new ss+/100 000 pop)	69	69	67	67
Incidence (new ss+/100 000 pop/year)	157	Detection of all cases (%)	45	49	46	45
Prevalence (all cases/100 000 pop)	524	Case detection rate (new ss+, %)	47	46	43	43
TB mortality (all cases/100 000 pop/year)	86	DOTS case detection rate (new ss+, %)	47	46	43	43
TB cases HIV+ (adults aged 15-49, %)	36	DOTS case detection rate (new ss+)/coverage (%)	47	46	43	43
New cases multidrug resistant (%)	1.2	DOTS treatment success (new ss+, %)	78	81	80	_

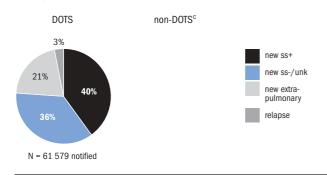
#### Notification rate (per 100 000 pop)



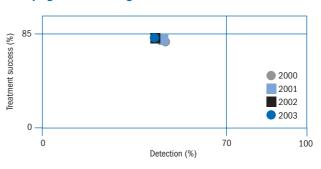
#### Notification rate by age and sex (new ss+)<sup>b</sup>



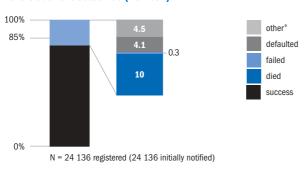
#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

Absence of a graph indicates that the data were not available or applicable.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

## UNITED REPUBLIC OF TANZANIA

care workers were trained on case detection and treatment. The central unit has produced training guidelines and a facilitator module for clinical officers and will develop training guidelines for nurses.

To improve programme performance, it will be necessary to strengthen HR capacity at the central level. Currently, there are only four people to supervise and monitor the TB control programme, to expand DOTS services, to implement training at lower levels and to develop and implement collaborative TB/HIV activities. PATH, with funding from USAID, has recently developed a plan to strengthen HR capacity at central, regional and district level.

The MoH is planning to establish and integrate a DOTS-Plus component within the NTP. Following a WHO mission in spring 2004, plans have been made to: introduce DOTS-Plus, including developing a computerized TB notification system to monitor treatment outcomes among re-treatment cases; construct a new MDR-TB ward within the national TB hospital; set up a technical committee to oversee future implementation of MDR-TB activities; and provide training for medical personnel in management of MDR-TB. A drug resistance survey is scheduled to start in the middle of 2005.

Three areas where programme performance needs to be improved are diagnostic and laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

The quality of the central laboratory services has been significantly improved in preparation for an application to the GLC. New equipment has been installed, and internal quality control is now mandatory; the mycobacterial culture contamination rate has been reduced from 15% to 10% in less than a year. However, laboratories at all levels are still short of qualified staff and the implementation of EQA for smear microscopy is still not

satisfactory. A further priority for the laboratory network is to improve the quality of supervision of the peripheral laboratories by the central unit.

#### TB/HIV coordination

A national TB/HIV strategic plan to cover all districts by 2007 has been developed and includes all the collaborative TB/HIV activities defined in the WHO interim policy. In 2003, the Tanzanian Government successfully applied to the GFATM (round 3) for resources to support collaborative TB/ HIV activities in 45 of 120 districts. There is a gap in funding to scale up TB/HIV activities nationally, and there is a need to align TB/HIV activities with the national plan for scaling up access to ART to ensure that HIVpositive TB patients are able to access ART. Implementation of collaborative TB/HIV activities is slow; in order to accelerate their implementation, additional financial resources will be needed as well as increased HR capacity, particularly at central level.

#### Links with other health-care providers

Anti-TB drugs may only be prescribed and dispensed with the approval of the NTP and using drugs procured and distributed by the NTP. As a result, non-DOTS treatment of TB is very limited in both the private and the public sector, which facilitates the implementation of PPM-DOTS strategies. The NTP has involved NGOs and private hospitals in TB control by providing training, drugs and supervision, and is now expanding this effort to include private clinics. Links with specialist TB clinics, medical colleges and prison health services are also being strengthened.

#### **Partnerships**

A range of technical and financial partners are involved in TB control and they have formed an Interagency Coordination Committee that meets once a year. Development Cooperation Ireland, the Government of the Netherlands and the Swiss Agency for Development and Cooperation are the main sources of funds for TB control

activities. GLRA, KNCV and WHO all support programme monitoring and offer other technical assistance.

#### **Budgets and expenditures**

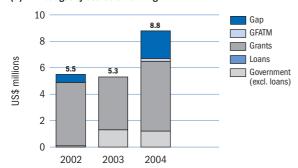
The NTP budget has increased from about US\$ 5 million in 2002-2003 to nearly US\$ 9 million in 2004 (from about US\$ 90 per patient in 2002-2003 to US\$ 133 in 2004). Budget data are not yet available for 2005 since the fiscal year starts in July. The budget was increased in 2004 to pay for dedicated staff, for the implementation of collaborative TB/HIV activities and for investment in buildings and equipment. The available funding has also increased, from around US\$ 5 million in 2002 and 2003 to US\$ 6.7 million in 2004. Most NTP funding comes from grants, with the government contributing US\$ 1.2 million (about 10% of the budget) in 2004. While a grant from the GFATM should make it possible to carry out the planned collaborative TB/HIV activities in selected pilot districts (provided that sufficient staff are available), a funding gap of US\$ 0.8 million remains.

In 2003, the government contribution was only US\$ 0.6 million rather than US\$ 1.3 million as anticipated. However, total available funding was higher than expected, at US\$ 5.6 million. Expenditures in 2003 were US\$ 3.8 million, i.e. 62% of the funds received. As more funding becomes available through grants, the capacity of the programme to absorb this money may become an important is-

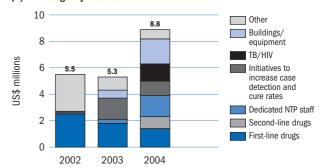
The total cost of TB control, which includes the cost of dedicated TB beds, clinic visits during treatment and items included in the NTP budget, was between US\$ 15 million and US\$ 16 million in 2002 and 2003 (about US\$ 250-275 per patient treated). If the 2004 budget is fully funded and the money is spent, this could increase to US\$ 21 million in 2004 (US\$ 320 per patient treated).

## UNITED REPUBLIC OF TANZANIA

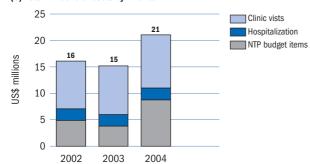
#### (a) NTP budget by source of funding



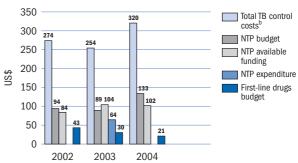
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a,b</sup>



#### (d) Per patient costs, budgets, available funding and expenditures<sup>a</sup>



<sup>&</sup>lt;sup>a</sup> No data available for 2005 - see text for explanation.

b Total TB control costs for 2002 are based on available funding, whereas those for 2003 are based on expenditures, and those for 2004 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

# **Viet Nam**

Viet Nam is the only member of the current group of HBCs1 to have reached the targets for DOTS implementation, which were achieved before 2000 and exceeded subsequently. This outstanding success was made possible by the effective integration of political commitment, international technical assistance and funding, and efficient community mobilization. Viet Nam has continued to expand the programme so as to reach remote population groups who have not had access to TB services, and to strengthen the diagnostic laboratory network. An urgent priority is the development of a national plan for improved TB/HIV coordination. A planned national TB prevalence survey will be of critical importance for measuring the impact of DOTS on the TB epidemic. Because of its success in achieving the targets, Viet Nam does not need substantial budget increases in 2005.

#### **System of TB control**

The National Hospital of Tuberculosis and Respiratory Diseases, in Hanoi,

is responsible for the activities for all of Viet Nam. Pham Ngoc Thach Hospital in Ho Chi Minh City is appointed to supervise the activities for the southern provinces. Each province has a provincial TB centre, under the direction of the provincial health service, which is responsible for the local implementation of the TB control programme. The district TB units, directed by the district health centres, coordinate the operation of peripheral TB activities. TB patients are referred to the district health centres from community health posts for sputum examination and initial treatment.

An effective national TB laboratory network operates under the supervision of the NTP. There are two reference laboratories (Hanoi and Ho Chi Minh City) that perform culture and drug susceptibility testing. Of the 64 provincial TB laboratories, nearly one quarter perform culture. Smear microscopy services are provided by more than 600 district TB laboratories.

#### Surveillance and monitoring

The best estimates of case detection

for 2003 (86%) and treatment success for the 2002 cohort (92%) suggest, as in previous years, that Viet Nam has comfortably exceeded the targets for DOTS implementation. Given that DOTS coverage and case detection and cure rates have been very high since 1997, a fall in the incidence rate could be expected, which should be reflected in the trend in case notifications. It is unclear why no such decline is visible in the nationally aggregated data, but analysis by province could be more illuminating. Case-notification rates are highest among elderly men and women, suggesting that TB incidence has been higher in the past. It is possible that incidence is not falling perceptibly in Viet Nam because the case detection rate may be lower, and the incidence rate higher, than the WHO estimates. In this context, Viet Nam's longplanned prevalence survey would help to establish the true burden of TB in the country, as well as providing a baseline against which to evaluate the impact of the programme on the TB epidemic.

## **Improving programme**

performance

Although all provinces maintain 100% coverage by the DOTS strategy, there are populations living in remote and mountainous areas with limited access to DOTS services. The NTP is expanding DOTS to reach these areas while maintaining excellent services. Efforts to reach these remote populations and other vulnerable groups started in 2003 and continued in 2004. Maintaining a consistent supply of high-quality anti-TB drugs for the entire country, especially in newly covered areas, is another important challenge being addressed by the NTP. A regulatory framework and enforcement mechanism have been developed to ensure the high quality of

#### PROGRESS IN TB CONTROL IN VIET NAM

## **Indicators**

DOTS treatment success, 2002 cohort 92% DOTS case detection rate, 2003 86% NTP budget available, 2004 98% Government contribution to NTP budget, including loans, 2004 78% Government contribution to total TB control costs, including loans, 2004 90% Government health spending used for TB control, 2004 6%

#### **Major achievements**

- Expansion of the TB network to cover remote and mountainous areas and increased access to DOTS for vulnerable groups
- Establishment of nationwide EQA system for smear microscopy
- Development of a regulatory framework and enforcement mechanism to ensure the high quality of anti-TB drugs
- Pilot testing of isoniazid preventive therapy for PLWHA infected with M. tuberculosis and co-trimoxazole preventive therapy for TB patients coinfected with HIV in An Giang province
- Studies on FDCs for patients in remote areas and on PPM-DOTS

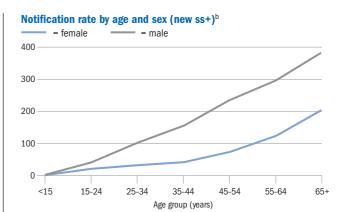
#### **Major planned activities**

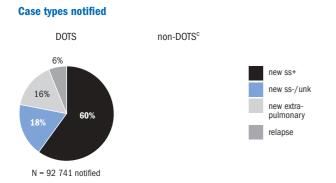
- Develop five-year plan for NTP for 2006–2010
- Train staff in EQA and maintain system throughout the country
- Carry out third national drug resistance survey

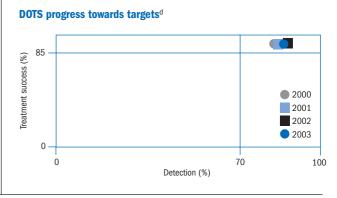
<sup>&</sup>lt;sup>1</sup> Peru was excluded from the original group of HBCs, having met the targets and successfully reduced incidence.

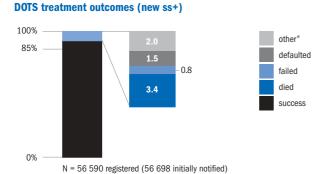
LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	81 376 724	DOTS coverage (%)	99.8	99.8	100	100
Global rank (by est. number of cases)	13	Notification rate (all cases/100 000 pop)	115	115	118	114
Incidence (all cases/100 000 pop/year)	178	Notification rate (new ss+/100 000 pop)	68	68	71	69
Incidence (new ss+/100 000 pop/year)	80	Detection of all cases (%)	63	63	66	64
Prevalence (all cases/100 000 pop)	240	Case detection rate (new ss+, %)	83	84	88	86
TB mortality (all cases/100 000 pop/year)	23	DOTS case detection rate (new ss+, %)	83	84	88	86
TB cases HIV+ (adults aged 15-49, %)	2.8	DOTS case detection rate (new ss+)/coverage (%)	83	84	88	86
New cases multidrug resistant (%)	2.3	DOTS treatment success (new ss+, %)	92	93	92	_

#### Notification rate (per 100 000 pop) = ss+ cases = all cases (92 741 in 2003) 120 40 0 1982 1985 1988 1991 1994 1997 2000 2003 Year









#### Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

Absence of a graph indicates that the data were not available or applicable.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

anti-TB drugs for TB services both within and outside the NTP. The feasibility of using FDCs for patients living in areas which are difficult to access is being explored. The last drug resistance survey was carried out in 1996 and estimated the prevalence of MDR-TB at 2.3% among new cases. A new survey is scheduled for 2005.

The NTP is developing the next fiveyear plan for TB control (2006-2010). Human resource capacity development will continue to be a priority, and the NTP will work with local authorities to recruit and maintain existing staff and to develop intensified training activities for staff at all levels.

Three other areas where programme performance needs to be improved are: diagnostic and laboratory services, TB/HIV coordination, and links with other health-care providers and the community.

#### Diagnostic and laboratory services

As DOTS services are expanded to remote and mountainous regions, diagnostic services also need to be provided to these areas. An EQA system for sputum microscopy based on new international guidelines is being

established in laboratories at district level throughout Viet Nam. In 2004 and 2005, staff in 20 of 64 provinces will be trained on the EQA system, and methods will be developed to implement and maintain EQA throughout the country.

#### TB/HIV coordination

In 2002, the prevalence of HIV in new TB patients was estimated to be 3% based on HIV sentinel surveillance among TB patients. This is somewhat higher than the WHO national estimate of 1.8%. In 10 provinces HIV prevalence exceeded 3%, and in two provinces (Binh Duong and Haiphong) the prevalence was more than 10%. In An Giang Province a pilot project included the use of isoniazid preventive therapy for PLWHA infected with M. tuberculosis, and co-trimoxazole preventive therapy for TB patients with HIV coinfection. ART for HIV-infected TB patients is not yet available. There is an urgent need for a well-defined national plan for TB/HIV coordination, including strategies for TB prevention and control for PLWHA, HIV/AIDS prevention, and health promotion and treatment for TB patients.

#### Links with other health-care providers

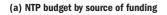
Private providers treat a considerable proportion of patients in metropolitan Ho Chi Minh City, but the situation is uncertain in other parts of the country. A project aimed at involving private providers in TB control in Ho Chi Minh was implemented from 2001 to 2004 with mixed results. Case notification increased, but the treatment success rate was poor in the private clinics involved, probably because anti-TB drugs were not provided free of charge. No other private sector initiatives have been undertaken.

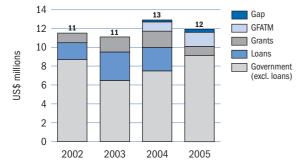
#### Links with the community

The community (i.e. villages, Women's Union, Farmer's Union) is involved in a successful IEC campaign for TB control activities, and there are plans to scale up these activities.

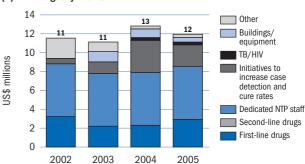
#### **Partnerships**

Effective international partnerships are a major feature of Viet Nam's TB control programme. Viet Nam's longstanding relationship with the Medical Committee Netherlands Viet Nam and, more recently, technical and funding partnerships with KNCV and

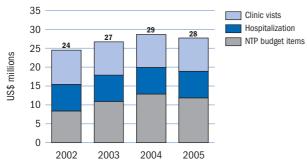




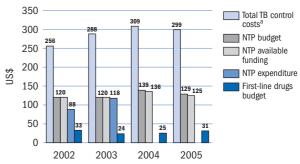
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>



#### (d) Per patient costs, budgets, available funding and expenditures



Total TB control costs for 2002 and 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

#### **VIET NAM**

the Dutch Government, have created nationwide TB services of high quality. A grant from the GFATM (signed in late 2003) is being used to reach TB patients among high-risk groups, remote populations and PLWHA. WHO and CDC provide technical and financial support for TB control activities, and CDC operates a Global AIDS Programme office in Viet Nam. A World Bank loan assists with purchase of anti-TB drugs.

#### **Budgets and expenditures**

The NTP budget has consistently been about US\$ 11-13 million per year between 2002 and 2005. Unlike most other HBCs, there has been no need for large increases in the budget during the period 2002-2005 because case detection and treatment success rates were already at target levels in 2002. Nevertheless, some budget increases have been planned, for example to allow better access to DOTS in remote areas and for a national prevalence survey.

Most funding is provided by the government (including loans), but grants also make an important contribution, and GFATM funding accounts for about 13% of the budget in 2005. Actual expenditures in 2003 were very similar to the planned budget. The NTP budget is consistently about US\$ 120-140 per patient treated, while the total cost of TB control (including a network of dedicated hospital beds for TB patients and visits to clinics for DOT and monitoring during treatment) is consistently about US\$ 250-300 per patient treated. The total estimated cost of TB control has remained stable at US\$ 24-30 million per year.

# **Zimbabwe**

Zimbabwe adopted the DOTS strategy in 1992 and has been reporting nationwide coverage since 2000. TB treatment is provided free of charge to all patients and an adequate supply of anti-TB drugs is assured until 2006. Nevertheless, Zimbabwe still has some way to go to reach the global targets for case detection and treatment success. Many difficulties face TB control efforts, including insufficient funding, severe staff shortages and the impact of the HIV/AIDS epidemic. WHO estimates that, in 2003, 69% of TB patients were HIVpositive. Efforts to address the needs arising from widespread TB/HIV coinfection are still in the developmental stage.

#### **System of TB control**

Zimbabwe's NTP was established in the 1960s. In 1983, the government introduced a policy of integrating all TB control activities into the general health services. The DOTS strategy was officially adopted by the NTP in 1997. The NTP operates at three levels: central, provincial/local authority and district. At the central level, the

NTP is part of the HIV/AIDS/STI and TB unit and is responsible for planning, coordination, monitoring, training and evaluation of programme performance. At the provincial level, training of staff and collection and analysis of TB data are the responsibility of the provincial epidemiology and disease control officer. Four local authorities (Bulawayo, Gweru, Harare and Mutare) run their own TB control programmes, but follow national guidelines and report to the NTP. Mission hospitals, health services of the uniformed forces and some large private organizations also provide TB control services according to national guidelines. The district is the basic management unit for TB control and is responsible for diagnosis, treatment and follow-up of patients, as well as supervision and monitoring of treatment, registration and compilation of quarterly and annual reports. There are rural health centres or municipal clinics in most urban localities that function as primary health-care facilities. These centres and clinics assist in the identification and referral of TB suspects, supervision and observation of treatment and follow-up of contacts and defaulters.

The laboratory network consists of an NRL, 10 intermediate (province/ city) laboratories and 96 peripheral laboratories. All intermediate and peripheral laboratories do smear microscopy and refer re-treatment and failure cases for culture and drug susceptibility testing to the NRL. In addition, the NRL is responsible for providing overall assistance and EQA to all laboratories in the network. There are more than 30 private laboratories that do smear microscopy for private and public providers and that participate in the NTP laboratory network, but they are not involved in the NRL EQA.

#### **Surveillance and monitoring**

The total number of TB cases reported in Zimbabwe rose from 6000 in 1988 to 60 000 in 2002. However, the rate of increase has been slowing since 1997, and the number of reported cases fell between 2002 and 2003. The smear-positive case notification rate has been fairly stable since 1997, so the proportion of cases diagnosed as smear-positive has fallen. This proportion was only 27% in 2003, indicating poor diagnostic technique. In 2003, Zimbabwe experienced nationwide industrial action in the public health sector for three months, which adversely affected diagnosis and treatment of TB. It is not clear whether these trends reflect the underlying trends in incidence or variations in the quality of reporting, but the pattern is similar in some other eastern and southern African countries with high rates of HIV infection. Case detection under DOTS was in the range 40-50% between 2000 and 2003, but further investigation is needed to verify this estimate.

The treatment success rate was 67% for patients registered in 2002 and has remained at this level since 1998. In the 2002 cohort, 11% of patients died and 22% either defaulted or were transferred between

### **PROGRESS IN TB CONTROL IN ZIMBABWE**

#### **Indicators**

DOTS treatment success, 2002 cohort	67%
DOTS case detection rate, 2003	42%
NTP budget available, 2004	58%
Government contribution to NTP budget, including loans, 2004	27%
Government contribution to total TB control costs, including loans, 2004	59%
Government health spending used for TB control, 2004	4%

#### **Major achievements**

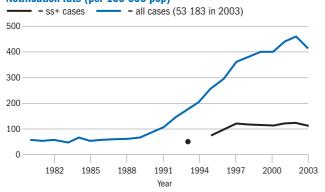
- Training of all laboratory staff and strengthening of laboratory supervision
- Training of prison health workers on DOTS
- Joint MoH/WHO review of the NTP in November 2003

#### **Major planned activities**

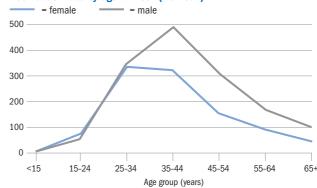
- Strengthen the EQA system in both public and private laboratories
- Improve the recording and reporting system that links the national reference laboratory and public and private laboratories
- Introduce DOTS to prison services and train prison health-care workers
- Introduce community-based DOTS in one pilot district
- Introduce FDCs
- Revise national TB manual
- Train TB microscopists

LATEST ESTIMATES <sup>a</sup>		TRENDS	2000	2001	2002	2003
Population	12 891 242	DOTS coverage (%)	100	100	100	100
Global rank (by est. number of cases)	19	Notification rate (all cases/100 000 pop)	402	441	461	413
Incidence (all cases/100 000 pop/year)	659	Notification rate (new ss+/100 000 pop)	114	120	124	112
Incidence (new ss+/100 000 pop/year)	265	Detection of all cases (%)	65	68	70	63
Prevalence (all cases/100 000 pop)	660	Case detection rate (new ss+, %)	46	46	47	42
TB mortality (all cases/100 000 pop/year)	153	DOTS case detection rate (new ss+, %)	46	46	47	42
TB cases HIV+ (adults aged 15-49, %)	69	DOTS case detection rate (new ss+)/coverage (%)	46	46	47	42
New cases multidrug resistant (%)	1.9	DOTS treatment success (new ss+, %)	69	71	67	_

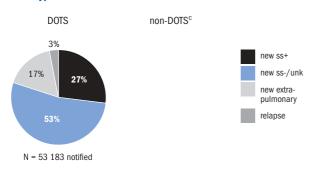
#### Notification rate (per 100 000 pop)



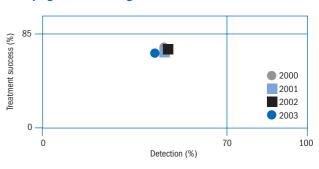
#### Notification rate by age and sex (new ss+)<sup>b</sup>



#### **Case types notified**



#### **DOTS** progress towards targets<sup>d</sup>



#### **DOTS** treatment outcomes (new ss+)



Non-DOTS treatment outcomes (new ss+)

ss+ indicates smear-positive; ss-, smear-negative; pop, population; unk, unknown.

Absence of a graph indicates that the data were not available or applicable.

- <sup>a</sup> See Methods for data sources. Prevalence and mortality estimates include patients with HIV.
- <sup>b</sup> The sum of cases notified by age and sex is less than the number of new smear-positive cases notified for some countries.
- $^{\mbox{\scriptsize c}}$  Non-DOTS is blank for countries which are 100% DOTS, or where no non-DOTS data were reported.
- <sup>d</sup> DOTS case detection rate for given year, DOTS treatment success rate for cohort registered in previous year.
- $^{\mathrm{e}}\,$  "Other" includes transfer out and not evaluated, still on treatment, and other unknown.

treatment centres without follow-up. Among patients registered for re-treatment, 20% were reported to have died, 16% defaulted or transferred without follow-up.

While it would be valuable to assess the impact of DOTS on the burden of TB in Zimbabwe, the immediate priority is to evaluate more accurately the progress made in programme implementation (case detection, treatment success) against the background of changing TB incidence, prevalence and death rates.

#### Improving programme performance

The high rates of HIV infection together with unfavourable socioeconomic conditions have had an impact on general health services in Zimbabwe in the past year, and will also affect TB control activities. A national review of the NTP by MoH/WHO carried out in November 2003 included a review of activities at the central level, in all eight provinces and the three major cities (Bulawayo, Chitungwiza and Harare). Recommendations were made on strengthening existing TB control and collaborative TB/HIV activities in order to reverse the downward trends in case detection and treatment success. Senior ministry officials are committed to improving TB control and a national TB policy, strategic plan and manual have been developed. However, the strategic plan for DOTS expansion has not been adopted nationally and there are serious financial and infrastructural deficiencies at all levels.

There is a severe shortage of human resources at all levels, especially at the central level. The NTP continues to be adversely affected by the departure of experienced staff from the public to the private sectors and to other countries. Five of the eight provincial TB coordinators were appointed in the past year and many districts have no TB coordinators. The NTP is planning to identify districts without coordinators, appoint new staff and ensure that all district hospitals have a staff member responsible for TB. Staffing at the central level has been strengthened by the appoint-

ment of a national TB coordinator to assist the NTP manager and by NTP advisers and officers that have been seconded by IUATLD and CDC. Training for staff has been intensified and efforts have been made to train prison health workers on the DOTS strategy.

IEC material is generally available at most facilities; however, it is produced centrally, which reduces its impact in areas where other languages are spoken. No national advocacy plan has been developed.

The supply of high quality anti-TB drugs is guaranteed until the end of 2006, with funding from the European Union, but FDCs and paediatric formulations are not available. The NTP intends to introduce FDCs in early 2005. The last national DRS was done in 1994-1995, when the prevalence of MDR-TB in previously untreated patients was 1.4%. No recent data on the prevalence of MDR-TB are available, but another DRS is planned for 2005. The draft policy document on MDR-TB management is awaiting finalization. Consequently, no secondline drugs are currently being used.

Other areas where programme performance needs to be improved include diagnostic laboratory services, TB/HIV coordination and links with other health-care providers.

#### Diagnostic and laboratory services

Training of laboratory staff and strengthening of laboratory supervision were undertaken in 2003-2004, but many facilities still have untrained staff. Similarly, while EQA systems were strengthened, financial and staffing constraints mean that some quality assurance activities were not routinely performed or have been suspended at national and provincial levels. A major problem for the laboratory services in Zimbabwe is the shortage of staff associated with the elimination of many posts for microscopists, and the movement of trained staff to the private sector or to other countries. The country is planning to train basic-level TB microscopists in 2005 to help to rectify this problem.

#### TB/HIV coordination

The number of AIDS cases and AIDSrelated deaths continues to increase in Zimbabwe. There is no routine HIV surveillance among TB patients, but WHO estimates that 69% of adult TB patients are infected with HIV. The government has set up units to manage opportunistic infections, including provision of co-trimoxazole and fluconazole to PLWHA, and plans to begin delivery of ART in Harare and Mpilo hospitals in the near future. The government has also signed a policy on the use of co-trimoxazole among HIV-positive TB patients, though not yet on the use of isoniazid preventive therapy in PLWHA.

A TB/HIV working group has been set up and collaborative TB/HIV activities have been planned. To date, few of these activities have started. WHO is funding a community TB/HIV care initiative in one district and HIV surveillance among TB patients is planned for 2005.

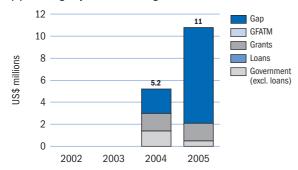
#### Links with other health-care providers

Private laboratories have been included in the NTP laboratory network. A small-scale PPM-DOTS project involving private practitioners and hospitals is being piloted in Harare. The NTP is involving medical colleges, specialist TB hospitals, prison health services, mission hospitals and health services operated by the police and the armed forces in DOTS implementation. A few large agricultural and mining companies also provide TB control services to their employees and dependants according to national guidelines.

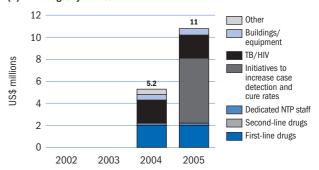
#### **Partnerships**

Technical assistance is provided by IUATLD and WHO. The CDC provides laboratory support (reagents and other consumables) and the EU provides funding for anti-TB drugs. There is a national TB expert committee that guides policy development and implementation, but there is currently no interagency body coordinating TB control. However, a country coordination committee meets monthly and functions as the national TB/HIV coordinating body.

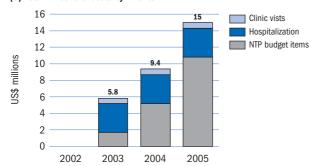
#### (a) NTP budget by source of funding

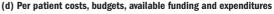


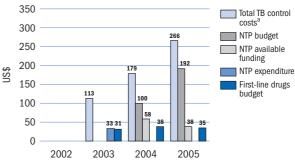
#### (b) NTP budget by line item



#### (c) Total TB control costs by line item<sup>a</sup>







a Total TB control costs for 2003 are based on expenditures, whereas those for 2004 and 2005 are based on budgets. Estimates of the costs of clinic visits and hospitalization are WHO estimates based on data provided by the NTP and from other sources. See Methods for further details.

#### **Budgets and expenditures**

The NTP budget for 2005 is US\$ 11 million, compared with US\$ 5 million in 2004 (about US\$ 200 per patient vs US\$ 100 per patient). The increased budget reflects plans to increase spending on various initiatives, including collaborative TB/HIV activities, training, monitoring and evaluation, and community TB care. However, available funding is limited, at only around US\$ 2 million in 2005. This is down from available funding of US\$ 3 million in 2004, but a slight increase compared with expenditures of US\$ 1.7 million in 2003. The government's contribution to funding is likely to be higher than reported, as finan-

cial support for buildings and equipment is not reflected in disease control programme budgets. Most of the grant funding for first-line drugs is provided by the EU through the essential drugs programme and has remained constant between 2003 and 2005 at around US\$ 1.6 million each year (equivalent to about US\$ 35 per patient treated). At around US\$ 9 million, the funding gap in 2005 is equivalent to 80% of the budget. Zimbabwe is likely to apply to the GFATM in round 5 to address this gap.

The total cost of TB control, which includes the cost of dedicated TB beds and clinic visits during treatment as well as items included in the NTP

budget, was about US\$ 6 million in 2003 (just over US\$ 100 per patient treated). If the 2005 NTP budget is fully funded and spent, this will increase to about US\$ 15 million in 2005 (about US\$ 270 per patient treated). The estimated cost of dedicated TB hospital beds, at US\$ 3.5 million, is based on an estimate of 1660 dedicated TB beds, including those in mission hospitals. However, since occupancy in district hospitals is decreasing (for example because of a new admission policy introduced in the late 1990s) and beds are being reallocated to other diseases, this may be an overestimate.

# Country data by WHO region

**Explanatory notes** 

**Africa** 

**The Americas** 

**Eastern Mediterranean** 

**Europe** 

**South-East Asia** 

**Western Pacific** 

# **Explanatory notes**

Country-specific data grouped by WHO region. For each country we present:

- Estimated burden of TB in 1990 (baseline year for MDG) and 2003 (the latest year covered by this report). Estimates of TB include incidence, prevalence and death rates.
- Population, TB notifications (numbers and rates), detection rates (countrywide and under DOTS), DOTS coverage and percentage of pulmonary cases that were smear-positive (DOTS and non-DOTS) in 2003.
- Treatment outcomes for cases registered in 2002 the new smear-positive cohort for both DOTS and (where available) non-DOTS programmes, and all re-treatment cases (taken together, where available) from DOTS programmes.
- Treatment outcomes for specific re-treatment cohorts from DOTS programmes in 2002 - relapse, after-failure and after-default cohorts.
- Trends in DOTS treatment success (1994–2002) and DOTS case detection (1995-2003).
- Age and sex distribution (numbers) of new smear-positive notifications from DOTS and non-DOTS programmes.
- Age and sex rates of new smear-positive notifications countrywide.
- Notifications (numbers and rates) of TB (all forms) since 1980.
- Notifications (numbers and rates) of new smear-positive cases since 1993.
- Country notes: remarks from respondents that may help to explain data in selected countries' reports.

#### **Notation for 1st table**

Country data ... estimated burden of TB

See Methods section for information on how estimates are derived.

"Incl. HIV+" means including HIV+ TB cases; "Excl. HIV+" means excluding HIV+ TB cases.

Rate: the number per 100 000 population. (For incidence and death rates, the number represents events occurring annually.)

#### **Notation for 2nd table**

Country data ... notification, detection and DOTS coverage, 2003

Pop thousands: the population of the country or territory, expressed in thousands, from the United Nations Population Division, World Population Prospects, 2002 revision.

All cases, country, number a: the total number of TB cases according to the

All cases, WHO, number b: the total number of TB cases notified to WHO (WHO definition of a TB case notification, which includes new and relapse cases and, for the WHO European region only, cases with previous history unknown).

All cases, rate: cases notified to WHO per 100 000 population.

New cases, smear-positive, rate: per 100 000 population.

New cases, pulm confirmed: new pulmonary laboratory-confirmed cases.

Re-treatment cases: see definitions in Methods section. "Other" re-treatment includes cases that were classified as re-treatment but were not specified as relapse, after-failure or after-default.

Other, number: TB cases that were not classified as new or re-treatment. These cases plus re-treatment cases other than relapse cases make up the gap (if any) between the country's count of TB cases and WHO notifications.

Detection rate, all cases: the proportion of all estimated cases (all forms) that were notified.

Detection rate, new ss+: the proportion of estimated new smear-positive cases that were notified.

DOTS % of pop: the percentage of the population living in geographical areas nominally serviced by health facilities implementing DOTS.

DOTS, DDR: the new smear-positive detection rate of DOTS programmes.

DOTS, % of pulm. cases ss+: the percentage of new pulmonary cases in DOTS programmes that are smear-positive.

#### **Notation for 3rd table**

#### Country data ... treatment outcomes for cases registered in 2002

Number of cases notified: the number of new smear-positive cases notified in 2002 and representing, in theory, a cohort of cases for which treatment outcomes should be monitored.

Number of cases registered: the number of new smear-positive cases ultimately reported as constituting the cohort for treatment outcome monitoring which, ideally, should be close the number of cases initially notified (see Methods sec-

% of notif registered: the percentage of notified cases that was ultimately represented by the number of cases registered

% not eval: the difference between the number registered and the sum of the six mutually exclusive outcomes (cured, completed, died, failed, defaulted, transferred-out, see definitions in the Methods section).

## **Africa**

**The Americas** 

**Eastern Mediterranean** 

**Europe** 

**South-East Asia** 

**Western Pacific** 

#### **AFRICA: SUMMARY OF TB CONTROL POLICIES** MONITORING STATUS MANUAL SCCd OUTCOME MICROSCOPY DOTe ALGERIA DOTS YES ANGOLA DOTS YES BENIN DOTS YES BOTSWANA DOTS YES BURKINA FASO DOTS YES BURUNDI DOTS YES CAMEROON DOTS YES CAPE VERDE NON-DOTS YES CENTRAL AFRICAN REPUBLIC DOTS YES CHAD DOTS YES COMOROS DOTS YES CONGO DOTS YES CÔTE D'IVOIRE DOTS YES DR CONGO DOTS YES **EQUATORIAL GUINEA** ERITREA DOTS YES DOTS **ETHIOPIA** YES GABON DOTS GAMBIA DOTS YES GHANA DOTS YES GUINEA DOTS GUINEA-BISSAU DOTS YES KENYA DOTS YES LESOTH0 DOTS NO LIBERIA MADAGASCAR DOTS YES MALAWI DOTS YES MALI DOTS YES MAURITANIA MAURITIUS DOTS NO MOZAMBIQUE DOTS YES DOTS YES NAMIBIA NIGER DOTS NIGERIA DOTS YES RWANDA YES DOTS SAO TOME AND PRINCIPE SENEGAL DOTS YES SEYCHELLES DOTS SIERRA LEONE DOTS YES SOUTH AFRICA DOTS YES **SWAZILAND** DOTS NO TOGO DOTS YES



UGANDA

ZAMBIA

ZIMBABWE

UR TANZANIA

- a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)
- b Manual: national TB control manual (recommended)

YES

YFS

YES

YES

DOTS

DOTS

DOTS

DOTS

- $\ensuremath{\text{c}}$   $\ensuremath{\text{Microscopy}}$  use of smear microscopy for diagnosis (core component of DOTS)
- d SCC: short course chemotherapy (core component of DOTS)
- e DOT: directly observed treatment (core component of DOTS)
- f Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)

**AFR** 

Country data for Africa: estimated burden of TB

	<u>-</u>	idence	1990		Prev	Prevalence, 1990	066	_		Death. 1990	06	L	Incide	Incidence, 2003			Prevaler	Prevalence, 2003			Death, 2003	3	
	All cases incl. HIV+	-	New ss+ incl. HIV+	₹	cases incl. HIV+ All cases excl	IIV+ All c		HIV+	cases incl. HIV+ All	IV+ All	cases excl. HIV+	_	All cases incl. HIV+	_	New ss+ incl. HIV+	₹	cases incl HIV+	All cases excl HIV+	cl HIV+	All cases incl HIV+		All cases excl HIV+	<b></b>
	number	rate	number rate		number	rate	number	rate	number	rate	number	rate	number rate	e unmper	ber rate	number	rate	number	rate	number	rate	number	rate
Algeria	9 094	36		16		83		83	2 383	10	2 383	10	16 790 53				53	16 757		609	2	209	7
Angola	19 729	211		32	•	453		445	5 520	29	4 802	21			15 424 113	.,		34 835		3 446	52	2 699	20
Benin	3 576	77		<b>4</b>		171		171	928	21	915	50			2 600 39			9 200	141	994	15	822	12
Botswana	3 266	241		35		496		480	934	69	704	25					515	6 105		1 530	98	610	8
Burkina Faso	13 142	147		7.7		285		271	4 066	46	2 790	31						39 370		5 912	45	4 342	33
Burundi	6 727	120	2 876 5	51		234		223	2 041	36	1 353	24	23 585 346		10 083 148	38 090	558	35 441	519	6 308	95	4 011	29
Cameroon	7 304	63		72		138		137	1 901	16	1 731	15	ľ					35 443		2 606	35	3 441	21
Cape Verde	263	161	252 7	72	1 277	366	1277	366	147	42	147	42	778 168		349 75	1 526	329	1519		186	40	180	39
Central African Republic	3 318	113		17		225		216	982	33	889	23	12 558 325		5 2 1 9 1 3 5			19 039	493	4 360	113	2 100	25
Chad	10 722	184	4 666 8	90	23 085	397	22 731	390	2 986	51	2 624	45	19 385 225		8 437 98	39 188	3 456	37 752	439	5 645	99	4 151	48
Comoros	449	85		88		193		193	118	22	118	22			172 22	795		794	103	09	00	29	00
Congo	3 285	132	1 423 5	25	6 310 2	253	5 974	240	1 013	41	647	56	14 134 380		6 124 164	19 382	521	18 202	489	2 937	79	2 132	22
Côte d'Ivoire	17 182	137		29	34 571			266	5 038	40	3 602	58						102 713		17 678	106	11 323	89
DR Congo	47 828	128	20 806 5	99			95 349	255	13 661	37	10 328	28	194 627 369		84 667 160	297 861	564	283 284	537		81	31 314	29
Equatorial Guinea	258	158	238 6	37		341		337	154	44	137	39			407 82			1 733			63	191	39
Eritrea	6 874	222						490	1845	69	1 755	22			4 946 119			17 861	431	2 556	62	2 162	52
Ethiopia	60 373	124	26 255 5		131 545	569	129 730	266	16 028	33	14 052	59	251 685 356	109 452		377 030	533	358 001	202	56 146	62	42 508	09
Gabon	1 300	136						295	343	36	304	32			1 326 100			3 2 1 8	242	502	38	302	23
Gambia	1 783	190	8 962	35	ľ	432		432	468	20	466	20	3 325 233		1 485 104			4 808	337	591	14	552	39
Ghana	33 599	220		37		484		480	9 136	09	8 456	22					380	77 193	369	10 572	21	8 513	41
Guinea	7 224	118		22		264		262	1 931	32	1 854	30	19 999 236		8 784 104			33 451	394		25	3 668	43
Guinea-Bissau	1 647	162		14		364		363	437	43	426	42						4 4 7 6			43	203	8
Kenya	26 329	112		48		233		227	7 399	31	2 187	52			83 822 262			262 708			133	28 527	68
Lesotho	2 918	186		4		416		414	744	47	704	45						7 030	390	Ì	601	823	46
Liberia	4 363	204		88		430		420	1 246	28	1 035	48	8 422 250		3 633 108	17 073	3 507	16 287	484		28	1 797	23
Madagascar	22 936	192		32	51 917	134		434	6 032	20	2 987	20						56 494	325		14	6 201	36
Malawi	24 845	263	10 259 10	80		523		502	7 369	78	5 137	72						56 754	469	Ì	107	6 337	52
Mali	26 883	297		31		922		649	7 299	81	6 777	75			571 127			75 738	582	662 6	75	8 346	25
Mauritania	4 755	234		)2		531		531	1 248	61	1 244	61	(1				_	19 220	664	2 232	11	2 120	73
Mauritius	716	89		ω Ω	.	154		154	188	18	188	18						1 664			11	138	=
Mozambique	21 336	158		90		332		323	5 970	44	4 711	35	•	_		`		105 146		•	129	11 708	62
Namibia	3 530	251		102		295		529	838	64	854	61	14 351 722					9 486			122	1 029	25
Niger	9 833	129						290	2 591	34	2 562	33				32 998	_	32 571	272		33	3 603	30
Nigeria	87 333	102				221		218	23 154	27	20 354	24		_	358 126	676 879	546	642 327			82	70 864	22
Kwanda	8 789	300	3 /83	9 7	010 /1	752	16 146	238	2 689	040	1 /49	9 %	31 353 374		13 497 161	55 701		52 643	979	8 732	45 8	5811	9 6
Sopogal	14 703	200		- 00		152	33 187	452	3 874	200	2 824	2 2		11 055		73		410	420	F 061	20 20	C+ 707 V	9 7
Sevchelles	33	44		2 2		1 8		9 6	, «	1 2	- «	1 5			13 16			55.	2,4	4	3 10	4	- LC
Sierra Leone	8 950	221		2 86		497	20 115 4	496	2 368	28	2 322	22	4		9 404 189	40	w	39 485	794	5 047	102	4 386	8
South Africa	68 593	186		92		ľ		411	17 682	48	16 407	45	"				Ĺ	153 694	341	١.	73	12 459	28
Swaziland	2 259	267	888 10	92	4 724	929	4 601	543	633	75	498	59	11 666 1 083		4 585 426	10 687	992	7 361	683		237	893	83
Togo	12 276	355		22		662		797	3 254	94	3 178	95	17 221 351		518 153	34 188	969 8	33 029	673		86	3 633	74
Uganda	27 641	159		66		270		244	9 547	22	4 589	56				168 387	652	160 315	621	24 692	96	18 233	71
UR Tanzania	47 831	183		82		371		358	13 942	53	10 095	39				193 610	524	175 953	476	31 745	98	19 101	25
Zambia	24 333	297		122	40 326 4	492	36 042	440	8 567	104	3 904	48	70 975 656		29 130 269	966 89		54 954	508	13 152	122	6 584	61
Zimbabwe	91/ 21	131	2 207	2		230		710	4 6 14	44	7 381	53	85 015 65	\$		821.28	099	64 4 / 4	000	19 /49	23	698 /	٥.
Region	724 602	146	310 745 6	63 1	525 020	307 1 4	489 099	300	203 451	41	164 627	33	2 371 745 345	1 012 416	416 147	3 486 914	1 507	3 2 1 2 5 5 8	467	538 212	78	351 432	21

								Whole country	Ž.									DOTS	s				non-DOTS	
	G	VIIV	00000	Ü	Smear-positive		New cases	five Evtre	i	Re-treatment cases	Re-treatment cases	ises		Other	Detection rate	te %		Notifications	tions	900	% of		Notifications	% of
	1	Country	WHO	5	ical-positive	0	or unknown	ğ	_	אונפו ומ	mare viter		Ē.	Č	dada New			cases	oo Mak				S MAN S	
	thousands	number	number ra	rate nu	number rate	number	number		per number		number nu	number nu	number nu	number			-	- 1	number	-	cases ss	ss+ number		number cases ss-
Algeria	31 800	19 730				8 924		12 8 861		508	38	98						19 730 62	8 549	27 1	113 8			
Angola	13 625	37 053		_	-		12		_	397	433	541							18 164			31 1 987		807 44
Benin	6 736	3 172	- 1		2 438 36	3 2 506				109	63	177				$\dashv$		- 1	2 438	- 1	Ì		35	
Botswana	1 785	10 030			•		2			208	22	113							3 050			38		
Burkina Faso	13 002	2 765	2 620	50						116	85	09							1 703			98		
Burundi	6 825	6 846		ľ		, TU &				36	٥	18		007		+			3 01 /			20/		
Cameroon	16 018	16 509		1001			.7			672	82	ı		463				15 964 100	10 692					
Cape Verde	463	323	316		165 36	165	111			19	2	2							c			316		165 60
entral Arrican Republic	2 000	71.87								001	40	Ogc				+		1 140 13	320					
Crad	0 220	9644			•					001	7/	007							660					
Comoros	3 724	91	7 782	200	3 477 93	50 0 7	0		20 20	5 75	o 4	7 2			23	57	80	7 782 209	3 477	0 6	57 8	22 22		
Côte d'Ivoire	16 631	18 2 13	1	ľ			2 250			557	274	157				+			10.915	1			784 4	496 85
DR Congo	52 771	86 715				53 578				100	929	965		387				84 687 160	53 578					
Equatorial Guinea	494																							
Eritrea	4 141	4 7 08			887 21	887				101	15	31		3				4 708 114	887			27		
Ethiopia	70 678	118 276	117 600 1				35 141			378	242	434							39 698			23		
Gabon	1 329						683			109	27	55				_			1 233			34		
Gambia	1 426	1 985				3 1 040				24	3	37							1 040			27		
Ghana	20 922	11 891					2 860	30 759		558					27	40	100 11	11 891 57	7 714	37	40 7	73		
Guinea	8 480	6 570	- 1							213	43			213		+			4 495					
Guinea-Bissau	1 493	1 647				963		20 29		128		47						1327 89	715				320 2	248 83
Kenya	31 987	95 3 10	91 522 2				(*)			826	104	1 023		2 661					38 158			21		
Lesotho	1 802	13 341	- 1	999	3 652 203	3 652	5 404			549		371		887		+			3 652			40		
Liberia	3 367										;													
Madagascar	17 404	19 799	19 309 1	111	7 246 64	12 881	1 910	3 580	•	938	134	356		0	51	77	100 19	19 309 111	12 881	4 2	77	87		
alawi	12 103	40.204								000	,	3		2 393		+		- 1	0 1 7 0			- 6		
Mauritania	2 893	4 040	4 430			2013	100			077	0	-							0000			2		
Mauritius	1 221	137	137	-	8	66		17	17	4	-	-					00	137 11		00		35		
Mozambique	18 863	29 107	1		"	16.1	7			176	182	323				H		1				37		
Namibia	1 987	12 931								765				1 155				11 776 593				25		
Niger	11 972	7 423					1 070			397	156	189							4 505			31		
Nigeria	124 009	46 596		.,	28 173 23		13		1 525 1 2	210	888	1 263		261	12	18	60 44	44 184 36	28 173	23	18 6	89		
Rwanda	8 387	6 046	2 895		3710 44	3 710	652			339	45	34		72								35		
Sao Tome & Principe	161																							
Senegal	10 095	96.46			6 587 65		1 421		951 4	421		367						9 380 93	6 587			32		
Seychelles	81	10				6				0	0	0							2			99		
Sierra Leone	4 971	5 421						388 388		93						+		5 289 106	3 113					
South Africa	45 026	255 422		÷		3 116 364				730 1			23 182						116 331				42	33 87
Swaziland	1 077	7 869	7 749 7	719	1 585 147		4 582	32 1012		570		83			99	35	100 7	7 749 719	1 585	147	35 2	56		
logo	4 909	42 004				20 320	125			101				1 006		+			702 00			- 45		
Ugarida IID Tanzania	36 977	64 665						_		310	128	250		2 708					27 800			3 2		
Zambia	10.877	58 032								2 168	43.1	2004		3,669				53 932 499	18 934			43.2		
Zimbabwe	12 891	57 117								000				0000					14 488					
		5								222				‡5000		_			25++			<u> </u>		

See Explanatory notes, page 151.

Country data for Africa: treatment outcomes for cases registered in 2002

New smear-positive cases – DOTS	100. 1100	וופוור סמ	New Year	v smear	positive	New smear-positive cases – DOTS	DOTS				-			New sme	ar-positi	New smear-bositive cases – non-DOTS	- non-D	STC			$\mid$		Sme	Smear-positive re-treatment cases - DOTS	e re-treat	ment cas	ses – DO	12		Г
			%	%	%	%	1	%	%	%				%	%	%	%	%	%	%	%		%	%	%	%	%		%	Τ
	Number of cases		of notif		compl-				trans- not			Number of cases		of notif	com				trans-	not		Number		-ldmoo			Ħ	- not		
	notified	regist'd r		cured		died fa	failed de	default fer		al success	Ľ	notified reg	9	ist'd cured	red eted	ed died		failed default ferred	ferred	- 1	saccess		cured		died fail	failed default	- 1		saccess	SS
Algeria	8 246	9 200	112	72	17	2	0	4	4	w	39											644	92	13	4	1	3			
Angola Benin	17 345	17 345	100	65	9	ი ყ	- 0	17 50	7 7	~ 8	7.4	742										2 320	40		ĸ	9 5	0 %	58		- v
Botswana	3 334	3 458	104	35	36	6	-		12		2,1										$\perp$	294	30		15				99	2
Burkina Faso	1 544	1 518	98	37	7	12	e c	5 4	9 -	w r	64											185	51	8 %	11 5	8 13	3 10		58	00 6
Cameroon	7 365	7 365	100	61	8	9	2	20	-	2 7	0,	556										185	29		10	6 12			5   59	, o
Cape Verde Central African Republic	2 657											111	34	31	53	6	•	18	21		23	12	29			w	.,	10	67	
Chad	3 4 1 7	791	23	22	20	2	-	1	11		72	102 2	2 630 2	2 578 1	14 4	40 8	3 18	2	15		54	88	42	47	8	-	2 2	4	89	6
Comoros	5.019	5.019	100	8 %	- 4	നെ	-	- 6	4	J, [	96											132	5	17	œ	4		_	7.	00
Côte d'Ivoire	10 255	10 236	100	25	13	7	- 2	16	- 8	9	37	771	336		6	7 3	3 2	29	15		16	486	38	12		11 26		. 9	20	
DR Congo Equatorial Guinea	44 518	45 013	101	71	00	7	-	00	2	2 7	82											4 618	61	9		4 10		ю.		
Eritrea	646	853	132	73	6	9	2		4	3	32											119	62	8	3		3			_
Ethiopia	36 541	36 541	100	59	17	<b>~</b> 2		5	10	2 2 7	76											1 716	52	6	7	8		2 22	09	0
Gambia	1 035	1 035	100	67	7	2	6		7		7.4											149	41		20	-	3 35		4	T-
Ghana	7 7 32	7 732	100	22	2	80	2	15		10 6	30																			
Guinea	4 300	4 246	66	61	10	6	-		6	-	72											340	20	1	12	5 12		6	6	_
Guinea-Bissau	532	532	100	33	4	e .	0	19	30	7	48	367	367	100	38 3	30 13	~	4	2		89	103	38	28	7	16	17		99	9
Kenya	34 337	30 966	90	92	4 6	ω ;	0 1	<b>ດ</b> ເ	٠ ـ ١		62											2 4 76	92	12	10	0		·	-	_
Lesotho	3.16/	3 16/	100	RZ	23	-	-	٥		2/ 2	70										+									Т
Liberia Madagascar	11 387	11 145	86	92	6	2	9	15	m	7	4		69	4)	57 1	10 7		23	ю		29	975	19		4				99	0
Malawi	7 703	7 703	100	20	3	19	-	4			72											862	99	4	24	1		_	71	_
Mali	2 7 5 7	2 757	100	33	18	2	2	19			20											207	8	13	9	1 29		8	47	_
Mauritania	98	86	100	88	c	2	2	c		O.	- 20											4	20		4.	50			22	
Mozambique	15 236	15 236	100	11	-	11	-	7			78											1721	92	-					19	_
Namibia Niger	4 690	4 649	66	45	19	ω	2	13	7	6	32											849	45	24	12	4	12	6		9
Nigeria	19 596	20 559	105	69	10	2	7	1	_			2 340										2 373	63	=	2					<sub>0</sub>
Rwanda Sao Tome & Principe	3 956	3 975	100	48	10	7	-	4	21	6	28	42										376	40	4	10	4	4 22	17	44	4
Senegal	5 796	5 796	100	59	∞	4 [	_	17	7 20	5	36											924	43	9	2	2 20	0 7	, 17	49	6
Sierra Leone	2 938	2 915	66	60	21	, T	2		0	, ω	÷ ÷											170	52	=	9			e	9	0
South Africa	929 26	060 86	100	72	14	6	-		6	9		1 143 1	1 239	108 5	59	6 5	5 2	13	80	9	65	28 755	43		11		_	7 (		<sub>6</sub>
Swaziland	1410	1 412	100	16	£ -	o £	0 0	4 t	30	4 C	17	782										367	35 2		∞ m	1 14		47		o 10
Uganda	19 088	19 098	100	30	31	9	0		7		30											2 555	58		10					2
UR Tanzania	24 136	24 136	100	92	4	, <del>L</del>	0	4	4	. 0												2 081	17		13			-		
Zambia Zimbabwe	11 694 15 941	11 694 15 941	100 100	67	16 6	2 5	0 2	2 7	15	w w	83 4	4 657 4	4 657	100	22	16 13	2	7	2		71	1 577	63 58	5 5	14 20	2 -	£ 80	4 6	76	ဖွာ
Region	442 729	437 873	66	09	13	7	-	1	7	,	73 11	11 714 9	9 332	80 4	42 2	22 10	8	10	œ	-	63	59 574	49	1	10	2 14		8	59	- a
Neglo:	741 444	5	3	3	2	-	-	-	-		4		400					2	,	-	3	1 20 00	ř		2	,			1	_

See Explanatory notes, page 151.

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				Kelabs	Kelabse - DOIS								Arrerran	Atter failure - DO IS	200			_					Atter detault - DOTS	200			
	number	%	%	%	%	%		%	%	number	%		%	%		%	%	%	number	%			%	%	%	%	%
	regist'd	Ď	compl- eted	_	ъ	≒	trans- ferred	not eval su	SS	regist'd	φ	compl- eted	died	ъ	=	trans- ferred	not	snccess	regist'd	cured	compl- eted	0	ъ	=		not eval su	snccess
Algeria Angola Benin	510	70	13	ю	-	0				46	41	1	7				15	52	88	45	17	9		30	-		63
Botswana Burkina Faso Burundi	66	54	ю	12	10	တ	13		25	54	52	თ	7	=	15	9		61	38	42	16	13		21	∞		28
Cameroon Cape Verde Central African Republic																											
Chad	24	79	13	4	4				95	2	80	20						100	09	23	63	က		က	7		87
Congo	132	61	17	9		15	-		78																		
Côte d'Ivoire DR Congo Equatorial Guinea	3 124	64	ß	თ	2	7	80	4	02	581	49	2	13	12	12	∞	-	54	913	56	∞	თ	2	17	2	2	64
Eritrea Ethiopia	82	83	7	-	4	4	-		06	12	29	∞	∞	∞			∞	75	25	72	12	4	4	4	4		84
Gabon	34	50		26		œ	8		20	e	67		33					67	112	88		4	-	ď	41		8
Ghana Guinea	5	3		2			2		3	<b>o</b>	5		3					5	1	3		2	-	<b>o</b>	F		3
Guinea-Bissau Kenya	67 2 476	42	31	ε Q	c	6	15		73	-	100							100	35	29	23			59	20		21
Lesotho			!																								
Liberia Madagascar Malawi	862	99	4	24	-	4	-		12																		
Mali	207	34	13	9	_	59	80	8	47																		
Mauritania Mauritius	2	20			20				20	-				100					-	100							100
Mozambique Namibia Nisor	1 148 849	68	1 24	13	2 4	12	9	1 3	99	202	62		œ	7	11	2	9	62	371	09	2	11	2	12		13	62
Niger									$\dagger$																		
Rwanda Sao Tomo & Dringing																											
Special Car Integral									$\dagger$																		
Sevchelles																											
Sierra Leone	72	69	10	4	9	7			62	22	4	2	18	6	27			45	9/	38	13	4	6	59	7		51
South Africa	12 562	44	13	10	-	12	6	12	99	1 010	33	11	12	2	13	14	15	44	3 075	34	12	6	2	24	80	12	46
Swaziland	43							100		-							100		16							100	
Jaanda	1771	29	25	6	-	14	4	19	54								2		2							2	
UR Tanzania	1 703	92	ю	13	0	4	ю	0	79	140	09	2	17	4	2	7	7	65	238	42	30	1	-	1	9		71
Zambia Zimbabwe	1 577 1 371	63 58	13	14 20	7 -	ကထ	4 თ		63																		

See Explanatory notes for previous table, page 151.

Country data for Africa: trends in DOTS treatment success and detection rates, 1994–2003

•		TOT	S Mon S	no ar-no a	itivo tros	OTS was seen and still a transfer of the second sec	(70) 3303				F	S Mon S	noor-noon	osco oviti	NOTS now smear-neatting association rate (0/)	n rate (0/		
1	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
Algeria			86			87	87	8	68			134			126	115	114	113
Angola				15	89		89	99	74			69	44	28		85	118	118
Benin	92	73	72	73	77	77		79	80	94	92	92	91	6	98		96	94
Botswana	72	29	20	20	47	71	77	78	71	9/	87	87	88	72	9/	71	9/	89
Burkina Faso		25	59	61	29	61	09	92	2	7	19	15	17	18	19	18	17	18
Burundi	44	45		29	74		80	80	79	20	25	31	19	39		35	30	30
Cameroon				80	75	75	77	62	02		2		7	21	35	43	63	98
Cape Verde																42		
Central African Republic		37					22	61			64					<b>о</b>	54	9
Chad	63	47			64				72	34	14			42			42	=
Comoros	94	06		82		93	93	95	96	24	22		24		49	25		37
Congo	69					61	69	99	71	99			49		98	81	88	22
Côte d'Ivoire	17	89	26	61	62	63		73	29	54	53	49	49	46	36	10	39	39
DR Congo	71	80	48	64	20	69	78	77	78	43	49	46	22	26	53	28	22	63
Equatorial Guinea	88	88	77	82						75	69	71	98					
Eritrea				83	73	4	9/	8	82			က	4	13	14	16	14	18
Ethiopia	74	61	73	72	74	9/	80	92	76	16	21	23	25	56	35	35	36	36
Gabon								48	4/								83	93
Gambia	74	9/	80	20				71	74	9/	69	72	9/				73	02
Ghana		54	51	48	29	22	20	26	09	16	14	32	33	31	39	41	41	40
Guinea	78	78	75	74	73	74	89	74	72	46	53	52	54	53	22	53	52	51
Guinea-Bissau						35		21	48						46		43	22
Kenya	73	75	77	92	77	78	88	8	62	23	54	20	24	25	44	47	46	46
Lesotho	26	47	71	63		69		71	52	62	72	83	75		74		99	70
Liberia		79		75							31		46					
Madagascar	51	22		64			20	69	74	25	99		20			72	71	77
Malawi	22	71	89	71	69	71	73	20	72	38	40	42	46	41	40	40	36	35
Mali	89	29	92	62	20	89		20	20	14	16	18	17	16	15		17	18
Mauritania Mauritius	96				91	87	93	68	26	34			32	35	33	24	25	28
Mozambione	67	30	54	67		71	75	77	78	56	50	48	48		45	44	45	45
Namibia	;		99	64	69	. 89	25	. 69	29	23	85	98	87	84	. 5	. 22	85	98
Niger			57	99	1	09		1			1	21	17		37			72
Nigeria	99	49	32	73	73	75	6/	6/	6/	12	12	12	12	14	14	14	13	18
Rwanda			61	89	72	29	61		28	32	35	4	54	45	34	27	31	27
Senegal	35	30	41	52	48		52	53	99	67	71	61	60	53		9	54	50
Severales		68	100	100		06	8	67	45	5	. 8	000	69	)	85	9 6	71	40
Sierra Leone	75	69	74	79		75	1 12	8	. 18	28	41	40	37		34	34	34	33
South Africa			69	73	74	09	99	65	89			9	22	70	75	81	105	118
Swaziland								36	47								34	35
Togo	45	09	99	99	69	9/		22	89	15	15		14	13	14		9	17
Uganda			33	40	62	61	63	26	09			28	28	28	20	46	46	44
UR Tanzania	80	73	9/	77	92	78	78	81	80	22	54	51	52	20	47	46	43	43
Zambia					20	73	69	75	83				5	49	46	46	41 74	42
						)	8		5				5	2	2	2		į
Region	29	62	22	63	20	89	72	74	73	24	26	29	35	36	37	40	47	20

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				MALE						щ	EMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	e2+
Algeria	40	1316	1 633	902	429	231	328	74	1 0 1 7	702	326	242	241	356	114	2 333	2 335	1 032	671	472	684
Angola	380	2 282	2 484	1841	1 018	483	344	551	2 983	2 525	1 660	1111	374	128	931	5 265	600 9	3 501	2 129	857	472
Benin	20	266	504	370	188	117	95	32	226	304	150	93	47	25	52	492	808	520	281	164	117
Botswana	22	203	552	446	244	136	78	32	338	524	276	104	52	43	54	541	1 076	722	348	188	121
Burkina Faso	14	148	313	321	162	129	80	19	102	131	132	20	46	36	33	250	444	453	232	175	116
Burundi	32	348	572	488	260	106	35	75	308	361	276	119	27	10	107	656	933	764	379	133	45
Cameroon	100	1176	2 2 7 4	1516	788	330	160	136	1 273	1 542	745	363	217	72	236	2 449	3 816	2 261	1 151	247	232
Cape Verde																					
Central African Republic																					
Chad	39	64	105	137	92	48	19	28	52	91	124	65	38	13	29	116	196	261	141	98	32
Comoros	_	7	12	2	_	ဇ	က	0	2	7	~	~	_	_	~	12	19	9	2	4	4
Congo																					
Côte d'Ivoire	205	1 200	2 009	1 473	799	402	362	141	1152	1 546	845	425	213	143	346		3 555	2 318	1 224	615	202
DR Congo	854	5 885	8 427	6 193	3 776	1 836	1 047	1 233	6 630	7 711	4 826	2 866	1 457	265	2 087	12 515	16 138	11 019	6 642	3 293	1 639
Equatorial Guinea	-	8	L	L	,			10	00,							0.70	, 00	L	007	0	i.
Eritrea	\r	96	82	22	46	44	36	17	120	149	100	09	36	77	44	017	734	155	106	200	200
Ethiopia	1 110	6 923	6 648	3 7 3 7	2 022	926	483	1 387	5 936	2 908	2 780	1 239	412	137	2 497	12 859	12 556	6 517	3 261	1 388	620
Gabon	14	165	225	149	103	48	22	16	138	144	107	51	33	18	30	303	369	256	154	81	40
Gambia	3	162	236	149	83	52	31	8	81	85	62	39	27	17	11	243	321	211	122	6/	48
Ghana	79	579	1 265	1234	924	609	44	83	487	744	586	380	200	203	162	1 066	2 009	1 820	1 304	400	644
Guinea	8	617	1 052	671	368	172	134	53	353	451	307	137	106	40	87	970	1 503	876	202	278	174
Guinea-Bissau	2	81	116	94	73	44	17	9	9/	63	28	36	28	18	11	157	179	152	109	72	35
Kenya	341	4 918	8 515	4 560	2 167	928	292	487	5 003	6 023	2 618	1 171	551	309	828	9 92 1	14 538	7 178	3 338	1 479	876
_esotho	10	219	614	265	466	219	83	32	328	292	313	219	69	33	42	547	1 181	902	685	278	116
Liberia																					
Madagascar	123	1 249	1830	1839	1413	723	438	216	1164	1 578	1 240	743	326	191	339	2413	3 408	3 079	2 156	1 049	629
Malawi	43	296	1374	936	489	209	128	9/	963	1 531	790	374	155	52	119	1 559	2 905	1 726	863	364	180
Mali	32	348	619	438	330	201	115	58	172	278	212	123	73	45	61	520	897	099	453	274	160
Mauritania	•	C	,	7	7	7	C	•	(	c	•	•	c	L	,	Ļ	ć	,	č	,	7
Mozambigue		D	71	2	=	=	D	-	D	0	4	4	2	n	-	0	707	<u>+</u>	7	<u>+</u>	4
Namibia	24	344	1 033	770	383	168	89	41	413	862	494	202	83	6	65	757	1 895	1 264	585	251	160
Niger	14	485	1 051	277	512	299	169	30	201	356	279	177	83	45	71	989	1 407	1 058	689	382	211
Nigeria	267	3 263	5 388	3 590	2 106	1 139	719	356	3 394	3 956	1 973	1 159	536	327	623	6 657	9 344	5 563	3 265	1 675	1 046
Rwanda	32	364	517	424	270	83	48	36	312	340	161	62	41	17	68	929	857	585	349	124	65
Sao Tome & Principe																					
Senegal	20	1 005	1 438	968	531	293	250	7.7	629	009	398	212	122	98	127	1 634	2 038	1 294	743	415	336
Seychelles	0	~	0	0	_	2	0								0	_	0	0	-	7	0
Sierra Leone	19	351	564	481	264	149	77	56	308	394	249	122	77	32	45		928	730	386	226	109
South Africa	1 767	10 105	20 389	17 858	9 535	3 600	1 491	2 341	12 599	16 863	9 204	4 080	1 972	1 171	4 108	22 704	37 252	27 062	13 615	5 572	2 662
Swaziland	15	120	298	171	96	48	19	4	242	325	145	09	20	∞	59	362	623	316	156	89	27
Togo	10	126	229	192	120	99	22	15	102	149	80	22	26	28	25	228	378	272	175	95	82
Uganda	261	1 643	4 142	3 0 1 1	1 578	719	201	377	1770	3 1 7 6	1 815	749	356	214	638	3 413	7 318	4 826	2 327	1 075	715
UR Tanzania	181	2172	4 964	3 7 2 8	2 166	1 237	1 025	244	2 063	3 504	1 833	929	509	84 44	425	4 235	8 468	5 561	3 095	1 746	1 369
Zambia	302	1 733	4 182	2 390	995	386	308	292	2 061	3 4 39	1 626	680	297	243	594	3 794	7 621	4 016	1 675	683	551
Zimbabwe	133	8/4	3 048	7 7 7 8	198.1	307	502	180	1 232	7 850	1 480	202	677	114	313	2 100	5 904	3 /08	1 246	285	3.18
Region	6 620	51 437	88 719	64 478	35 780	16 513	6866	8 771	54 239	69 793	38 275	19 104	690 6	5 227	15 391 1	105676 1	158 512 1	102 753	54 884	25 582	15 216
															l	ı	ı			l	

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

Country data for Africa: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers)

				;   <u>.</u>			+							-							
	0-14	15-24	25-34		45-54	55-64	+59	0-14	15-24	25-34 3		45-54 5	55-64 6	ę2+	0-14 1	15-24 25	25-34 35-	4	45-54	55-64	65+
Algeria Angola Benin	59	73	114	29	72	59	17	40	92	116	87	46	21	-	69	168	230	154	118	20	18
Botswana Burkina Faso Burundi																					
Cameroon Cape Verde Central African Republic	ъ	12	32	32	6	7	8	-	9	7	13	7	4	11	4	18	39	45	16	11	19
Chad Comoros Condo	116	192	323	412	227	143	69	84	154	272	373	194	113	88	200	346	595	785	421	256	97
Côte d'Ivoire DR Congo Equatorial Guinea	7	49	94	65	32	21	41	o o	39	89	44	56	15	13	16	88	162	109	28	36	27
Eritrea Ethiopia Gabon																					
Gambia Ghana Guinea																					
Guinea-Bissau Kenya Lesotho	4	20	37	24	35	19	10	-	21	19	20	22	10	9	ω	14	26	44	22	59	16
Liberia Madagascar Malawi																					
Mali Mauritania Mauritius																					
Mozambique Namibia Niger																					
Nigeria Rwanda Sao Tome & Principe																					
Senegal Seychelles Sierra Leone																					
South Africa Swaziland Togo	2	2	က	4	5	4	4	0	-	4	က	0	0	-	2	က	7	7	5	4	2
Uganda UR Tanzania Zambia Zimbabwe																					
Region	161	348	603	604	380	223	112	135	316	486	540	295	163	70	296	664 1	1 089 1	1 144	675	386	182

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

178 73 274 45 60 79 80 50 111 111 80 85 176 176 319 77 68 35 35 42 234 158 48 132 128 170 55-64 45-54 43 76 289 71 297 43 99 62 77 111 117 95 93 193 182 590 127 75 354 250 250 200 154 308 308 35-44 323 84 424 47 47 148 157 89 259 106 200 200 140 90 90 157 157 264 656 175 66 110 51 84 388 62 62 286 180 180 407 57 75 25-34 304 96 409 26 1112 174 60 229 135 195 161 65 65 121 121 316 520 186 55 516 454 56 219 164 164 343 8 8 8 133 133 133 133 133 143 173 2 8 2 32 16 14 23 25 63 27 27 140 17 8 8 82 82 74 12 56 61 63 30 94 62 70 105 33 33 71 71 74 + Country data for Africa: smear-positive notification rates (per 100 000 population) by age and sex, 2003 146 40 40 20 20 21 59 45 142 28 122 73 73 63 63 104 114 33 83 22 28 151 151 87 82 82 71 71 137 30 32 33 32 285 444 444 23 23 70 70 43 57 113 73 54 50 50 130 121 310 103 37 206 164 35 35 122 88 88 236 157 38 38 43 89 80 80 57 78 78 122 185 358 41 325 297 297 36 214 116 455 321 25-34 48 49 54 302 71 71 15 140 140 218 126 149 84 48 79 79 83 83 259 430 192 33 433 444 190 134 479 337 234 32 32 8 8 40 76 124 85 102 60 60 42 69 69 131 13 18 27 32 266 193 20 20 68 68 53 77 78 0-14 222 107 386 63 48 60 105 34 178 52 85 130 130 120 130 131 131 154 40 52 220 118 82 82 171 171 262 263 213 64 87 214 108 118 118 1100 143 212 73 177 152 120 110 232 232 198 615 91 114 186 49 60 336 238 238 72 190 193 168 170 56 128 293 102 450 71 71 165 94 99 228 164 136 262 262 252 252 154 119 510 369 82 288 226 389 310 124 202 202 124 124 194 350 345 130 71 71 194 133 200 93 161 66 130 671 522 90 359 248 671 494 25-34 305 122 423 37 37 141 101 240 144 243 240 82 82 181 160 373 643 179 76 137 65 102 479 69 248 196 196 349 99 123 121 72 72 73 129 104 181 38 99 11 46 70 23 51 26 41 25 44 98 25 25 63 63 55 149 56 Cape Verde Central African Republic Rwanda Sao Tome & Principe DR Congo Equatorial Guinea Botswana Burkina Faso uinea-Bissau Seychelles Sierra Leone South Africa JR Tanzania Sôte d'Ivoire **Aozambique** ladagascar **Aauritania** Somoros lauritius lamibia thiopia Sambia esotho ambia. Suinea /alawi Iganda Surundi Shana iberia enya

83 93 17 17 17 18 13 13

51

88 27 27 31

148 148 156 168 168

8 2

Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available

Region

1985   1985		1980	1981	1982	1983	1984	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002
1,005   1,00	Algeria	2 702		13 916	13 681	13 133	13 832	12 917	11 212	11 325	11 039	11 607	11 332	11 428	13 345	13 345	13 507						18 250 1	
1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1,	Angola	10 117	7 501	7 911	6 625	10 153	8 653	9 363	8 510	8 184	9 587	10 271	11 134	11 272	8 269	7 157	5 143							29 996
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1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1,	Botswana	2 662	2 605	2 705	2 883	3 101	2 706	2 627	3 173	2 740	2 532	2 938	3 274	4 179	4 654	4 756	5 665	6 636						
The control of the	Burkina Faso	2 577	2 391	2 265	3 061	877	4 547	1018	1 407	949	1 616	1 497	1 488		1 443	861	2 572	1 814	1 643				2 406	2 376
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Figure   F	Cape Verde	516	344	393	230	285	259		285	276	210	221					303	179	196	205				195
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10   10   10   10   10   10   10   10	OR Congo	5 122	3 051	9 902	13 021	20 415	26 082	27 665	27 096	30 272	31 321	21 131	33 782	37 660	36 647		42 819							
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State   Stat	Ethiopia	40 096	42 423	52 403	56 824	65 045	71 731	80 846	85 867	95 521	80 795	88 634	900 09	900 09	Î		26 034							
State   Stat	Gabon	865	96/	161	725/	654	855	69/	864	127	912	91/	906	976	972	1 034	1 115				1 598			
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1,000   1,00	Ghana	5 207	4 041	4 345	2 651	1 935	3 2 3 5	3 925	5 877	5 297	6 0 1 7	6 407	7 136	7 044	8 569	17 004	8 636						11 923 1	11 723 11 891
11   10   10   10   10   10   10   10	Guinea	245	1 884	205	376	203	131/	1 240	752	770	1 262	1 163	1 246	1 050	3 107	3 300	3 523							
The color   The	Guillea-Dissau	11 049	10 027	502	11 966	000	10.460	10 022	10 515	10 957	12 592	11 788	12 320	14 599	20.451	22 030	28 142						73 017 B	
T74         102         885         885         425         232         334         884         417         6 07         1 06 <td>-esotho</td> <td>4 082</td> <td>3 830</td> <td>4 932</td> <td>3 443</td> <td>2 923</td> <td>2 927</td> <td>21</td> <td>225</td> <td>2 346</td> <td>2 463</td> <td>2 525</td> <td>2 994</td> <td>3 327</td> <td>3384</td> <td>4 334</td> <td>5 181</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>10 111 12 007</td>	-esotho	4 082	3 830	4 932	3 443	2 923	2 927	21	225	2 346	2 463	2 525	2 994	3 327	3384	4 334	5 181							10 111 12 007
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132   157   121   152   148   149	Mauritania	7 576	9 427	2 327	2 333	3 977	4 406	2 2 5 7	3 722	3 928	4 040	5 284	3 064	4 316	3 996		3 849	3 837				3 067		
que         7 457         6 884         5 787         5 204         6 645         8 263         1 5 89         1 5 89         1 6 60         1 5 88         1 7 18         1 7 88         1 7 88         1 7 88         1 7 88         1 8 84         1 8 84         1 8 84         1 6 80         1 8 84	Mauritius	132	157	121	152	118	111	119	117	114	129	119	134	130	159		131							
The color of the	Mozambique	7 457	6 984	2 787	5 937	5 204	5 645	8 263	10 996	13 863	15 958	15 899	16 609	15 085	16 588		17 882	_					22 094 2	25 544 28 602
717         2871         754         673         665         670         556         671         686         670         668         570         668         670         680         680         680         680         680         680         680         680         680         680         680 <td>Namibia</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>4 840</td> <td>4 427</td> <td>3 640</td> <td>2 815</td> <td>3 703</td> <td>2 671</td> <td>2 500</td> <td>1 756</td> <td>2 200</td> <td></td> <td>1 540</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>	Namibia						4 840	4 427	3 640	2 815	3 703	2 671	2 500	1 756	2 200		1 540							
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1 058 1170 497 2 029 1 1 392 1 4 6 4 3 066 1 0 4 5 1 1 4 7 4 0 1 9 0 1 6 20 662 2 1 5 7 9 6 9 4 25 3 16 27 3 8 3 9 3 7 5 7 3 8 1 3 9 3 3 7 5 7 3 8 4 1 3 9 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Тодо	208	126	204	174	343	745	296	1 184	1 071		1 324	1 243	1 223	1 005		1 520							
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4 US7 4 US1 4 577 3 881 5 694 4 759 5 233 5 848 6 UO2 6 822 9 132 11 710 16 237 2 0 125 2 3 959 3 0 8 3 1 3 5 7 3 5 4 3 7 6 2 8 5 5 5 5 5 5 0 8 8 7 3 5 9 5 9 3 6 8 7 7 3 5 9 9 7 5 0 138 5 0 8 5 5 5 5 5 0 8 8 7 3 5 9 7 5 0 7 8 7 5 0 7 8 7 5 7 5 7 7 7 7 7 4 4 4 4 4 4 4 4 4 7 3 7 4 4 7 5 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	Zambia	5 321	6 162	6 525	0 8 9 0	7 272	8 246	8 716	10 025	12 876		16 863	23 373		30 496		35 958						46 259 5	54 220 53 932
219 802 224 102 240 263 258 842 264 928 296 627 301 683 333 842 373 550 365 432 418 530 412 414 432 997 418 995 550 183 504 309 585 773 598 024 687 391 750 780 782 291 40 41 43 44 41 43 44 41 43 49 41 37 41 38 45 44 42 45 41 37	Zimbabwe	4 057	4 051	4 577	3 881	5 694	4 759	5 233	5 848	6 002		9 132	11 710		20 125		30 831		762	077				
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	1980	1981	1982	1983	1984	1985	1986	1987	1988 1	1989 1	1990 1	1991	1992 19	1993 19	1994 19	95 19.	96 19	97 18	98 19	399 20	00 20	01 20	02 2	2003
Algeria	4		20	99	62	63	22	48	48															62
Andola	144	103	105	25	126	104	110	80	6													170		965
Benin	53	52	49	84	49	51	52	45	49	43	45	45	49	46	40		. 14	3 68	39	42	43		43	44
Botswana	270	255	256	264	275	232	218	256	214															252
Burkina Faso	38	34	32	42	12	29	13	17	7													20		20
Burundi	19	15	22	23	40	48	51	53	20															00
Cameroon	28	25	41	36	8	34	21	36	45															00
Cape Verde	178	117	132	9/	95	82		87	82															89
Central African Republic	28	32	61	29	18	20	59	18	59															102
Chad	2	9	က	4	53	30	25	20	25															54
Comoros									43															12
Congo	41	9	193	209	136	125	143	154	166															603
Côte d'Ivoire	20	20	54	62	09	55	56	57	56											97	82 1	103		107
DR Congo	18	7	34	43	65	81	84	80	98															091
Equatorial Guinea					61	2	0	က	9															
Eritrea																								14
Ethiopia	112	116	139	146	162	173	189	194	209													141		99
Gabon	124	11	103	66	83	105	92	100	8	66			91							130			156	164
Gambia	37	6																						36
Ghana	47	35	37	22	15	24	59	42	37	40														57
Guinea		39	30	16	23	25	21	22	8	32														77
Guinea-Bissau	81	57	25	4	42	59	144	80	81	138	114	119									93			19
Kenya	89	29		92		53	49	49	20	22														983
Lesotho	320	293	367	250	207	203	1	15	154	159														999
Liberia	41	52	42	43		20	1	18																
Madagascar	100	80	37	36	98	31	35	36																
Malawi	77	79	89	02	63	74	82	93														224 2	207	213
Mali	12	13	က	7	24	20	23	30																35
Mauritania	471	572	138	135	225	243	122	196																
Mauritius	14	16	12	15	12	11	12	11																11
Mozambique	62	99	46	46	40	43	62	83															138	152
Namibia						424	373	294		273		171	117							542 5		9 029		263
Niger	13	20	13	7	=	=	∞	∞																29
Nigeria	15	16	16	15	16	20	18	25					16				15					39		36
Rwanda	59	26		54	22	22	40	21																20
Sao Tome & Principe	139	38	41	29	48	39	80	51																
Senegal	36	45	28	40		17	14	91																93
Seychelles	25	0	25	54	15	15	35	20																12
Sierra Leone	23	26	26	6	23	24	10	က																90
South Africa	190	201	209	199	195	180	163	166	174															202
Swaziland		23	483	299			:	144	171									308 3						19
Togo	∞	2	∞	9	12	25	19	37	33															36
Uganda	∞ ;	6	4	15			တ	თ	19															162
UR Tanzania	61	62	28	24	26	62	89	72	72															191
Zambia	88	100	102	104	107	117	120	134	167	179	206	277	293	342	385	384 47	421	360	7 7	442 4	478 47	438 5	507	499
ZIIIDabwe	2	5	60	7	ò	5	5	70	5															2
Region	29	29	61	4	49	69	89	73	8	92	84	81	83	78	66	89 1	100	100	112 1	120 1	122 1	130	148	156

Country data for Africa: case notification rates (per 100 000 population), 1980-2003

116 59 87 77 227 20 40 27 27 63 63 189 121 88 88 39 50 101 69 123 120 **60** 37 48 38 94 170 21 22 42 42 
 Rate (per 100 000 population)

 1997
 1998
 1999

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 61 12 12 154 154 17 17 61 17 128 128 80 68 25 104 104 104 104 104 107 118 184 184 70 15 15 232 232 232 233 234 100 100 45 21 35 35 35 34 10 10 10 10 10 17 17 69 69 69 69 47 2003 8 549 18 971 1 703 3 050 1 703 3 017 10 692 1 65 2 818 3 599 63 3 3 477 11 411 887 39 698 1 233 1 040 7 714 4 495 963 38 158 3 652 99 16 138 5 004 4 505 28 173 3 710 646 36 541 1 033 1 035 7 732 4 300 899 34 337 3 167 2002 8 8 246 18 087 2 415 3 334 1 544 2 791 7 921 111 111 2 758 3 519 3 519 4 207 9 667 44 518 86 15 236 4 535 4 319 10 920 42 054 3 057 1 522 3 040 4 695 1 382 87 4 218 8 497 36 123 7 316 3 920 526 28 773 3 041 2 725 2 920 112 2 222 10 047 34 923 527 21 597 916 861 6 877 3 563 704 27 197 2 729 72 098 1 781 904 18 463 24 125 11 645 8 132 2 690 2 051 122 12 825 3 751 2 631 15 903 4 298 Number of cases 14 492 109 12 116 3 593 2 189 13 161 4 417 7 462 7 462 7 333 1 1988 3 112 3 112 2 782 2 782 4 374 1 04 Country data for Africa: new smear-positive cases, 1993-2003 7 740 8 246 1 939 2 824 1 126 2 022 2 022 3 548 1 03 7 587 3 178 2 519 112 11 116 3 223 1 970 2 820 1996 6 556 8 016 8 016 1 1868 2 530 1 133 2 312 1 17 1 1992 8 70 1 17 2 505 2 24 125 2 29 13 160 263 263 6 474 2 844 2 844 1 788 1 788 6 6 8 8 8 4 5 6 6 7 0 3 2 17 3 9 040 486 778 2 263 2 263 9 56 1 361 1 154 1 8 026 6 285 6 285 1 8 66 2 074 1 1361 1 1492 1 1492 1 1840 1 1840 1 454 23 112 660 887 13 631 19 955 10 038 8 965 11 324 1 330 1994 6 793 4 337 1 618 1 668 561 1 527 1 883 5 752 395 9 677 7 012 10 149 1 405 1 547 6 881 5 692 545 11 949 15 569 9 526 463 4 874 1 653 1 508 5 331 2 316 Rwanda Sao Tome & Principe R Congo quatorial Guinea 3urkina Faso Mauritius Mozambique Namibia Seychelles Sierra Leone South Africa te d'Ivoire auritania

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324 648

121 005

107 012

## **Notes**

#### **Cape Verde**

The NTP notes that age and sex data are incomplete.

#### **Ethiopia**

The NTP notes that treatment outcome data are provisional.

#### Gabon

Selected data received in nonstandard format. DOTS coverage was assumed to be same as in the previous report.

#### Kenya

The treatment outcome cohort excludes nomadic patients (roughly 10% of patients registered) who are not evaluated at the end of their treatment regimen.

#### Madagascar

Data are considered preliminary (83% of expected quarterly reports received). Treatment success for a subset of 11 214 patients on shortcourse chemotherapy was 74% (with 15% defaulting). Treatment success for a subset of 69 patients on longcourse chemotherapy was 67% (with 23% defaulting).

#### Mali

Some 100 patients classified and registered as new smear-positive were excluded from the cohort analysis of treatment outcomes after a chart audit revealed that the initial smear result for these patients could not be identified.

#### Mozambique

Age and sex data are not available according to the categories requested. Among smear-positive cases in 2003, 1.9% were in children. Aside from DOTS coverage, an estimate of "access" to TB services is reportedly 57%.

#### **Seychelles**

DOTS coverage was not reported, but assumed to be same as in the previous report.

#### **South Africa**

Data on age and sex of new smearpositive cases are available as follows:

Age (years)	Male	Female
0-14	38	36
15-19	115	132
20-39	1 143	837
40-59	549	359
60+	81	49

#### **Swaziland**

The NTP notes that age and sex data are incomplete.

#### **Uganda**

The NTP notes that age and sex data are incomplete.

#### **United Republic of Tanzania**

Aside from DOTS coverage, an estimate of "access" is reported in terms of distance from health facility: 70% of the population lives with 5 km from a health unit, and 90% lives within 10 km from a health unit.

**AFR** 

## **Africa**

## **The Americas**

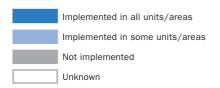
**Eastern Mediterranean** 

**Europe** 

**South-East Asia** 

**Western Pacific** 

#### THE AMERICAS: SUMMARY OF TB CONTROL POLICIES MONITORING OUTCOME<sup>f</sup> **STATUS**<sup>a</sup> MANUAL MICROSCOPY SCCd DOT ANGUILLA NON-DOTS YES ANTIGUA AND BARBUDA DOTS NO ARGENTINA DOTS YES BAHAMAS DOTS YES BARBADOS BELIZE DOTS YES BERMUDA BOLIVIA DOTS YES BRAZIL DOTS YES BRITISH VIRGIN ISLANDS NON-DOTS NO CANADA DOTS YES CAYMAN ISLANDS DOTS YES YES CHILE DOTS COLOMBIA DOTS YES COSTA RICA DOTS YES CUBA DOTS DOMINICA DOMINICAN REPUBLIC DOTS YES YES **ECUADOR** DOTS DOTS YES EL SALVADOR GRENADA NON-DOTS YES GUATEMALA DOTS YES GUYANA DOTS YES HAITI DOTS YES HONDURAS DOTS YES JAMAICA DOTS NO YES MEXICO DOTS MONTSERRAT DOTS YES NETHERLANDS ANTILLES NICARAGUA DOTS YES DOTS YES PANAMA PARAGUAY DOTS YES PERU DOTS YES PUERTO RICO DOTS YES SAINT KITTS AND NEVIS DOTS YES SAINT LUCIA DOTS YES ST VINCENT & GRENADINES DOTS YES SURINAME NON-DOTS NO TURKS & CAICOS ISLANDS NON-DOTS NO TRINIDAD AND TOBAGO NON-DOTS YES URUGUAY DOTS YES



US VIRGIN ISLANDS

USA

VENEZUELA

- a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)
- b Manual: national TB control manual (recommended)

YES

YES

DOTS

DOTS

- c Microscopy: use of smear microscopy for diagnosis (core component of DOTS)
- d SCC: short course chemotherapy (core component of DOTS)
- e DOT: directly observed treatment (core component of DOTS)
- f Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)

**AMR** 

Country data for Americas: estimated burden of TB

data juliante data de la company data de la company de la		poidono 1990	1000	;  -	!	Droviolopoo 1000	1000			Dooth 4000	000		000000000000000000000000000000000000000	2003			Drougloppo	2002	-	2	2003		Γ
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	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number rate	$\vdash$	rate	number	rate	number rate		number rate	edmun	ber rate	<u>a</u>
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Bolivia	19 334	290	8 692	130	30 266	454	30 251	454		42	2 812	42	19 849 225	00		26 548	301	26 508 301	1	696	2	945 33	33
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								Whole country							_		DOTS	S.				non-DOI 3		
						New cases	Se.			Re-treatment cases	cases		Other	Detection rate		,o	Notifications	tions		% of		Notifications	% of	<b>-</b>
	Pop	All	All cases	   	Smear-positive	Pulm Sm	ar-negative	!	Relapse After	After failure After default		Other	₹	cases New ss+	wss+ of		All cases	New ss+	+ DDR	R pulm	■	New ss+	mlnd .	_
	thousands	Country	WHO	ate	number rate	confirmed	or unknown p	pulmonary	nımber	nımber	nımber	nimber	nimber	%	%	non	her	nimber	rate %	+88 8985	s+	number	+ss sessor	9
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Bermuda	82																							
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Paraguay	5 878	2 116					712	225	72	2			119					337						92
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See Explanatory notes, page 151.

Country data for the Americas: treatment outcomes for cases registered in 2002

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Afrilles  4 681 4 680 239 83 2 7 1 6 1 6 6 4 78 1507 1507 1507 1507 1507 1507 1507 1507						15	2	7
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S Antilles         1 1066         1 2 86         99 76         1 4         2 6         97 6         1 4         2 6         1 4         2 6         1 4         2 6         1 4         2 6         1 4         2 6         1 4         2 6         1 4         2 6         1 7         4 4         2 6         1 7         4 4         2 6         1 7         4 4         2 6         1 7         4 4         2 6         1 7         4 4         4 6         4 6         4 89         427         87         8 1         4 4         2 6         1 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8         5 7         4 8	2	9	55 283	63 5	4 3		8	69
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3015 98 82 5 0 10 4 82 381 368 97 79 2 0 16 2	2 0 16	2 7	79 222	78	6 1	13	2	78
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Country data for the Americas: re-treatment outcomes for cases registered in 2002

See Explanatory notes for previous table, page 151

Country data for the Americas: trends in DOTS treatment success and detection rates, 1994–2003	Americas	s: trend	s in DO	TS trea	tment su	s secon	and dete	ction ra	tes, 199	1-2003								
		0	TS new s	smear-po	DOTS new smear-positive treatment success (%)	atment su	%) ssecon				DOT	S new S	near-pos	itive cas	e detection	DOTS new smear-positive case detection rate (%)	(9)	
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
Anguilla					ğ	G G	7	6	0					7	6	Ţ	ç	46
Argentina					22 22	20 20	54	94	28			4	7	74	3 8	‡ %	e 2	65 5
Bahamas					72	99		64	59					62	96		26	52
Barbados																39	8	
Belize						88	78	99	85		100	29			80	95	118	86
Bermuda	99	C	74	1	63	77	70	00	0	0	70	7.5	76	76	7.0	75	76	7
Brazil	99	70	-		2 6	t 6	73	67	75	<del>1</del>	0	2	5 4	5 4	, «c	<u>,</u> «	9 4	- 42
British Virgin Islands						3		5	)				-	-	,		2	2
Canada					32	6/	80	29	8					45	09	61	26	9/
Cayman Islands								100								126		
Chile	83	62	80	77	83	83	82	83	86	70	9/	84	91	95	88	102	117	115
Colombia					74	82	80	85	84					28	83		∞	7
Costa Rica						81	92	72	85					31	120	87	9/	117
Cuba	98	06	92	06	94	91	93	93	92	81	88	87	92	96	86	88	91	93
Dominica			100					100				84	51				36	
Dominican Republic						8	79	82	78					6	9	6	41	92
Ecuador								82	84							2	30	37
El Salvador					77	78	79	88	88			42	52	26	26	28	28	23
Grenada																		
Guatemala	62	61	81	73	62	8	98	82	84	42	26	22	99	26	51	41	46	4
Guyana						91	91	06	82						9	50	10	31
Haiti				73	79	20	73	75	78			2	12	24	22	30	40	46
Honduras					93	88	88	98	87				2	15	62	107	117	28
Jamaica		29	72	79	88	74	42	78	49		95	92	91	102	101	8	29	6
Mexico			75	65	78	80	92	83	84			15	30	39	89	91	20	84
Montserrat Netherlands Antilles																		069
Nicaragua	81	80	79	8	82	81	82	83	82	72	83	84	98	82	84	6	82	91
Panama				51	51	80	29	65	73				13	80	43	99	8	92
Paraguay	46	21					77	98	95	4	22				4	6	00	18
Peru	81	83	88	06	92	93	06	06	92	101	88	94	86	91	87	87	98	81
Puerto Rico		65	89	89	89	77	72	80	62		29	73	99	72	61	22	99	29
Saint Kitts & Nevis					25	20							174	88			48	
Saint Lucia				29	82	88	100	20	25			118	82	77	09	52	20	71
St Vincent & Grenadines				98		100	100	80					18		26	19		38
Suriname																		
Trinidad & Tobago																		
Turks & Caicos Islands	;				71					i	,		į	116		i	i	1
Uruguay US Virgin Islands	83	20	80	1	2	83	82	82	82	92	93 75	94	84	88	80	79	72	80
USA	0	72	71	72	72	76	82	83	70	1	84	82	84	84	83	28 2	98	88
Venezuela	20	14	8	7/	8	78	9/	200	82	/3	(2)	(2)	8/	82	2	/9	62	28
Region	77	77	8	8	8	83	81	83	8	23	28	30	34	37	4	43	46	20

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Barbuda         69-14         15-24         26-34         35-44         45-54         55           Barbuda         89         574         565         413         461         6           1         2         5         6         3         461         3           1         4         8         10         3         461         3           1         4         8         10         3         461         3           In Islands         3         45         58         50         52         1076           In Islands         3         45         58         59         52         1076         1076           In Islands         3         45         58         59         52         1076         10						l					ALL			
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156   164   742   501   438     18														
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49     309     429     290     173       18     310     266     194     125       7     75     105     103     81       29     175     200     169     156       66     769     774     470     314       66     769     774     470     314       10     117     49     1387     1272       10     174     1429     1387     1272       14     179     210     135     60       6     72     93     73     60       7     50     31     31     34       101     758     506     355     206       0     0     0     1     2       0     3     5     8     10       0     0     0     1     2       0     0     0     1     2       13     36     50     35     42       13     360     518     750     821		62	0 11	14	20	23	13	30	2	34 104	111		92	109
49         3109         423         290         173           7         75         105         103         81           7         75         105         109         156           0         13         28         36         166           0         13         28         36         16           66         769         774         470         314           66         769         774         470         314           10         171         14         122         227           10         174         1429         1387         1272           0         0         1         0         0           0         1         0         0         0           0         3         5         8         10           0         3         5         8         10           0         3         5         8         10           13         360         518         750         821					0	ć	ç			020			2	7
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29 175 200 169 156 66 769 774 470 116 66 769 774 470 314 11 1 9 14 172 180 1174 1429 1387 1272 0 0 0 1 13 36 518 750 821		0 0		71	47	8 8	4	2 2	14	145 176	150		103	153
29         175         200         169         156           0         13         28         36         16           66         789         774         470         314           52         20         344         235         227           180         1174         1429         1387         127           0         0         0         0         0         0           14         179         210         135         103         60           6         72         93         73         60         6         7         60         10         0		3				2	;						2	3
0         13         28         36         16           66         769         774         470         314           52         20         344         470         314           1         11         14         174         1429         1387         1272           1         10         0         1         0         0         0         0         0           1         179         210         135         103         60         0         0         0         0         0           6         72         93         73         60         34         34         60         36         506         365         206         306         306         365         206         306         306         306         306         366         306 <td></td> <td>128</td> <td></td> <td></td> <td>157</td> <td>104</td> <td>88</td> <td>75</td> <td>.,</td> <td>361 379</td> <td></td> <td></td> <td>213</td> <td>203</td>		128			157	104	88	75	.,	361 379			213	203
66         769         774         470         314           52         20         344         235         227           1         10         14         122         127           180         1174         1429         1387         1272           0         0         1         0         0           14         179         210         135         103           6         72         93         73         60           7         50         31         31         34           101         758         506         355         206           0         3         5         8         10           0         0         0         1         2           3         46         50         35         42           13         360         518         750         821		9		13	19	6	9	2	,		1 55		17	00
52         20         344         235         227           1         11         9         134         125           180         1174         1429         1387         127           0         0         1         0         0           14         179         210         135         103           6         72         93         73         60           7         50         31         31         34           101         758         506         355         206           0         3         5         8         10           0         0         0         1         2           3         46         50         35         42           13         360         518         750         821		114	101 859		440	289	135	77	_	_			297	191
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4   179   210   153   103     6   7   2   31   31   34     101   788   506   885   206     0   0   0   1   2     0   0   0   1   2     3   46   50   35   42     13   360   518   750   821   4		u d		4	5	7	7			000			5	4
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Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

Country data for the Americas: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers)

	6	,		L										<u> </u>							
	0-14	15-24	25-34	35-44	45-54	55-64	65+	0-14	15-24	25-34	35-44	45-54	55-64	65+	0-14	15-24	25-34	35-44	45-54	55-64	65+
Anguilla Antigua & Barbuda Argentina																					
Bahamas Barbados Belize																					
Bermuda Bolivia Brazil	264	3 511	4 491	4 712	3 787	1 954	1 470	292	2 838	2 745	2 004	1 316	729	711	556	6 349	7 236	6 716	5 103	2 683	2 181
British Virgin Islands Canada Cayman Islands																					
Chile Colombia Costa Rica	224	611	727	783	779	595 4	683	158	265	625	464	349	257	365	382	1 208	1 352	1 247	1 128	852 4	1 048
Cuba Dominica Dominican Republic	ဇ	55	88	41	21	16	16	0	36	38	25	23	14	16	က	91	127	99	4	30	32
Ecuador El Salvador Grenada																					
Guatemala Guyana Haiti	70	43	83	78	42	16	7 23	10	31	48	37	18	4 %	es 25	20	74	131	115	9 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 50	10
Honduras Jamaica Mexico	2	3 8	32	30	17	2 8	3 25	07	02	5 5	2	3 45	25	5 %	. 4	53 53	47	500		2 2	080
Montserrat Netherlands Antilles																					
Panama	4 4	19	27	20	16	16	15	w *	2 5	18	12	9 7	12	ر د د	7	21	45	32	22	28	50
ralaguay Peru	1	2	5	0	0	98	6	<u>+</u>	6	0	c c	ţ	9	5	0	2	00	2	27	06	9
Puerto Rico Saint Kitts & Nevis																					
Saint Lucia St Vincent & Grenadines																					
Suriname	0	2	0	10	2	-	2	-	က	2	-	_	0	_	_	80	2	7	9	~	9
Trinidad & Tobago	0	6	13	10	13	10	9	-	2	2	0	80	-	2	-	=	15	10	21	=	80
Turks & Calcos Islands Uruguay	0	0	7	0	0	0	0	0	0	7	0	8	0	0	0	0	4	0	N	0	0
USA Venezijela		α	Ľ		+	C	c	,	Ę	α	u	ď	+	c	~	ά.	4	12	7	-	
Region	541	4 642	5 820	5 924	4 888	2 746	2 359	511	3 851	3 824	2 787	1 867	1 103	1 205	1 052	8 493	9 644	8 711	6 755	3 849	3 564

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

**AMR** 

5 5 72 16 39 17 6 41 18 31 46 98 100 109 3 32 65 49 47 47 4 53 16 17 17 15 30 1 123 33 55-64 45-54 37 37 15 17 29 36 13 29 17 16 40 26 35 127 145 104 7 7 15 3 17 21 35-44 11 22 9 2 6 8 2 15 48 18 22 29 165 163 65 5 17 17 47 34 28 28 18 2 7 2 ALL 25-34 c 1 0 5 5 12 1 19 28 15 2 2 8 27 99 31 22 124 173 56 4 4 37 29 20 15-24 16 16 2 2 38 o 4 15 8 116 25 27 27 27 27 4 0-14 10 Country data for the Americas: smear-positive notification rates (per 100 000 population) by age and sex, 2003 9 10 2 113 50 27 50 13 22 7 36 **65**+ 92 140 18 34 11 32 23 60 17 2 29 23 7 12 7 106 86 25 3 37 48 81 81 94 0 26 24 33 44 45 0 2 19 45-54 3 7 5 10 18 24 21 21 0 9 1 19 24 65 18 27 75 128 105 3 17 Ξ 0 2 Ξ FEMALE 35-44 13 , 16 7 3 2 38 8 8 8 5 2 2 2 - 5 23 23 4 4 22 25-34 15 <u>ο</u> 6 α 5 5 5 10 15 8 21 86 177 45 45 37 29 17 18 2 8 84 24 15-24 16 ر م 17 11 15 46 115 30 18 15 25 7 4 22 97 0-14 10 0 0 % + 0 0 0 0 0 , 0 2 24 **65**+ 39 147 252 48 27 79 30 14 31 59 60 79 100 7 7 57 26 100 26 5 59 46 15 15 58 1 25 27 57 25 10 55-64 8 \$ 23 8 55 161 123 125 6 39 76 75 68 7 56 11 22 27 24 164 51 4 4 63 51 18 5 45-54 24 21 43 10 10 51 23 35 43 165 103 11 31 35 24 138 56 35 37 23 34 111 31 241 173 66 9 18 15 29 10 57 24 33 53 33 33 33 11 25 32 11 17 3 27 23 35-44 73 25-34 19 39 39 က 12 23 25 25 19 0 5 23 163 168 66 4 4 2 4 4 5 20 33 33 22 13 72 103 15-24 15 2 9 9 8 23 45 7 17 2 4 16 136 31 27 28 0 0 0 0 0 0 % 0 0 0 Vincent & Grenadines Frinidad & Tobago Turks & Caicos Islands Sritish Virgin Islands ominican Republic Jetherlands Antilles Puerto Rico Saint Kitts & Nevis Antigua & Barbuda JS Virgin Islands Sayman Islands El Salvador Sosta Rica Suatemala **Jontserrat** Saint Lucia icaragua Solombia **3ahamas 3arbados** Dominica **Srenada** londuras araguay ermuda cuador amaica anama Jruguay Suvana Region **∂**exico

33 7 119 31

**65**+

8 2 2 o

8 8 4

45 78 78 4 42

49 13 3

92 42 9

3 16

Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available

0 2 226 11 640 527 844 2 642 631 14 004 3 102 120 17 078 2 283 1 585 2 175 2 175 31273 1 147 1 147 6 643 14 861 6 734 4 696 6 442 1 383 227 2 909 590 12 066 4 579 106 17 790 2 092 1 514 2 107 36 092 3 129 1 129 9 3 17 1 133 5 3 6 4 040 5 829 1 550 3 006 11 480 630 929 2 419 422 10 224 4 435 121 18 879 **229 874** 40 91 2001 2 913 422 10 420 3 984 127 18 434 16 362 6 466 236 183 38 86 3 021 11 630 585 1 135 5 291 5 908 1 485 2000 2 820 407 9 124 4 568 115 1999 1 806 2 3 429 10 999 851 1 135 240 650 39 89 5 767 5 756 1 623 17 521 6 598 2 755 318 9 770 4 916 121 21 514 1998 3 3 652 3 455 9 155 7 30 1 234 5 114 7 164 1 700 2 604 1 422 1 831 1 831 201 20 20 8 85 199 668 18 199 6 273 2 948 407 10 116 4 030 118 23 575 2 806 1 473 1 946 42 062 257 22 22 22 22 22 22 22 22 22 76 76 76 76 76 76 76 76 77 76 1997 0 3 880 8 042 692 1 346 6 5 381 9 435 1 662 3 003 1 314 2 072 41 739 110 3 35 6 6 6 53 21 119 5 650 256 467 39 89 3 232 314 6 632 4 176 121 20 722 1 868 0 0 0 9 702 636 1 465 10 6 302 8 397 1 686 1996 1 931 2 2 2 4 150 9 912 586 1 553 8 8 8 8 8 7 893 2 422 3 95 4 14 422 91 013 3 119 296 6 212 4 984 109 11 329 22 860 5 578 258 331 39 89 2750 827 1850 1850 274 274 274 274 129 129 166 666 666 8677 4 291 109 16 353 13 683 78 59 9 431 75 759 1994 2 074 4 138 8 901 325 1 681 1 2 4 337 9 685 3 901 2 508 266 13 887 60 2 012 2 4 598 11 043 313 790 7 4 033 7 050 3 347 3 745 115 25 287 5 169 1993 2 798 1 146 2 037 51 675 6 2 108 3 3 3 11 199 11 199 410 410 13 3 490 2 495 2 495 4 155 111 14 446 3 5 498 12 263 201 514 14 1 837 6 879 2 304 10 237 134 10 237 4 560 121 15 216 1 997 2 6 151 12 447 230 546 6 2 597 8 243 2 367 3 647 123 14 437 2 944 846 2 167 37 905 159 0 13 82 120 0 120 0 888 12 636 52 52 5 30 2 12 563 80 048 2 035 2 6 728 111 329 311 581 13 3 145 5 480 617 4 900 120 8 100 4 026 8 86 15 489 3 106 672 2 270 35 687 314 0 0 28 70 124 987 0 5 739 150 8 054 3 962 65 15 371 13 267 51 51 28 28 10 664 82 395 1 947 0 6 324 11 469 442 628 7 7 3 081 5 497 2 378 **241 834** 41 93 1 972 0 6 280 111 437 434 630 27 27 2 459 5 867 1 647 5 700 117 8 514 4 227 1 133 14 631 Country data for the Americas: number of TB cases notified, 1980–2003 14 681 52 7 7 23 6 6 837 83 731 2 046 6 854 111 639 111 639 656 35 2 634 5 687 1 659 4 806 8 583 4 213 88 13 180 6570 215 4959 3377 130 6 644 12 024 376 680 8 2 335 4 798 1 461 6 586 165 5 803 2 120 160 14 531 1 6 561 12 792 393 705 5 3 100 4 301 1 564 16 359 53 53 14 14 35 35 4 131 88 365 6 013 149 6 839 1 935 157 22 795 2773 429 1 800 22 753 2 452 4 48 4 48 4 48 1 112 5 5 1 359 4 2 846 4 4 266 2 515 0 0 6 941 12 126 459 815 18 2 457 3 880 2 171 7 277 135 3 337 1 714 153 24 853 2 554 2 7 337 11 483 521 833 26 1 778 3 966 2 091 6 641 117 6 550 1 696 178 32 572 3 723 580 1 388 21 925 521 16 406 70 64 21 2 885 0 0 8 523 11 589 1 1589 1 133 2 174 1 2 255 1 2 255 1 2 255 1 1 3 3 950 1 1 7 5 624 1 674 1 674 1 1674 1 1674 1 1674 1 1674 1 4 412 72 608 Suriname Trinidad & Tobago Turks & Caicos Islands aint Lucia t Vincent & Grenadir fontserrat letherlands Antilles Puerto Rico Saint Kitts & Nevis r reporting t reporting Virgin Islands ayman Islands Oominica Oominican F cuador

	1980	1981	1982	1983	1984	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002
Anguilla	0		57	0	0		0	0	0	0			0			20		0				0	0
Antigua & Barbuda	13		0		5		=	0	2	2			6			0	4	9	9	4	9	-	9
Argentina	58		09		22		48	43	42	39			38	41	40	39	38	35	34	32	32	31	30
Bahamas	33	31	25	26	23	27	22	18	21	21	18	20	24	22	28	20	20	30	25	25	27		14
Barbados	26		12		9		က	_	2	7			7			_	~	2	က	<del>-</del>	_	5	7
Belize	15		29		22		14	24	16	17			33	40	28	45	45	48	24	4	44	22	24
Bermuda	_		7		4		80	က	-	3			5			5	0	5	0	0	0	0	0
Bolivia	82		85		71	`	112	144	167	193	`	•	136	120	129	193	133	126	127	121	122	124	118
Brazil	09		69	29	99		09	28	22	22			26		48	22	24	20	22	47	45	43	46
British Virgin Islands																		16				0	5
Canada	12		10	6	6		80	7	7	7			7	7	7	7	9	7	9	9	9	2	2
Cayman Islands	0		0	2			2	0	0	8			11	7		9	0	0	6	9	14	3	0
Chile	9/		09	09			26	20	20	52			39	33	30	59	29	27	25	23	20	19	16
Colombia	4		41	45	4		36	35	34	33			31	30	24	56	25	20	22	27	28	27	26
Costa Rica	17		18	19			15	15	15	10			4	6	10	17	18	19	19	22	15	16	13
Cuba	12		8			7	9	9	9	9			4	7	15	14	13	12	11	10	10	8	8
Dominica	27	35	24	22		7	48	37	10	18	00	19	18	10	16	7	13	00	7				ო
Dominican Republic	38		41				40	37	45	45			48	24	22	23	81	99	63	20	63	26	47
Ecuador	20		46	46			61	61	26	22			99	64	87	69	72	80	09	47	26	48	46
El Salvador	49		46	44	33		8	34	48	12			47	62	20	43	29	28	28	27	24	23	24
Grenada	19		_				-	2	0	2			4	0	4	2	0	2	2	9	0		-
Guatemala	82	92	101	82	87		61	70	69	22			27	26	56	31	32	28	25	22	26	21	24
Guyana	16		18				25	16	20	16			25	12	36	40	42	24	42	72	26	22	77
Haiti	152		28				136	132	122	120						83	87	131	125	115	130	126	147
Honduras	47	46	45	49			86	92	98	82	75		80	20	78	88	72	89	80	73	62	29	89
Jamaica	8		7	7			4	9	က	4			2	2	4	4	2	2	2	4	2	2	4
Mexico	46		35	32	20		17	19	19	19			17	17	18	12	22	25	22	20	19	19	17
Montserrat	∞	0	0	6			45	118	22	46			0		0				16	41	0	0	0
Netherlands Antilles																						7	က
Nicaragua	45	`	66	86			75	84	75	83	77		71	29	2	25	99	09	54	25	47	47	39
Panama	33		28	21			35	34	33	28	32		30	45	35	49	48	23	20	48	40	22	49
Paraguay	43	43	43	53	49	24	4	39	36	22	21	23	43	4	36	36	45	38	32	9	36	37	37
Peru	92	`	119	122		`	124	150	1//	167	1/4		233	225	208	190	1/2	1/0	1/4	158	149	141	135
Puerto Rico	21		4 :	4 1	12		Ξ,	<b>ກ</b> (	00 (	თ (	2			<b>-</b> :	00 1	<b>-</b> :	က၊	- :	c i	2	2	က၊	က၊
Saint Kitts & Nevis	16		14	Ω :	<b>-</b> !		0 (	0 ;	O !	0	) )		10	4	υ į	Ξ'	- 1	87 !	75	- ;	0 (	٠ <u>:</u>	- :
Saint Lucia	36		32		46		78	20	25	22	10		19		14	80	25	15	14	=	9	10	11
St Vincent & Grenadines	78		4		22		თ	က	9	က	2		4	12	0	=	2	2	7	∞	4	∞	∞
Suriname	22	23	15	21	20		15	20	19	18	8	12	4	=	13		13	9	20	23	21	19	22
Trinidad & Tobago	7		9		6		10	10	6	10	10		12	6	10	13	16	20	16	12	15	16	10
Turks & Caicos Islands	27	0	24	28	0		20	113			0		0	0						92		16	12
Uruguay	64		49	46	46	40	36	34	31	32	29		22	22	21	19	22	22	20	19	19	20	16
US Virgin Islands	0	-	-	2	3		-	2	9	4	4				10	4	80						
USA	12	12	7	10	6	0	6	თ	თ	თ	10		10	10	တ	∞	00	9	_	9	9	9	2
Venezuela	28	56	26	26	28		28	27	25	24	78		27	25	23	22	25	26	27	78	27	25	25

2 4 15 2 4 15 41 69 83 15 15 2001 15 40 35 4 4 4 12 4 4 4 12 14 14 32 15 20 96 15 16 Rate (per 100 000 population) 1997 1998 1999 15 49 16 110 83 0 2 0 1 2 5 6 6 2 4 9 5 6 4 3 28 28 1 2 2 2 38 29 19 145 13 19 94 25 1994 8 1 4 2 4 2 1 8 9 4 25 4 41 21 155 244 7 015 1 956 81 1 404 788 1 166 18 504 62 5 303 3 882 0 592 0 1 276 7 972 346 511 2 806 1 488 870 125 803 1 865 138 6 188 2 956 60 11 555 1 320 709 1 004 20 533 5 380 3 444 445 0 1 412 7 787 328 538 2 179 1 223 980 127 357 1 669 174 5 607 2 839 75 15 103 1510 575 915 21 685 5 600 3 476 1 355 8 022 385 562 2 622 4 439 1 003 129 536 1471 410 900 22 580 82 5 865 3 525 506 1 290 8 358 349 677 130 251 432 1 041 24 511 106 6 252 3 670 395 2 1 1 497 8 329 458 720 3 278 4 300 1 023 2 264 178 6 828 2 415 135 068 Country data for the Americas: new smear-positive cases, 1993-2003 Number of cases 1 648 1 393 850 27 707 106 6 630 3 450 471 2 1 576 6 969 353 746 139 286 1998 1 582 6 090 320 765 6 882 3 234 3 162 7 214 882 142 512 3 46 6 949 44 503 5 787 3 733 6 426 965 1 722 904 894 26 800 110 7 401 3 195 156 0 1 562 7 572 302 835 426 136 657 1 568 1 066 748 32 096 128 2 306 93 9 220 8 013 3 056 404 0 1 561 7 530 245 834 1995 5 787 5 890 138 820 6 905 1 951 6 532 230 914 8 3 177 6 674 2 144 2 385 61 9 726 1 615 748 873 33 925 14 346 2 738 1994 142 405 1 714 1 046 985 35 646 542 2 2 629 6 987 565 6 2 297 5 325 2 471 2 016 83 8 164 16 046 2 849 5 937 20 2 128 51 104 931 1993 6 833 388 Suriname Trinidad & Tobago Turks & Caicos Islands Vincent & Grenadines ominican Republic uador British Virgin Islands Canada Nontserrat Vetherlands Antilles Puerto Rico Saint Kitts & Nevis JS Virgin Islands Sayman Islands Colombia Costa Rica Salvador aint Lucia icaragua Sarbados araguay ominica renada onduras Suyana Iruguay Region

# **Notes**

#### **Brazil**

The NTP notes that age and sex data are incomplete and that treatment outcome data are provisional.

#### **Ecuador**

Treatment outcomes were available from only 3 provinces.

### Guyana

Age and sex data are for all new cases, not just smear-positive.

#### Peru

Age and sex data are for quarter 3 only.

#### **Uruguay**

Treatment outcomes listed under new smear-positive patients are for laboratory-confirmed cases.

### **USA**

The NPT notes that treatment outcome data are provisional. Treatment outcomes listed under new smearpositive cases are for laboratoryconfirmed cases.

**AMR** 

**Africa** 

**The Americas** 

**Eastern Mediterranean** 

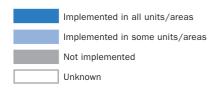
**Europe** 

**South-East Asia** 

**Western Pacific** 

## **EASTERN MEDITERRANEAN: SUMMARY OF TB CONTROL POLICIES**

	STATUS <sup>a</sup>	MANUAL	MICROSCOPY	SCC <sup>d</sup>	DOT®	MONITORING OUTCOME <sup>f</sup>
AFGHANISTAN	DOTS	YES				
BAHRAIN	DOTS	YES				
DJIBOUTI	DOTS	YES				
EGYPT	DOTS	YES				
IRAN	DOTS	YES				
IRAQ	DOTS	YES				
JORDAN	DOTS	YES				
KUWAIT	DOTS	YES				
LEBANON	DOTS	YES				
LIBYAN ARAB JAMAHIRIYA	DOTS	YES				
MOROCCO	DOTS	YES				
OMAN	DOTS	YES				
PAKISTAN	DOTS	YES				
QATAR	DOTS	NO				
SAUDI ARABIA	DOTS	YES				
SOMALIA	DOTS	YES				
SUDAN	DOTS	YES				
SYRIAN ARAB REPUBLIC	DOTS	YES				
TUNISIA	DOTS	YES				
UNITED ARAB EMIRATES	DOTS	YES				
YEMEN	DOTS	YES				
WEST BANK AND GAZA STRIP	DOTS					



- a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)
- b Manual: national TB control manual (recommended)
- c Microscopy: use of smear microscopy for diagnosis (core component of DOTS)
- d SCC: short course chemotherapy (core component of DOTS)
- e DOT: directly observed treatment (core component of DOTS)
- f Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)

**EMR** 

Country data for the Eastern Mediterranean: estimated burden of TB

	_	ncidenc	Incidence, 1990		Pre	Prevalence, 1990	, 1990			Death, 1990	0661	Ī	=	cidenc	Incidence, 2003			revalenc	Prevalence, 2003	r	ă	Death, 2003	003	
	All cases incl.	+ All	All cases incl. HIV+ New ss+ incl. HIV+		All cases incl. HIV+ All cases excl	HIV+ A	Il cases excl.	+AIIA+	All cases incl.	4/I/	HIV+ All cases excl. HIV+	+ AIIV	All cases incl.	± MH	New ss+ incl.	+ AIN	All cases incl HIV+	-	All cases excl HIV+		All cases incl HIV+	HV+	All cases excl	+   
	number	rate	unmber	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate
Afghanistan	45 996	333	20 698	150	95 688	693	95 688	693	10 370	75	10 370	75	79 656	333	35 845	150	160 303	671	160 302	671	22 125	93	22 123	93
Bahrain	332	89	149	30	691	141	691	141	75	15	75	15	330	46	148	20	374	52	373	52	33	2	33	2
Djibouti	3 161	599	1 395	264	7 060 1 338	338	7 028 1	1 332	845	160	811	154	5 152	733	2 2 7 3	324	7 166	1 020	6 939	988	805	115	069	86
Egypt	23 323	42	10 492	19	48 521	87	48 521	87	5 259	6	5 259	6	20 233	28	9 102	13	26 148	36	26 133	36	2 350	က	2 343	က
Iran	23 179	4	10 424	18	48 221	82	48 221	85	5 226	6	5 226	6	19 423	28	8 735	13	25 165	37	25 137	36	2 259	က	2 246	က
Iraq	15 983	92	7 192	14	33 250	192	33 250	192	3 604	21	3 604	21	39 552	157	17 797	71	59 4 19	236	59 411	236	8 298	33	8 293	33
Jordan	408	13	184	9	849	56	849	56	92	က	92	က	270	2	121	2	283	2	283	2	27	0.0	27	0.0
Kuwait	848	40	381	18	1 764	82	1 764	82	191	6	191	6	671	27	302	12	774	31	773	31	89	က	29	က
Lebanon	1 158	43	521	19	2 409	68	2 409	89	261	10	261	10	448	12	201	9	492	13	491	13	45	_	45	_
Libyan Arab Jamahiriya	1 333	31	669	14	2772	49	2772	64	300	7	300	7	1 156	21	519	6	11151	21	1 145	21	8	2	83	-
Morocco	32 960	134	14 822	09	68 269	279	68 269	279	7 431	30	7 431	30	34 327	112	15 437	51	32 211	105	32 159	105	3 099	10	3 084	10
Oman	316	17	142	00	657	36	657	36	71	4	71	4	307	7	138	2	350	12	349	12	8	_	34	_
Pakistan	201 031	181	888 06	82	418 218	377	418218	377	45 325	41	45 325	41	278 392	181	125 172	82	550 948	329	550 424	358	66 503	43	66 037	43
Qatar	248	53	112	24	516	110	516	110	26	12	26	12	372	61	167	27	440	72	439	72	41	7	40	7
Saudi Arabia	9 065	22	4 079	25	18 858	114	18 858	114	2 044	12	2 044	12	9 751	40	4 387	18	13 720	22	13 718	22	1 132	2	1 131	5
Somalia	24 042	336	10 738	150	53 815	751	53 598	748	6 408	89	6 187	98	40 645	411	18 154	184	74 655	755	73 972	748	12 468	126	11 695	118
Sudan	44 706	179	19 756	79	100 764	404	100 558	403	11 818	47	11 607	47	73 802	220	32 614	26	122 277	364	119 285	355	20 960	62	18 067	25
Syrian Arab Republic	8 673	89	3 903	31	18 043	142	18 043	142	1 955	15	1 955	15	7 551	42	3 3 3 8	19	9 292	25	9 290	25	814	2	813	2
Tunisia	3 122	38	1 405	17	6 495	79	6 495	79	704	6	704	6	2 170	22	926	10	2 374	24	2 373	24	227	7	227	2
United Arab Emirates	538	56	242	12	1119	22	1119	22	121	9	121	9	532	18	239	∞	778	26	776	56	62	7	61	7
West Bank and Gaza Strip	771	36	347	16	1 603	74	1 603	74	174	∞	174	∞	856	54	385	7	1 332	37	1 332	37	150	4	150	4
Yemen	16 431	138	7 386	62	34 182	286	34 182	286	3 705	31	3 705	31	18 514	93	8 323	42	30 299	151	30 257	151	2 353	12	2 332	12
Region	457 623	120	205 352	25	964 066	253	963 610	253	106 036	28	105 570	28	634 112	122	284 434	22	1119950	216	1115362	215	143 937	28	139 623	27
Legion	431 023	77	700 007	ŧ	304 000	232	200 0 10	700	000 001	07	0.00 CO	707	211 400	77	404 407	2	0000	710	2	205		213 143	143 337	213 145 95/ 20 159

See Explanatory notes, page 151.

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							1	A - I - I										2	F			ŀ		O.F.O.	Γ
							\$	whole country	2									3	200				2	non-DOI 3	
						New	New cases			Re-	Re-treatment cases	S	Other		Detection rate	%		Notific	Notifications		-	% of	Notifications	ons	yo %
	Pop	A	All cases	Š	Smear-positive	Pulm	Smear-negative	ve Extra-	- Relapse		After failure After default	ault Other		¥ا ا	All cases New ss+	Г		All cases	New ss+		DDR	/ mlnd	All cases 1	New ss+	mlnd
	l	Country	WHO	l		confirmed	d or unknown	n pulmonary	ary																
	thousands	number	number ra	rate n	number rate	number	r number	r number	ber number		number number	iber number	ber number	per	%	d %	pop number	nber rate	number	rate	% cas	cases ss+	number	number cases ss+	ises ss+
Afghanistan	23 897		13 808	58	6 510 27	6 510	0 3 440		3 254 6	604	141						53 13	13 808 58	6 510	27	18	65			
Bahrain	724	263	263	36	73 10		99		124						80	49	100	263 36	73	10	49	23			
Djibouti	703		3 231 4	460	1 202 171		828		1 186	184	38	23				53 1	100 3	3 231 460	1 202	171	53	28			
Egypt	71 931	11 747	11 490	16	5 118 7	5 118	8 2 764		3 045 5	163	144	113			57		100 11	11 490 16	5 118	7	26	65			
Iran	68 920	10 857	10 857	16	5 166 7	5 166			3006	319	113		,-	113	56	59 1	100 10	10 857 16	5 166	7	29	69			
Iraq	25 175	11 656	11 656	46	3 577 14		3 727		3454 8	868					29	_	85 11	11 656 46	3 577	4	20	49			
Jordan	5 473	310	310	9	108 2	119			139	2	2	0		10	115	_	100	310 6	108	2	68	64			
Kuwait	2 521	299	999	22	201 8	201	1 73		288	4		30			84	67 1	100	566 22	201	80	29	73			
Lebanon	3 653	380	380	10	134 4	156			157	6	0	0					100	380 10	134	4	29	63			
Libyan Arab Jamahiriya	5 551	1 559	1 917	35	764 14		351		795	7	2	131					100 1	1917 35	764	14	147	69			
Morocco	30 566	26 789	26 789	88	12 842 42	13 009	9 2 2 7 1	11 676	376						78	83 1	100 26	26 789 88	12 842	45	83	85			
Oman	2 851	257	257	6	112 4	112	2 35		103	7	0	0					100	257 9	112	4	81	92			
Pakistan	153 578	73 130	73 100	48	20 962 14		34 447	7 12874		1 633	39 3	3 145				H		73 100 48	20 962	14	17	38			
Qatar	610		276	45	95 16	276	6 63		117	_					74	57 1	100	276 45	95	16	22	09			
Saudi Arabia	24 217	3 3 1 7	3 317	14	1646 7	1 646	6 546		1 0 1 0 1	115						_	100 3	3 317 14	1 646	7	38	75			
Somalia	068 6	9 2 7 8		94	5 190 52		2 193		1388 5	507		0			23	29 1		9 278 94	5 190	25	59	20			
Sudan	33 610	25 095	25 095	75	10 993 33	20 429	962 2 6		4666 16	640								25 095 75	10 993	33	8	29			
Syrian Arab Republic	17 800	4 966	4 820	27	1545 9	1 545	5 1 026		2173	92	56	35		22	64	45	100 4	4 820 27	1 545	6	45	09			
Tunisia	9 832	1 965	1 965	20	6 068		194		840	41							100 1	965 20	890	6	91	82			
United Arab Emirates	2 995		117	4	77 3	7	77	9	33	_	0	0			22	32 1	100	117 4	77	က	32	93			
West Bank and Gaza Strip	3 557	37	36	<del>-</del>	15 6			+	17						4	4	100	36 1	15	0	4	79			
Yemen	20 010	10 413	10 413	52	3 793 19		3 435		2759 4	426					56	46	9 86	6 632 33	3 602	18	43	73	3 781	191	80
Region	518 063	192 585	209 941	14	81 013 16	54 364	4 65 802	2 52 905	7 0 37	137	538 3	3 477	-	178	33	28	86 206 160	160 40	80 822	16	28	26	3 781	191	œ

Country data for the Eastern Mediterranean: notification, detection and DOTS coverage, 2003

See Explanatory notes, page 151.

Country data for the Eastern Mediterranean: treatment outcomes for cases registered in 2002

Carolin Management			1		1		2		,							144		1				-			141			Ė		
			Z	ew sme	New smear-positive cases - DOLS	ve case	S - DOL	0						New	smear-pc	New smear-positive cases - non-DOLS	u - sast	SIDOI-NO	_					Smear-	positive	re-treath	Smear-positive re-treatment cases - DOLS	es - nor	٥	
			%	%	%	%	%	%	%	%	%			%	%	%	%	%	%	%	% %		%	% %	%	%	%	%	%	%
	Number of cases	of cases	of notif		-ldmoo				trans-	not		Number of cases	of cases	of notif	J	-ldmoo			tra	trans- no	not	Number	Je.	compl	-6			trans-	not	
	notified	regist'd	regist'd	cured	eted	died		failed default ferred	ferred	evals	saccess	notified	regist'd	regist'd	cured	eted	died f	failed de	default fer	ferred ev	eval success	ss regist'd	st'd cured	ed eted	died died	d failed	ed default	It ferred	eval	saccess
Afghanistan	6 208	7 780	120	09	27	4	2	2	ю		87																			
Bahrain	17	17	100	88		12					88																			
Djibouti	1 253	1 256	100	99	16	~	-	15	-		82											2	268 4	43 29	29 2	2	3 21	2		73
Egypt	4 889	4 605	8	9/	12	2	9	3	က		88																			
Iran	5 335	5 366	101	81	4	9	2	က	က	0	85											2	599 6		13 8	8	5 8	2	_	74
Iraq	3 895	3 895	100	98	2	က	2	က	2		91											2	553 6		7 5	5 11	1 8	_		75
Jordan	91	91	100	80	6	2		2			88												12 5	58 17	7 17	7	8			75
Kuwait	_	206		45	13			25	4	16	22																			
Lebanon	148	147	66	80	=		-	7	-		91												<b>—</b>	100	0					100
Libyan Arab Jamahiriya	722	716	66	48	13	-	0	17	18	က	61												14						100	
Morocco	12 914	12 830	66	83	9	2	_	80			88											17	1727 6	69	5 5	5	5 12	2		74
Oman	151	119	79	92		7		~	-		92												1	100						100
Pakistan	15 331	14 314	93	92	13	က	-	14	4	-	77	934										2 871		33 40	43 4	4	2 11	2	2	9/
Qatar	25	49	100	69	9	9			19		75																			
Saudi Arabia	1 674	1 365	82	69	7	7	0	14	2		9/											_	112 5	51 16	16 4		2 13	4	#	29
Somalia	4 8 1 8	4 8 1 8	100	98	3	3	-	3	4		88											4	411 5	¥	4 5	2	3 4	30		29
Sudan	10 338	10 993	106	29	19	က	-	7	2	80	78											16	640 5	51 26		3	0 5	5	12	77
Syrian Arab Republic	1 447	1 447	100	69	18	က	2	7	-		87											_	167 3	35 21		7 17	17 18	2		26
Tunisia	927	913	86	68	က	2	-	2	2		92												53 7	75		2 8	9 11	2		75
United Arab Emirates	22	22	100	70	6	4	2	2	2	12	79												က						100	
West Bank and Gaza Strip		10		100							100												9		7 5	5	5	2		9/
Yemen	3 870	3 790	86	73	6	4	~	6	4		82	389	334	98	28	49	-	-	15	5	78		393 5	50 20	20		20		10	70
Region	74 450	74 799	100	72	12	ო	-	80	က	-	84	1 323	334	25	28	49	-	-	15	2	78	8 825		51 2:	23 4	4	4 10	2	က	74

See Explanatory notes, page 151.

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				Relaps	Relapse - DOTS	LS						1	After failure - DOTS	Ire - DO	TS						After	After default - DOTS	- DOTS			
	number	%	%	%	%	%	%	%	- %	number	%	%	%	%	%	%	% %	number	% it	%	%	%	%	%	%	%
			-ldwoo				trans-	not			J	compl-				trans- n	not			comp				trans-	not	
	regist'd	cured	eted	died	failed	failed default ferred		eval su	saccess	regist'd	cured	eted	died fa	failed de	default fer	ferred e	eval success	ss regist'd	d cured	d eted	died		failed default	t ferred	eval	saccess
Afghanistan																										
Bahrain Diibouti	268	43	29	2	က	21	2		73																	
Egypt																										
Iran	363	71	7	œ	4	2	2		62	103	29	က	80	6	00	2	1 70	133	3 28	38	80	4	15	2	2	99
Iraq	553	89	7	2	1	00	-		75																	
Jordan	6	78	22						100	3			29		33											
Kuwait																										
Lebanon	_		100						100																	
Libyan Arab Jamahiriya	14							100																		
Morocco	1372	71	9	4	4	6	2		78	96	26		10	13	14	4	69	9 259	9 29	,,	5	6	24	9		26
Oman	_	100							100																	
Pakistan	2198	28	53	4	-	7	2	-	81									673	3 48	3 10	9	3	22	7	4	28
Qatar																										
Saudi Arabia	112	51	16	4	2	13	4	11	29																	
Somalia	411	24	4	2	က	4	30		29																	
Sudan	1640	51	56	က	0	2	2	12	77																	
Syrian Arab Republic	82	4	30	7	7	12	_		72	61	33	1	7	31	16	2	44	1 24	4 21	13	4	13	42	00		33
Tunisia	53	75		2	6	11	2		75																	
United Arab Emirates	က							100																		
West Bank and Gaza Strip		89	7	2	2	6	2		9/																	
Yemen	393	20	20			20		10	70																	
Region	7 473	51	26	4	က	œ	2	3	11	263	26	4	6	15	12	4	0 59	1 089	9 47	Έ	9	2	22	7	က	28

See Explanatory notes for previous table, page 151.

Country data for the Eastern Mediterranean: trends in DOTS treatment success and detection rates, 1994–2003

		DOT	S new sr	DOTS new smear-pos	itive treat	itive treatment success (%)	(%) ssec				۵	OTS new	smear-po	sitive ca	se detec	DOTS new smear-positive case detection rate (%)	(%)	
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
Afghanistan				45	33	87	98	8	87			2	9	2	0	14	19	18
Bahrain					13	92	73	87	88					14	62	29	11	49
Djibouti		75	77	9/	79	72	62	78	82		103	107	06	79	89	61	22	53
Egypt	52		81	82	87	87	87	82	88	43	1	11	17	31	45	49	53	99
Iran			87	84	83	82	85	8	85	47		13	35	53	58	61	09	29
Iraq					83	85	92	68	91				2	9	22	23	23	20
Jordan	06				92	88	06	98	68	112			73	71	64	71	72	88
Kuwait									55									29
Lebanon	88				73	96	92	91	91	53				90	79	72	68	29
Libyan Arab Jamahiriya					89	29			61					148	113		138	147
Morocco	98	06	88	88	88	88	89	87	89	94	94	93	88	88	84	83	84	83
Oman		8	87	91	98	92	93	06	92		114	115	110	85	117	112	109	81
Pakistan	74	70		29	99	70	74	77	77	1	2		4	2	3	5	13	17
Qatar	83	81	72	79	8	74	99	09	75	46	34	28	48	39	34	48	39	22
Saudi Arabia					22	99	73	77	92					22	37	39	38	38
Somalia		98	84	06	88	88	83	98	68		19	24	23	24	25	29	28	29
Sudan				20	92	81	79	80	78		2	<b>~</b>	29	59	34	31	33	34
Syrian Arab Republic			92	88	88	84	79	81	87			8	21	29	42	43	42	45
Tunisia					91	91	91	06	92					96	103	104	92	91
United Arab Emirates							74	62	79						29	28	24	32
West Bank and Gaza Strip									100						10	13		4
Yemen			92	81		83	75	80	82		80	30	37		54	51	47	43
Region	82	87	98	79	9/	83	83	83	84	1	6	10	17	17	22	23	27	78

**EMR** 

														-							
				MALE						_	FEMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	65+
Afghanistan	127	511	436	284	256	288	203	245	1152	1 287	814	462	305	158	372	1663	1723	1098	718	593	361
Bahrain	0	2	2	_	_	က	4	0	_	0	0	~	_	0	0	က	2	_	2	4	4
Djibouti	10	222	288	132	9/	42	24	19	127	123	55	38	28	00	29	349	411	187	114	20	32
Egypt	42	286	814	675	631	404	195	57	463	338	268	282	175	71	66	1049	1152	943	913	629	566
Iran	29	404	524	393	281	292	673	78	445	278	254	304	438	773	107	849	802	647	585	730	1446
Iraq	30	629	876	355	293	168	143	43	258	241	154	160	143	8	73	917	1117	609	453	311	177
Jordan	0	19	20	17	∞	13	0	-	9	7	2	က	12	0	-	25	27	19	11	25	0
Kuwait	_	14	39	33	26	1	2	_	16	31	18	2	3	_	2	30	20	51	28	14	9
Lebanon	0	19	26	22	9	2	7	ဇ	14	12	6	2	2	4	က	33	38	31	11	7	7
Libyan Arab Jamahiriya	0	108	266	142	32	25	19	4	43	28	30	25	21	21	4	151	294	172	25	46	40
Morocco	91	2 2 2 5	2 347	1 667	1 004	525	220	168	1455	1 029	633	431	366	351	259	3680	3376	2300	1435	891	901
Oman	0	12	10	20	19	5	11	2	11	9	0	9	9	4	2	23	16	20	25	11	15
Pakistan	278	2 582	2 2 1 9	1 763	1 623	1 268	844	613	3 032	2 371	1 595	823	478	478	891	5614	4590	3358	2446	1746	1322
Qatar	_	10	27	17	16	2	2	0	4	9	0	2	0	2	-	14	33	17	18	2	7
Saudi Arabia	5	150	285	200	145	102	107	18	210	181	75	28	51	69	23	360	466	275	203	153	166
Somalia	118	1 054	850	513	319	250	214	106	535	462	333	171	161	104	224	1589	1312	846	490	411	318
Sudan	489	1 195	1 644	1271	856	645	473	443	881	1 052	879	562	384	219	932	2076	2696	2150	1418	1029	692
Syrian Arab Republic	10	343	279	127	86	75	64	26	242	66	68	48	33	33	36	585	378	195	146	108	97
Tunisia	3	102	166	131	96	29	75	7	28	99	37	35	24	33	10	160	222	168	131	91	108
United Arab Emirates	2	10	80	12	က	2	10	4	6	2	က	3	2	4	9	19	13	15	9	4	4
West Bank and Gaza Strip	0	_	-	_	က	0	2	0	_	0	0	က	0	က	0	2	-	<del>-</del>	9	0	2
Yemen	37	555	551	376	238	148	93	72	451	404	300	196	111	02	109	1006	922	929	434	259	163
Region	1 273	10 783	11 678	8 152	6 030	4 343	3 721	1910	9 4 1 4	8 016	5 527	3 620	2 744	2 430	3 183	20 197	19 694	13 679	9 650	7 087	6 151

Country data for the Eastern Mediterranean: age and sex distribution of smear-positive cases in DOTS areas, 2003 (absolute numbers)

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

Country data for the Eastern Mediterranean: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers)

MALE FEMALE				MALE						E	FEMALE	,	,								
	0-14	15-24	25-34	35-44	45-54	55-64	<b>65</b> +	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59
Afghanistan																					
Bahrain																					
Djibouti																					
Egypt																					
Iran																					
Iraq																					
Jordan																					
Kuwait																					
Lebanon																					
Libyan Arab Jamahiriya																					
Morocco																					
Oman	5	16	22	11	10	80	4	80	15	12	12	7	5	က	13	31	34	23	17	13	7
Pakistan																					
Qatar																					
Saudi Arabia																					
Somalia																					
Sudan																					
Syrian Arab Republic																					
Tunisia																					
United Arab Emirates																					
West Bank and Gaza Strip																					
Yemen	8	26	36	23	12	9	10	2	19	22	17	80	3	4	2	45	58	40	20	6	14
Region	00	42	28	34	22	4	4	10	34	34	29	15	00	7	8	92	92	63	37	22	2
											í	4	,			2	1	3	;		

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

**EMR** 

Country data for the Eastern Mediterranean:	e Easte	rn Me	diterra		smear-	positiv	e notit	smear-positive notification rates (per 100 000 population) by	ates (	per 10(	d 000 o	opula	ion) by	age a	age and sex,	, 2003					
				MALE						III.	FEMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59
Afghanistan	2	21	26	23	31	99	61	5	52	81	72	09	61	46	4	36	52	47	45	58	25
Bahrain	0	က	2	~	2	20	42	0	2	0	0	4	6	0	0	က	~	_	က	15	21
Djibouti	7	326	592	395	346	280	237	13	187	251	160	162	170	99	10	257	421	275	251	223	4
Egypt	0	80	16	17	20	22	13	0	9	9	7	6	6	4	0	7	1	12	14	15	∞
Iran	0	2	10	10	10	21	42	_	2	2	7	11	28	20	0	2	00	00	1	25	46
Iraq	_	25	46	27	34	33	43	-	10	13	12	19	27	6	-	18	30	20	27	30	54
Jordan	0	3	4	2	5	11	0	0	1	2	1	2	11	0	0	2	3	3	4	11	0
Kuwait	0	7	1	6	14	20	23	0	6	15	10	2	6	9	0	00	13	6	11	16	15
Lebanon	0	2	∞	6	4	9	7	_	4	4	က	က	7	က	0	2	9	9	4	4	2
Libyan Arab Jamahiriya	0	16	54	46	13	15	17	0	7	9	10	12	16	20	0	11	31	29	13	15	19
Morocco	2	69	88	98	75	80	06	4	46	40	32	32	48	46	က	28	64	29	53	63	99
Oman	_	10	10	12	24	23	47	2	10	10	11	19	28	24	-	10	10	11	22	25	36
Pakistan	-	16	21	23	28	36	30	2	21	24	21	15	14	16	-	18	22	22	22	25	23
Qatar	_	24	40	17	23	24	78	0	10	22	0	∞	0	51	~	17	35	12	19	17	89
Saudi Arabia	0	7	12	7	15	23	31	0	10	10	9	6	13	19	0	∞	=	6	13	18	52
Somalia	9	110	135	124	118	166	201	4	99	72	78	59	96	82	2	83	103	101	88	129	136
Sudan	7	36	64	73	73	8	82	7	27	42	20	47	47	34	7	32	23	61	09	65	22
Syrian Arab Republic	0	16	20	14	18	25	26	-	12	7	7	80	10	11	-	14	13	11	13	17	18
Tunisia	0	10	19	20	21	27	27	-	9	7	2	80	6	11	0	80	13	13	14	17	18
United Arab Emirates	_	4	2	2	_	2	4	_	4	က	7	က	7	23	~	4	2	2	_	က	8
West Bank and Gaza Strip	0	0	0	~	က	0	4	0	0	0	0	က	0	4	0	0	0	0	က	0	4
Yemen	-	28	48	49	45	51	48	2	24	37	38	36	37	31	-	56	42	43	41	44	39
Region	-	20	30	28	30	38	40	2	18	22	20	19	24	24	2	19	26	24	25	31	32

Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available.

2003 13 808 263 3 281 11 490 11 656 566 566 566 566 567 19 17 19 17 73 100 74 100 75 1 2002 13 794 44 3 191 11 177 11 436 11 898 312 **190 394** 19 90 437 1 824 29 804 290 52 172 278 3 374 7 391 24 554 4 766 1 1 885 2001 10 139 120 4 198 10 549 11 780 11 780 342 28 285 292 34 066 284 3 327 6 852 23 997 4 997 1 945 516 2000 7 107 94 3 971 10 762 9 697 306 571 1 341 1 341 1 1050 2 29 2 686 5 686 5 686 5 686 5 686 5 79 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 52 8 68 8 52 8 68 8 68 8 72 8 **171 052** 19 90 **233 878**21
100 **136 226** 16 76 1997 1 290 4 5 3 830 2 6 6 5 2 6 6 5 3 97 7 01 7 01 2 98 2 12 2 8 4 4 5 0 2 9 8 9 4 4 4 5 0 2 0 8 9 4 4 9 7 2 49 3 332 12 338 14 189 29 196 400 836 1 282 31 771 4 307 257 3 920 20 230 5 200 2 387 507 40 14 364 **145 373** 19 90 1996 **121 745** 17 81 1995 11 145 15 936 9 697 4 98 336 983 1 1 440 29 829 276 13 142 304 2 504 14 320 4 404 2 383 3 311 3 911 113 021 19 733 443 237 940 30 316 2 518 2 023 23 178 5 127 2 376 426 1994 **864** 15 71 107 114 3426 20 569 18 553 427 217 **190 544** 14 67 1993 27 626 281 73 175 200 2 386 140 2 884 8 876 14 121 504 282 884 1 164 25 403 367 84 069 16 76 19 503 5 437 2 164 227 97 23 067 142 2 900 3 634 14 246 390 330 884 239 27 638 442 194 323 195 2 221 **295 112** 19 90 16 423 5 651 2 064 234 89 1990 4 332 117 2 100 2 142 9 255 9 255 4 439 2 77 442 27 658 482 156 759 184 2 415 **229 919** 19 90 212 6 018 2 054 285 64 1989 14 386 122 2 040 1 492 12 005 8 032 484 468 265 26 756 478 170 562 2 583 1 323 701 5 504 2 403 308 250 248 20 95 Country data for the Eastern Mediterranean: number of TB cases notified, 1980-2003 16 051 142 2 030 1 378 9 967 6 504 553 480 416 25 717 477 194 323 2 233 2 2433 2 728 693 4 952 2 309 339 85 **271 800** 20 95 **288 805** 21 100 1987 18 091 120 120 120 10 034 6 517 537 537 548 616 779 480 7 322 800 4 290 4 290 7 322 800 800 818 818 230 427 21 100 1986 10742 1942 1938 1938 1938 1943 325 26 790 861 11144 111419 20 3 966 2 722 2 722 1 698 2 722 **186 344**21
100 **171 652**20
95 2 111 2 501 534 123 **234 482** 19 90 1 867 3 062 507 136 **433 271** 19 90 1 838 2 554 597 136 **514 791**20
95 47 431 1 908 2 316 638 139 71 685 219 1 637 1 637 1 809 298 847 718 24 878 1 872 316 340 257 10 956 **522 110** 18 86 32 971 1 689 2 504 522 191 Sudan Syrian Arab Republic Tunisia United Arab Emirates Vest Bank and Gaza Region number reporting percent reporting Egypt Iran Iraq Jordan Kuwait Lebanon Libyan Arab Ji Qatar Saudi Arabia Oman Pakistan

**EMR** 

	1980	1981	1982	1983	1984	1985 1	1986	1987 ′	1988 1	1989 ,	1990 19	1991 1	1992 1	1993 1	1994 1	1995 1	1996 1	1997 1	1998 1	1999	2000 20	2001 2002	12 2003
Afghanistan	474	480	286	369	136	79	108	138	123	108		158						9	15	16	33		00
Bahrain	63	73	42	09	52	47	36	27	31	26	24	28	27	21		7	80	7	13	2	14		9
Djibouti		299	193		408	593	458	451	429	405	398	535	524	630	593		571	634	604	638		617 4	31 460
Egypt	4	က	4	4	က	က	2	42	က	က	4	9	15	9	9	18	20	22	19	18	16		91
Iran	109	59	22	19	23	18	16	19	19	22	16	25	24	8	21	26	22	20	18	18	18	18	17
Iraq	91	79	99	49	46	43	44	41	40	48	85			86	101	48	140	124	134	132	42		61
Jordan	13	28	36	34	26	28	21	19	18	16	13	11	14	11	1	12	1	6		∞	9	7	9
Kuwait	62	22	69	22	20	42	33	28	23	22	13	16	4	12	14	20	23	59	29				
Lebanon		က	က	7	15	73	85	93				32	31		31	31	26	21	19	20	16	15	12
Libyan Arab Jamahiriya	24	15	15	17	10	6	7	80	10	9		5	26			30	26		31	31	26		*
Morocco	128	144	138	129	104	122	122	118	109	111	, 113	110	100	106	115	111	116	109	103	104	66	96	66
Oman	158	74	89	28	28	99	79	37	78	27	56	23	18	14	4	12	13	12	12	10	12		0
Pakistan	392	389	379	132	66	117	152	177	186	158		171		61		11	က		99	15	80		35
Qatar	112	85	62	89	61	69	22	61	52	42	39	40		40		28	48	39	45	45	48		91
Saudi Arabia	114	81	79	99	29	31	27	21	16	16	15	13	1	13	4			16	16	16	16		4
Somalia				42	41	41	46	107	39	19					28	34	52	22	54	22	65	75	78
Sudan	170	237				7	=======================================	က	က	က	_	64	75	140	85	51	20	71	74	87	79		75
Syrian Arab Republic	19	21	19	19	20	20	35	37	4	45	47	43	40		36	30	35	32	34	34	31		7.5
Tunisia	39	35	37	4	32	34	33	29	29	30	25	25	25	30	27	27	26		24	23	21	20	19
United Arab Emirates	51	22	48	38	37	37	28	47	18	16	14	1	10		18		20		59	2	4	က	ဗ
West Bank and Gaza Strip	13	6	6	œ	7	9	က	4	4	7	က	4	4			က	<b>—</b>		_		က		
Yemen																92	91	74	73	75	92	20	09
í	:	į	:																				

31 51 35 Rate (per 100 000 population) 1997 1998 1999 29 13 11 52 10 33 16 1994 2003 6 510 7 3 1 202 5 118 8 516 8 510 1 201 1 34 1 12 82 1 12 82 1 12 12 1 12 81 013 6 509 17 1 253 4 889 5 335 3 895 91 12 914 15 151 16 265 64 1 674 4 729 10 338 1 1 447 74 962 12 804 156 10 935 77 77 1 686 4 640 11 136 1 507 1 077 69 69 69 005 2001 4 639 89 1 312 4 514 5 523 3 559 94 Country data for the Eastern Mediterranean: new smear-positive cases, 1993-2003 202 607 12 872 164 3 285 53 1 595 1 595 1 1594 1 1099 249 803 13 420 120 6 248 58 1 680 3 461 11 047 1 1577 68 971 1999 1 669 21 1 564 5 094 5 426 9 908 102 13 426 156 14 974 69 1 644 3 121 10 820 1 593 1998 1 833 25 1 690 4 915 5 105 8 933 110 1185 74 923 4 896 618 22 1 904 5 469 5 253 8 164 136 201 206 39 1 568 3 093 10 835 1 423 14 134 165 4 717 57 947 24 4 371 2 894 8 978 1 523 1 005 58 720 1996 4 229 5 347 3 194 187 175 197 14 171 135 2 578 60 1 572 8 761 1 295 1 243 46 851 3 681 1 743 1 811 4 615 5 781 161 155 148 1 168 3 728 20 428 135 1994 123 173 148 20 260 1993 82 668 800 1 006 United Arab Emirates West Bank and Gaza Strip Lebanon Libyan Arab Jamahiriya Morocco yrian Arab Republic Qatar Saudi Arabia Afghanistan Bahrain Djibouti **Suwait** udan

# **Notes**

#### **Bahrain**

Of the 263 total TB cases, 73 were in nationals.

#### **Kuwait**

Treatment after default cases were included under new smear-positive cases.

#### Libyan Arab Jamahiriya

"Transfer out" treatment outcomes were all non-nationals who left the country. Among the remaining patients, the success rate was 74%.

#### Morocco

Among 13 009 new pulmonary laboratory-confirmed cases, 167 cases were smear-negative but culture-positive.

#### **Oman**

Treatment outcomes excluded cases in non-nationals who must leave the country after conversion to smearnegative status.

#### **Oatar**

From 95 notified cases, 14 were in nationals. Among the cohort of 64 cases evaluated for treatment outcome, 10 were in nationals.

#### **Saudi Arabia**

Treatment outcomes excluded 276 cases in non-nationals that left the country, and 33 cases that were not evaluated for other reasons.

#### **Somalia**

There is a discrepancy between the population estimate used by the national TB programme (6 992 904) and that used by the UN (9 890 068). Using the country's population estimate, there would be 40 645 estimated TB cases (of which, 18 154 smear-positive cases), and DOTS detection of smear-positive cases would be 40% (instead of 29%).

#### Sudan

DOTS coverage was not reported, but assumed to be same as in the previous report.

#### **Syrian Arab Republic**

Notification data do not include 26 cases diagnosed in prison.

#### **West Bank and Gaza Strip**

DOTS coverage was not reported, but assumed to be same as in the previous report.

**EMR** 

**Africa** 

**The Americas** 

**Eastern Mediterranean** 

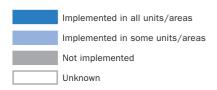
**Europe** 

**South-East Asia** 

**Western Pacific** 

## **EUROPE: SUMMARY OF TB CONTROL POLICIES**

	STATUS <sup>a</sup>	MANUAL <sup>b</sup>	MICROSCOPY°	SCC <sup>d</sup>	DOT°	MONITORING OUTCOME <sup>f</sup>
ALBANIA	DOTS	YES	MICROSCOPI	300	DOI	OOTCOME
ANDORRA	DOTS	YES				
ARMENIA	DOTS	YES				
AUSTRIA	5010	120				
AZERBAIJAN	DOTS	YES				
BELARUS	DOTS	YES				
BELGIUM	DOTS	NO NO				
	DOTS	YES				
BOSNIA & HERZEGOVINA BULGARIA	DOTS	YES				
		153				
CROATIA CYPRUS	NON-DOTS	NO				
	DOTS	NO VEC				
CZECH REPUBLIC	DOTS	YES				
DENMARK	DOTS	YES				
ESTONIA	DOTS	YES				
FINLAND	NON-DOTS	NO NO				
FRANCE	NON-DOTS	YES				
GEORGIA	DOTS	YES				
GERMANY	DOTS	YES				
GREECE	NON-DOTS	NO				
HUNGARY	DOTS	YES				
ICELAND	DOTS	NO				
IRELAND	NON-DOTS	YES				
ISRAEL	DOTS	YES				
ITALY	DOTS	YES				
KAZAKHSTAN	DOTS	YES				
KYRGYZSTAN	DOTS	YES				
LATVIA	DOTS	YES				
LITHUANIA	DOTS	YES				
LUXEMBOURG	DOTS	NO				
MALTA	DOTS	YES				
MONACO						
NETHERLANDS	DOTS	YES				
NORWAY	DOTS	YES				
POLAND	DOTS	YES				
PORTUGAL	DOTS	YES				
REPUBLIC OF MOLDOVA	DOTS	YES				
ROMANIA	DOTS	YES				
RUSSIAN FEDERATION	DOTS	YES				
SAN MARINO	DOTS	NO				
SERBIA AND MONTENEGRO	DOTS	YES				
SLOVAKIA	DOTS	YES				
SLOVENIA	DOTS	NO				
SPAIN	NON-DOTS	YES				
SWEDEN	DOTS	YES				
SWITZERLAND	NON-DOTS	YES				
TAJIKISTAN	DOTS	YES				
TFYR MACEDONIA	DOTS	YES				
TURKEY						
TURKMENISTAN	DOTS	YES				
UKRAINE	DOTS	YES				
UNITED KINGDOM	NON-DOTS	YES				
UZBEKISTAN	DOTS	YES				



- a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)
- b Manual: national TB control manual (recommended)
- c Microscopy: use of smear microscopy for diagnosis (core component of DOTS)
- d SCC: short course chemotherapy (core component of DOTS)
- e DOT: directly observed treatment (core component of DOTS)
- f Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)

**EUR** 

	Inck	×۱			ž -	Prevalence, 1990				Death, 1990				cidence	_		ď.				Death,		
	All cases life. HIV+	- 1	new sst IIIG. nivt	₹	number	rafe A	number rate number ra	Tafe A	number	₹	cases exci.	- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	All Cases IIICI.	rate 1	new ss+ IIIci. n	rate	number rate	All cases e	<u> </u>	All Cases	er rate	All cases exc	
Albania	1	27				46		46	_			4		23		9 9				L	اءً		l
Andorra		37		16	22	14		14	က	9	3	9	13	19	9	80					1 2		2
Armenia		26		12	1 569	44	1 569	44	162	2	162	2	2 146	20	964	32							3 11
Austria	1 535	20		6	1 718	52	1718	22	246	က	246	က	1 172	4	524	9						117	+
Azerbaijan		35	1 133 1	16	4 238	20	4 238 6 525	59	437	9 ^	437	9 ^	6 351	76	2 858	8 8	9 122 109		9 121 109	914	4 0	913	2 11
Spinos.		3 0		. 0	2 1/10	5 8		200		- ("	308	- 0	1 416	3 5	634	1 4						177	
Bosnia & Herzedovina		2- 70		12 0	6 920	161	`	161		ر د	938	, <del>L</del>	2 2 7 9	י ב	1 025	2 2						7 6	
Bulgaria	2 459	78 2	1 106	13	4 138	47		47	427	2 5	427	2 5	3 423	8 2 8	1 540	2 2	3711 47		3710 47	7 466	9 9	466	9
Croatia		74		33	6 123	126	6 123	126		12	565	12	1 909	43	859	19						295	5 7
Cyprus		2		2	28	6		6		8.0	2	_	34	4	15	2	32				8	.,	~
Czech Republic		37		16	4 222	41		41		9	909	9	1 191	12	535	2					5 1	13	1
Denmark		16		7	868	17	868	17	129	က	129	3	429	∞	192	4					3 1	42	1
Estonia		32		14	840	23		23		2	87	2	658	20	293	22					2 6	6	3 7
Finland		18		8	286	20		20		3	142	3	483	6	217	4					0 1	20	1
France	17 776	31	7 907	4	19 896	35	19 896	35	2 854	2	2 854	2	7 257	12	3 228	2	7 4 13 12		7 226 12			716	
Georgia		88		17	3 511	4		4	362	_	362	_	4 244	83	1 906	37					0 14	69	13
Germany		78		13	25 228	32		32		2	3 619	2	6 749	∞	3 027	4						999	
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Kyrgyzstan	2 290	52		23	3 854	88		88	398	6	398	6	6 391	124	2 874	26		7 200				926	9 18
-atvia		33		15	1 525	26		26	157	9	157	9	1 735	75	922	8						246	
Lithuania	1 416	8 8	637	17	2 382	<b>4</b> 8		29 8	246		246		2 398	02 9	1 078	٤,		2,		320	0 1	319	o ,
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Poland		53		24	34 655	91		91		80	3 195	∞	11 878	31	5 342	14		Ĺ				1 58.	
Portugal		71		31	7 848	79	7 848	79		1	1 126	7	4 497	45	2 000	20						456	
Republic of Moldova		64		59	4 700	108	`	108		11	485	1	5 918	139	2 657	62						84	
Romania	17 668	92 9	7 949		29 729	128	` .	128		5 0	3 067	5 0	33 276	149	14 972	29	43 382 194	43 370				4 48	
Kussian Federation		8 6			1203//	14	120 377	14		» ς	12 417	» ς	160 688	21.	786 17	2 %	229 046 160		761 184	29 024	4 6	26 /94	+ 19
Serbia & Montenedro		2 65		27	10 323	102	1	102		1 6	952	1 6	3 648	35	1 639	16						525	
Slovakia	2 187	42	984	19	3 743	71	3 743	71			345	7	1 312	24	591	7 =	1 569 29	1 569	569 29		9 4	208	
Slovenia		45		20	1 476	77		77		7	136	7	366	18	165	8						5	3
Spain		62		28	27 399	20		20	3 930	10	3 930	10	11 237	27	4 974	12		10 932				1 082	
Sweden	713	ω ;		4	798	စ		0	115	_	115	_	393	4	176	7					6	8	_
Switzerland	ľ	198		ω <u>ς</u>	1 395	8 3	ľ	5 20	200	e (	200	e (	529	/ 50,	235	e	ľ		ľ	_			
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Turkey	26 321	t 4	11 844	-2.4	45 042	78		78	4 153	٥ /	4 153	0 ~	18 555	26	8 349	± 2	28 872 40	28		3 166	+ (0		
Turkmenistan		59			3 639	66		66	375	10	375	10	3 272	29	1 472	30							
Ukraine		39			33 892	65		65	3 525	7	3 491	7	44 674	95	19853	41			•	9		5 69	
United Kingdom	6 546	12	2 936	3.7	7 327	13	7 327	13	1051	2 5	1 051	2 5	30 001	112	3 165	22	7 160 12	7 108	108 12	2 706	3 1	696	7 1
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45 0 4 to 48 Notifications
All cases New ss+ 138 2,219 0 185 4,661 22,546 482 12,785 1,455 1,948 157 353 438 234 141 107 392 5,740 0 342 288 571 2,033 554 3,378 170 48 39 52 46 33 38 38 100 49 49 61 67 34 26 37 67 100 52 33 55 86 87 57 126 126 DDR 44 28 0 6 2 282 52,983 1,742 1,029 5,757 6,322 6,322 292 319 200 116 0 715 989 1,679 0 526 150 1,481 3,665 1,643 1,643 31 DOTS 25 0 8 82 0 All cases 1,282 320 3,677 3,861 14,909 21,064 1,127 1,127 1,127 275 904 275 882 882 483 201 10 1,250 1,615 1,030 1,740 3,069 3,069 3,069 1,101 378 378 505 4,234 26,936 6,172 1,686 2,586 54 36 15 0 52 102 128 81 81 86 61 61 77 77 77 69 75 75 98 98 105 101 83 91 69 77 27 135 135 1,338 5,691 10 10 10 10 10 14 44 558 10,659 728 3,485 31,797 206 ,987 572 281 188 188 3,164 5,477 318 208 208 0 15,706 142 441 119 860 1,055 196 37 122 147 145 895 Whole country 124 1 599 1,871 406 951 1,274 756 756 1,919 1,919 1,919 1,919 1,919 1,620 233 1,620 244 253 1,620 274 284 1,919 2,067 161,308 541 149 5,055 876 458 101 2,731 125 302 250 Country data for Europe: notification, detection and DOTS coverage, 2003 Pulm confirmed number 336 205 2,906 211 211 1,197 1,621 91,809 282 52 2,983 1,742 1,742 10,418 28,868 29,868 319 200 200 200 116 2,816 319 200 100 100 543 10 1,538 1,282 320 3,677 3,619 124,041 1,276 28,335 1,124,041 1,276 27,63 2,763 2,763 3,895 1,176 2,763 3,895 4,260 4,260 653 421 529 4,518 32,169 7,027 1,726 2,821 54 3,931 1,128 1,780 3,263 3,263 3,263 3,263 3,263 3,263 3,263 3,263 3,263 6,350 4,076 1,157 2,919 983 293 5,918 410 623 697 561 11 1,570 3,166 1,100 1, 5,402 1,984 41,060 8,876 7,169 6,245 2,056 71,325 71,325 71,325 74,867 48,523 59,251 26,093 · Kosovo - Serbia & Montenegro Belarus Belgium Bosnia & Herzegovina Bulgaria Croatia ortugal tepublic of Moldova ithuania uxembourg

See Explanatory notes, page 151.

Country data for Europe: treatment outcomes for cases registered in 2002

Country data for Europe, treatment outcomes for cases registered in 2002	rope. rican		No.	Smear	-positive	New smear-positive cases - DOTS	DOTS	100			-			New sme	New smear-nositive cases – non-DOTS	e Cases	- non-D	STC			-		Sme	Smear-nositive re-treatment cases - DOTS	ve re-trea	tment ca	Ses - D(	TS		
			%	%	%	%	%	%	6 %		%		%	°	% %	%	%	%	%	%	%		%	. %	%	%	%	%	%	ļ
	Number of cases		of notif		-Idmoo			ţ				Number of cases	of	otif	-ldwoo	_			trans-	not		Number		-ldwoo			trans-	_		
	notified	regist'd re	regist'd c	cured	eted	died	failed de	default fe	ferred ev		saccess	notified re	regist'd regist'd	t'd cured	- 1	died	failed	default	ferred	eval su	saccess	regist'd c	cured	eted di	died fail	failed default	ult ferred	d eval	saccess	ssac
Albania	93	93	100	29	24	-		4		4	06	132	132 10	100 2	27 52	2	2	80	-	00	80	10	20	20		2	20	10		02
Andorra Armenia	2 295	3 295	150	8 8	6	4	က	2	က	8	001	216										62	56	Ω	9	19	e	3 37		31
Austria	220																													Γ
Azerbaijan Belarus	1 310	748	22	81	ღ	2	2	6	က		28	351	401	114 8	80	9		10	4	4	80	29	22	7	7	13 1	15		9	28
Belgium	419	337	80	24	45	6	-	-		19	69											38	29	47	80			16		92
Bosnia & Herzegovina	526	526	100	8	7	~	2	_	~		92											28	74	19	2	3	2			93
Bulgaria	742	742	100	98		3	4	2	2		98	265	265 10	100 7	78	9	6	2	1		78	293	64	15	4		8 1		7	19
Croatia												437		47 6	32 8		-	6	က	7	70									
Cyprus	80	00	100	38	38	13			13		75											5	100							8
Czech Republic	329	320	26	29	13	4		_			73											8	25	25			13			20
Denmark		135		4	36	9	<del>-</del>	4	4	2	77	135										10	20	2				10		6
Estonia	203	203	100	62	2	4	0	ກ	0		19	130										92	28	2	_	4 21	<u>.</u>	37		¥
rimand											+	130									$\dagger$									Τ
riance	700	200	5	i c	c	c	c	0	ć		LI C	2770										4	ii c	-	и	0		ć		90
Germany	1868	1329	3 5	8 8	26	4 0	0	2 0		19	69											164	3 4	23	, <del>L</del>		,	212		67
Greece											-	212									H									Г
Hungary	556	549	66	39	16	6	6	2	· Ю	19	22	!										110	27	12	15	17	2	2 22		39
Iceland	2	5	100	20	20						100																			:
Ireland												100	95	95	5 72	4	2			17	77									
Israel	271	257	92	72	6	=	0	2			81											18	72	=	9		÷	_	80	83
Italy	1 275	185	15	30	49	2		4	4	. 11	79											22	32	23		1	14 9	9 23		22
Kazakhstan	9 452	9 185	26	9/	<del>-</del>	2	=	4	2		78											3 307	62					3		49
Kyrgyzstan	1 456	1476	101	8 i	က	4 (	۲.	ı 2	7		82	131										820	56	17	4 ;	7	ი	42		43
Latvia	636	636	100	44	7	5 9	- -	2	0		9/	0			9						+	192	3/			ľ				50 00
Lithuania	634	634	901	7.7		10	4	ກ	0	4	7.7	188	188	100	69	13	9	10		7	69	282	36		19	8	02	14		 65
Luxembourg	<u>-</u> 10	ĸ	100		9	40					9																			
Monaco	,	,	3		3	2					3										+									Τ
Netherlands	330	296	06	16	21	m		Ç		22	89											26	12	27		0	23	38		82
Norway	31	35	113	63	17	9		0	. 9		80											0	20		13		13			75
Poland	3 060	2 602	82	75	10	9	-	9	-		98											445	49		10	2 1				73
Portugal		1 902	96	6	73	9	0	2	4	e	82											247	10					2 6	2 9	15
Republic of Moldova		556	100	53	80	80	9	16	4		61									53	+	453	30							37
Romania Russian Federation	6 086 5 179	6 459 5 171	9 2	2 2	3 3	ა წ	മ		- 4	_	9/	4 61 7 22 686	5 141	111	56 15	2	_	00	-	10	0,	2 915 962	37	ر 9 9	9 21	11 26	9 6	18		46
San Marino											+																			
Serbia & Montenegro	402	664	165	25	26	ი (	<del>-</del>	4	-	0	91											94	48		o !	e (	6		00 0	80 5
Slovakia	202	730	99 0	8 8	20	9 %		ď			8 8 2											42	8 %	33	/ 2		ď		20 1-	7 87
Spain	25	000	3	67	99	0					3	3 317										-	+7		+7					-
Sweden	109	108	66		73	9			,	. 61	73											13		46		00		46		46
Switzerland											+	123																		
Tajikistan	100	107	107	6/	!	2	7	Ω :	<del>-</del>		79	587									-	;	;	;	,		,	•		- ;
TFYR Macedonia	143	143	100	62	17	<del>-</del>		9		-	6/	2/	57	100	46 39	2	2	2	2	2	84	58	8	\$	m	3 21	<u>.</u>	,	e e	69
Turkmenistan	735	735	100	20	7	2	13	4	-		77	519	519 10	100 6	69					31	69	529	42	21 1	12 1	16 8	-		63	8
Ukraine													1 031	76	62	9	-	ď	-	24	63									
Uzbekistan	2 766	2 544	92	99	14	2	8	7			80	2 017						,	.		3	519	38	23 1	10 1	12 17			61	-
Region	43 112 4	40 307	93	63	13	9	9	9	2	3	, 92	40 450	8 622	21 4	47 21	4	4	7	-	15	69	12 551	42	12 1	10 1	12 11		10		24
																														1

See Explanatory notes, page 151.

not eval % trans-ferred default 6 8 4 After default – DOTS % % failed ω σ died 9 7 % compl-eted 14 16 cured 4 15 1 145 regist'd saccess 41 72 6 2 8 % not eval default 9 8 After failure – DOTS % failed 44 4 died 5 4 11 22 completed cured regist'd 1 566 87 380 436 Country data for Europe: re-treatment outcomes for cases registered in 2002 saccess 63 65 83 46 46 75 73 80 80 41 51 81 79 % not eval 0 6 30 % trans-ferred က default α α 9 6 6 5 12 8 6 6 Relapse – DOTS failed e + 15 7 10 10 10 26 died 13 2 10 1 1 2 2 2 6 11 4 7 17 22 9 17 21 compl-eted 5 5 25 11 26 2 25 9 67 14 14 33 50 cured 70 4 4 37 63 41 46 46 50 64 12 38 37 37 48 29 regist'd 52 19 2 908 414 163 179 8 445 132 255 946 962 42 14 8 759 Belarus Belgium Bosnia & Herzegovina Bulgaria Croatia San Marino Serbia & Montenegro Slovakia Norway Poland Portugal Republic of Moldova Russian Federation Tajikistan TFYR Macedonia Cyprus
Czech Republic
Denmark
Estonia
Finland
France
Georgia Jnited Kingdom Turkey Turkmenistan Italy Kazakhstan Kyrgyzstan Latvia Lithuania Monaco Netherlands Albania Andorra Armenia Austria Azerbaijan Switzerland Jzbekistan Germany Greece Hungary Iceland Ireland Israel Romania Slovenia weden Jkraine Region Malta

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See Explanatory notes for previous table, page 151

Country data for Europe: trends in DOTS treatment success and detection rates, 1994–2003

DOTS new sme	200	DOT	S new sr	near-posi	DOTS new smear-positive treatment success (%)	ment suc	ccess (%)	- 1			DOT	S new sn	ear-posi	DOTS new smear-positive case detection rate (%)	detectio	n rate (%		
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
دند، طاح								ao	00							22	ac	g
Andorra					100	67	20	100	100			226	4	53	75	2 4	33.5	116
Armenia		83	77	82	8 2	8	87	06	62	13	59	20	46	42	5. 2	32	3 8	43
Austria						77	73	64							58	48	41	
Azerbaijan			98	87	98	88	91	99	84	2	6	7	7	7	9	0	46	78
Delarus								5	6							77	Į.	‡ [
Bosnia & Herzegovina				ö	œ	8	70	0 4 8	90 0				37	84	œ	75	0 0 0	2/48
Bulgaria				3	3	3	5	87	98				5	5	23	9 9	47	8 8
Croatia																		
Cyprus					42			95	75				94	47			23	91
Czech Republic	73	09	99	69	65	78	20	73	73	45	59	53	65	29	09	61	26	63
Denmark						8	î	č	77						č	Č	3	75
Estonia Einland						63	2	94	/9						94	S S	1.9	9
France																		
Georgia		28		92	28	61	63	29	65	17	33		31	40	31	52	51	52
Germany				72	72	28	77	29	69			63	64	99		23	26	22
Greece																		
Hungary					88		2	46	55					36	25	37	40	4
Iceland								29	100							74	53	28
Ireland																		
Israel		6			í	i	78	62	18		;	•		í	9 (	26	28	22
Italy		80	82	69	7.7	5	4/	40	6/		14	5	13	96	32	10	65	2
Kazakhstan			;	i	6 1	6/	6/	78	78				4	79	94	93	32	98
Kyrgyzstan		į	80 1	9/	82	. 33 1	82	ω i	82		ကျ	4 ;	33	61	44	i	51	27
Latvia		61	64	65	7.1	74	72	73	92		73	71	74	65	73	9/	78	83
Lithuania					62	\$	95	75	72					က	7	31	28	82
Luxembourg		100	100	100	001	75	100	100	09		34	22	45	7.1	41	74	67	126
Monaco		2	3	2	8	2	2	2	8		5	77	2		-	3	2	2
Netherlands	81	72	81	80	92	6/	9/		89	75	48	44	37	47	46	51	22	20
Norway		77	80	4	69	77	20	87	80		99	29	35	16	29	48	56	46
Poland					75	69	72	77	86				2	3	4	3	22	26
Portugal	48	69	74	78	74	82	79	78	82	77	77	29	82	62	84	92	92	87
Republic of Moldova							83	99	61							36	51	39
Romania				72	82	78	80	78	92				84	4	o	10	4	38
Russian Federation San Marino		9	29	100		ç	89	/9	/9		0	102	-	N	1 1 1 2	Ω	_	ກ
Serbia & Montenegro								88	91							26	24	37
Slovakia	96	64	73	29	82	79	82	87	85	9/	81	32	38	34	35	35	33	8
Slovenia		06	87	82	78	88	8	82	85		75	26	62	71	29	20	71	70
Spain							í		í							i	í	
Sweden							£	29	٤/							\$	69	
- Switzerland									0,0								c	
TEYR Macedonia							98	88	6/							75	7 84	49
Turkey																		
Turkmenistan							69	75	77						19	40	47	49
Ukraine																		
Uzbekistan					78	6/	80	92	80				0	7	4	7	22	20
c		8	1	f	1	1	1	1	1				;	;	,	;	8	8
Region	89	69	72	72	9/	11	11	75	9.2	3	က	2	11	1	12	14	22	23

65+ 17 0 4 051 42 42 123 147 - 48 E E 110 2 2 3 3 707 195 46 570 399 63 31 ۲, 17 0 7 117 19 75 153 14 26 7 4 76 127 2 0 370 370 126 62 62 537 512 23 7 55-64 3 245 6 785 266 206 1 281 39 25 62 62 141 212 1 252 51 51 66 244 79 28 47 206 6 821 286 45 83 241 35 49 245 183 212 8 8 588 415 256 205 746 41 27 7 871 263 12 342 460 238 1287 11 06 33 33 32 7 822 173 50 79 275 203 340 116 111 17 14 67 70 4 17 17 14 142 7 7 184 233 233 219 815 720 14 8 85 47 59 59 4 991 18 2 53 0 4 0 1 8 5 2 2 0 0 0 0 0 10 0 0 45 7 73 58 0 8 4 5 0 20 38 1 358 47 21 224 224 163 16 1 724 0 0 0 1 0 0 55-64 - 5 20 20 0 0 65 27 27 9 9 92 4 Country data for Europe: age and sex distribution of smear-positive cases in DOTS areas, 2003 (absolute numbers) 11 14 44 - 2<u>1</u> 9 2 43 24 35 38 0 2 132 39 39 37 220 282 3 3 3 1 317 ω 15 34 63 15 17 17 17 86 42 37 4 4 152 82 82 54 284 339 10 4 1 938 8 - 18 42 49 25-34 39 17 37 37 2 13 4 96 9 108 163 58 437 401 2 667 35 35 2 089 7 1 29 0 6 9 7 61 63 88 380 233 4 ∞ ռ Ţ 1 0 0 0 8 - - 8 0 2 327 ۲ 0 1 18 60 35 50 89 32 60 60 0 2 349 148 25 25 236 25 15 ∞ 155 146 55-64 15 0 96 17 45 19 7 48 24 24 59 107 2 2 305 305 99 99 53 437 420 2 541 103 2 58 67 22 35 4 653 227 169 1061 226 40 40 52 200 106 174 36 22 5 504 1 64 243 30 49 49 38 38 168 141 175 7 4 436 333 202 921 407 31 23 5 933 8 0 69 234 297 180 850 8 6 134 33 42 169 4 8 8 8 8 219 74 116 10 5 155 15-24 35 35 0 0 120 67 27 32 99 - = = -9 0 6 6 93 134 131 435 487 2 902 0 0 - 0 0 2 6 2 1 1 2 0 Belgium Bosnia & Herzegovina Serbia & Montenegro Republic of Moldova Russian Federation FYR Macedonia Szech Republic Jnited Kingdom Turkmenistan -uxembourg Vetherlands Austria Azerbaijan Kazakhstan Switzerland **Kyrgyzstan** San Marino Jzbekistan Denmark Estonia -ithuania Hungary Iceland Ireland Monaco Sulgaria Georgia Vorway Slovakia sermany Portugal Slovenia Sweden roatia -rance Jkraine nland Turkey Syprus Greece rael atvia /alta

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries

Country data for Europe: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers)

County data for Europe, age and sex distribution of sineal-positive	מאני מ	Vac nii	nginen	2	וופמו אי	Silive St	200	cases III IIOII-DOI 3 aleas, 2003 (absolute IIuIIIbels)	2000	2002 (a	Panice	5	6	-							
	0-14	15-24	25-34	35-44	45-54	55-64	65+	0-14	15-24	25-34 35-44		45-54 5	55-64	65+	0-14	15-24 2	25-34 35-	4	45-54 58	55-64	65+
Albania	0	16	11	16	6	10	12	2						10							22
Andorra	10	0	39	7	46	8	0	0	0	0	2	00	7	0	10	0	39	13	25	4	0
Austria	2		3	:	2	5	,				1 :			,			8 1		5 1	: :	
Azerbaijan Belarus	0	92	89	75	36	59	4	0	12	7	21	17	19	0	0	77	75	96	53	48	4
Belgium Bosnia & Herzegovina Bulqaria																					
Croatia	0	15	27	89	80	42	09	-	14	19	18	10	15	69	-	29	46	98	06	22	129
Cyprus Czech Republic																					
Denmark																					
Finland	0	2	3	8	19	17	59	0	2	10	3	9	5	31	0	4	13	11	25	22	9
France	18	129	249	223	190	127	210	16	114	129	79	44	32	159	34	243			234		369
Georgia Germany																					
Greece	2	20	28	25	23	25	36	0	7	6	7	2	2	18	2	27	37	32	25	30	55
Hungary																					
Ireland	0	10	11	13	14	7	11	0	4	7	9	4	-	10	0	14	18	19	18	8	21
Israel																					
Kazakhstan																					
Kyrgyzstan	0	7	0	ဇ	0	က	0	0	-	0	4	0	0	9	0	œ	0	7	0	3	9
Latvia							+							+							
Lithuania																					
Malta																					
Monaco																					
Norway																					
Poland																					
Portugal																					
Republic of Moldova	0	21	21	20	37	6	0	0	13	13	10	3	7	-	0						-
Komania Russian Federation San Marino	0 0	315 1 641	3 786	774 4 572	892 4 696	399 1 594	822	0 79	, 923	333 1 352 1	186 1 198	192 999	370	180 535	0 2	602 564	1 048 5 138 5	0420	1 084 5 695 1	495 964 1	428 1 357
Serbia & Montenegro																					
Slovakia																					
Spain	7	153	334	305	219	132	222	9	138	218	113	51	29	87	13	291	552	418	270	161	309
Sweden		;				,	;	, ,		) (	, ,	. (		;	! ,				i (		;
Swizerland	0			6	2	4	+	-	2	2	4		2	2	-	17	=	23	2		4
TFYR Macedonia	0	က	9	б	10	2	6	0	2	2	-	2	0	4	0	80	80	10	15	2	13
Turkey	(	C	9	007	L	,		(		0	3	3									
Turkmenistan	0 6	50	719	108 2 808	55 2 634	13	517	0 0	23	60	24	273	221	0 421	30 0			0	76		23 33
United Kingdom	13 5	101	182	128	81	59	95	23 14	108	148	è 88	47	17	55	27	209	330		128	76	147
Uzbekistan	0	149	373	298	234	124	88	0	63	265	162	86	54	39	0						128
Region	92	3 558	8 012	9 513	9 285	3 616	2 475	92	2 244	3 332 2	2 487 1	878	894 1	628	171 5	5 802 11	11 344 12 (	12 000 11	11 163 4	4 510 4	4 103

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

16 24 29 36 4 - 0 1 2 2 4 6 f 55-64 31 23 1 43 45-54 38 44 2 29 ALL 35-44 2 11 11 29 52 74 74 34 53 55 40 10 0 46 47 2 28 15 25-34 29 67 37 35 16 2 53 40 4 35 15 15-24 35 4 4 8 0 24 18 3 15 0-14 00000 0 0 22 8 22 2 21 10 15 0 +59 18 9 10 0 Country data for Europe: smear-positive notification rates (per 100 000 population) by age and sex, 2003 - 0 11 0 0 33 45-54 25 21 16 0 4 6 113 114 117 0 0 0 20 10 19 35-44 32 25 44 25 3 26 15 2 21 6 5 4 2 4 25-34 58 27 21 3 10 3 9 37 22 4 29 35 4 4 8 9 10 19 41 3 13 0-14 0 000 0000 15 55 27 34 8 0 0 t t 8 c 2 9 15 25 2 41 65+ 21 
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**%** 9 α α ο

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Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available

	1980	1981	1982	1983	1984	1985	1986	1987	1988		1990	1991	1992	1993	1994				1998 1			2001 20	2002 2003
Albania	1 050	954	876	891	975	916	686	915	759		653				707	641							594 543
Armenia	756	924	759	702	774	768	832	992	651		590												<b>~</b>
Austria	2 191	2 061	1 942	1 825	1 765	1 442	1377	1 390	1 402		1 521									1			
Azerbaijan Belarus	3 080	3 180	3 217 5 468	3 176 5 509	3 506 5 065	3772	3 804	3 677	3 340	2 989 3 708	2 620 3 039	2 771 3 745	2 821	3 036	2 839	1 630	2 480 4	4 635 4 5 985 6	4 672 4 6 150 7	7 339 67	5 187 4 8 6 799 5	4 898 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	5 142 3 840 5 139 5 106
Belgium	2 687	2 837	2 652	2 190	2 149	1 956		1 772	1 588		1 577							١.	1				
Bosnia & Herzegovina	4 421	4 376	4 678	4 468	4 691	4 666		4 522	4 093		4 073												
Bulgaria	3 280	3 007	2 999	2 892	2 856	2 555		2 352	2 387		2 256												
Croatia	3 388	4 021	3 718 86	3 632	3 612	3 605	3 355	3 326	2 973		2 576												
Czech Republic	4 962	4 312	4 146	4 016	3 653	3 117		2 196	2 047		1 937												
Denmark	430	394	378	348	302	312	1	322	304		350								1				
Estonia	614	260	563	287	546	541		446	471		423												
Finland	2 247	2 204	2 170	1 882	1 791	1 819		1 419	1 078		772												
France	17 199	16 459	15 425	13 831	12 302	11 290		10 241	9 191		9 030	8 510											
Georgia	20 008	2 124	2 168	1 881	1 855	1 822	1833	1 810	1 598														
Germany	29 991	27 003	180 07	116 77	4 050	4 550		1, 102	707 01						12 302 1								
Greece	5 412	455	5 193	3 880	1 956	1 226	1 266	1 193	907														
rungary Iceland	21412	222	25	5 U28 24	4 4 7 2	4 652		4 125	4 016 16			3 636 15	3 960 16	4 209	4 103 18			4 240 10				2 923 2	
Ireland	1 152	1 018	975	924	837	804		581	534														
Israel	249	227	232	222	257	368		184	226														
Italy	3 311	3 182	3 850	4 253	3 472	4 113	4 077	3 278	3 610														
Kazakhstan	14 442	13 876	13 808	13 357	12 563	12 423		13 286	13 501														
Kyrgyzstan	1 973	2 085	2 051	1 981	2 022	2 094		2 088	2 159			2 515											6 613 6
Latvia	1 194	1 140	1 077	1 072	1 054	1 223	985	948	938														
Lithuania	1 636	1 599	1 495	1 477	1 420	1 453		1372	1 339														
Luxembourg	24	26	- 4	24	δ <del>τ</del>	7 1	5 <del>1</del> 4 5	0 4 4	5 5														
Monaco	-	0	0	0	0	-		2	i ~														
Netherlands	1 701	1 734	1 514	1 423	1 400	1 362	1 238	1 227	1 341													1 408 1 3	
Norway	499	461	448	396	373	374	343	307	294														
Poland	25 807	24 087	23 685	23 411	22 527	21 650	20 603	19 757	18 537														
Portugal Penublic of Moldova	0 781	7.249	3 107	7 052	2 554	2 732	9 0 2 2	7 810	9 510	0 084	6 214 1 728	1 940	1 835	2447	5679			2 908	5 260 4		2035 31	3608 3769	381 3861
Romania	13 553	13 602	13 588	13 570	12 952	12 677	12 860	13 361	14 137														'
Russian Federation	74 270	73 369	72 236	73 280	74 597	64 644	71 764	70 132	67 553	62 987		50 407				84 980 11	111 075 119			134 360 140 (	•		873 124 041
San Marino Serbia & Montenegro	6 232	6 381	6 274	6 443	6.454	8 246	8 128	6 042	5 583	5 045	7 107												
Slovakia	2 465	2 304	2 263	2 252	2 152	1 989	2 0 2 2	1 830	1 651	1 501	1 448												
Slovenia	1 085	939	982	925	896	923	816	792	760	768	722				526								
Spain	4 853	5 552	7 961	8 987	10 078	10 749	13 755	9 468	8 497	8 058	2 600			9 441									
Sweden	926	875	784	832	754	702	640	545	536	595	557	521	610	616	537	564	497	456	446			394	375 386
Jaiikistan	2 647	2 631	2 628	2 509	2 427	2 485	2610	2 727	2 474	2 621	2 460			652									
TFYR Macedonia	1	9	1	1	i	1	2	1		1	2	5	1 602	1712									
Turkey	36 716	39 992	26 457	28 634	27 589	30 960	31 029	30 531	27 884										25 501 22				
Turkmenistan	1 677	1 625	1 559	1 541	1 604	1 607	1614	1 956	1 904	2 169		2 358											3 671 3 771
Ukraine	26 095	25 646	01/ 42	24 216	4 356	24 058	22.946	22 145	20 744														
Uzbekistan	9 163	9 682	8 697	8 817	8 544	8 717	9 427	9 794	10 134	10 632	9 414		9370	9774	14 890	9 866	11 919 13	13 352 14	14 558 15	15 080 157	15 750 17	17 391 20 588	588 20 700
Region	348 921		324 580	319 220		298 933	302 602													"			
number reporting	48					48	48	48	48	49	209	49	49	47	46	20	51	51	51	51 51	51 51	51 51	51 49
percent reporting	94	94	94	94	8	94	94																

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Country data for Europe: case notification rates	Europe:	caseı	notific	ation r		(per 100 000 population),	d 000 (	opula	_	1980-2003	2003												
	1980	1981	1982	1983	1984	1985	1986	, 286	1988	989 1	1990 19	1991 19	1992 19	1993 19	1994 1995	5 1996	6 1997	1998	1999	2000	2001	2002	2003
Albania	39	35	35	31	34	31	33	29	24	21											18	19	17
Andorra										24											15	7	14
Armenia	24	29	24	22	24	23	25	22	19	18		21									45	47	20
Austria	5 28	27	26	24	23	19	<del>2</del> 5	18	<del>0</del> 5	7											12	13	9
Azerbaijan Ralariis	2 2	2	2 2	84	22	70	20	30	37	38											55	20	52
Belgium	25	5 8	22	8 6		2 8	- 6	3 2	. 4	2 2											13 6	12	10
Bosnia & Herzegovina	113	111	117	111	115	113	110	106	94	96											61	1 4	42
Bulgaria	37	8	34	32	32	29	28	26	27	26											48	42	39
Croatia	91	95	82	83	82	81	74	72	63	26											31	33	31
Cyprus	7	=	14	12	9	6	7	2	9	က											5	က	4
Czech Republic	48	45	40	39	35	30	25	21	20	18											13	11	1
Denmark	∞ ,	∞	7	7	9	9	9	9	9	9											0	∞	7
Estonia	45	88	38	39	36	32	34	29	30	27											52	46	42
Finland	47	46	45	39	37	37	31	59	22	50											o	တ	∞
France	32	8 4	788	25	22	5 50	19	3 3	16	16											10	10	10
Georgia	14	242	25	30	35	4 %	22	22	23	30											20	/8/	82
Germany	8 8	8 4	2 6	000	0 6	2 4	2 4	7 7	- 0	5 5											υс	οц	о и
Hindary	27.5	5.05	84	47	42	46	5 4	39	n 80	- 95											60	27	25.
Iceland	7	10	1	10	1 =	2 42	2	5	9	2											4	i m	2
Ireland	8	98	78	26	24	23	17	16	15	19											10	10	ı o
Israel	7	9	9	9	9	0	9	4	2	4											6	80	00
Italy	9	9	7	8	9	7	7	9	9	7											7	7	7
Kazakhstan	26	95	90	98	80	78	81	81	82	80											169	178	175
Kyrgyzstan	22	99	54	51	51	52	52	20	51	49											133	131	120
Latvia	48	45	43	42	41	47	38	36	35	32											82	77	73
Lithuania	48	47	43	45	40	41	39	38	36	37											75	70	75
Luxembourg	207	12	-	- 1	13	11	12	13	4	17.											\	\	12
Malta	_ `	ω (	4 (	۲ (	4 (	ო •	4 1	4 1	ကေ	4 (											4 (	9 (	7
Monaco	4 (	o (	0 7	0 (	o (	4 (	<b>~</b> 0	<b>\</b>	m c	თ (											0 0	0 0	
Nemeriands	2 5	7 2	= =	2 2		n   c	0	0 1	2 2	n   u											n u	0 4	0 1
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Portugal	2 2	7 5	8 4	7 1	69	8 69	99	71	64	67	63	60									43	4 4	388
Republic of Moldova	69	70	78	69	61	65	71	99	58	52											84	88	85
Romania	61	61	61	09	22	26	26	28	61	63											127	133	127
Russian Federation	22	23	51	52	52	45	20	48	46	43											91	88	87
San Marino	19	9	ii d	9	ú	ç	C	4	Ü	C											) )	4 5	4 1
Slovakia	20 82	98	45	9 4	42	3 6	39	35	32	8 8											5 4	5 4	17
Slovenia	29	51	53	20	48	49	43	42	40	40											18	17	14
Spain	13	15	21	24	56	28	36	24	22	21											17	18	14
Sweden	7	#	6	10	6	∞	8	9	9	7											4	4	4
Switzerland	18	19	18	17	15	15	13	15	18	16											00	∞	∞
Tajikistan TEVD Macadonia	29	92	63	28	22	72	22	26	49	21			30	12	37 40	5 28	36	41	42	46	32	65	68
Turkey	80	85	55	58	54	09	59	56	50	47	42										25	26	3
Turkmenistan	29	22	52	20	51	20	49	28	55	61	63			69	4						84	77	77
Ukraine	52	51	49	48	48	47	45	43	40	39	32	32	35	38							75	82	92
United Kingdom	19	17	15	14	13	12	12	10	10	11	10		11	11	11 1						10	12	11
Uzbekistan	27	29	52	21	48	48	21	51	52	53	46		44	45							69	80	79
Region	4	43	40	39	38	36	36	35	33	32	28	27	29	28	28 3	3 37	7 41	40	43	43	42	43	39

Rate (per 100 000 population) | 2002 | 225 | 226 | 227 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 220 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 171 | 1999 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | 1989 | | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 Country data for Europe: new smear-positive cases, 1993-2002 | 1997 | 1997 | 1997 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | 1998 | | 1996 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 | 173 1 413 3 022 832 504 979 6 955 2 019 6 6 955 2 019 6 65 11 0 469 3 7 512 7 88 3 303 2 605 1 0 42 1 1 497 1 1 497 1 1 487 1 1 487 1 1 487 1 1 483 1 1 483 1 483 1 483 1 483 1 583 1 583 1 583 1 583 1 583 1 583 1 58 8 471 270 7 487 86 4 000 2 072 704 10 385 30 389 129 1 441 681 548 243 303 470 499 484 472 8 314 283 Russian Federation San Marino Serbia & Montenegro snia & Herzegovina rtugal epublic of Moldova urkmenistan

## **Notes**

#### **Belarus**

Treatment outcomes for the 2002 cohort of laboratory-confirmed cases were not included in this annex. Among 2145 cases, 1661 patients were cured, 0 completed treatment, 136 died, 194 failed, 44 defaulted, 0 transferred, and 110 were still on treatment.

#### **Belgium**

Treatment outcomes listed under smear-positive cases are for all laboratory-confirmed cases.

#### **Cyprus**

Data refer only to the Republic of Cyprus, i.e. the northern area is excluded (roughly 40% of the island).

#### Georgia

Regions Afkhazia and South Osetia are not represented in the data.

#### Israel

Treatment and re-treatment outcomes listed under smear-positive cases are based on cohorts of culture-positive cases, where cure is based on culture results.

#### Kazakhstan

Under new smear-positive outcomes, the cohort excluded 185 cases of TB (2% of the cohort) because of resistance to TB drugs, intolerance to TB drugs, deterioration of condition, and loss to follow-up. Under relapse outcomes, 526 cases of TB (18% of cohort) were similarly excluded.

#### Latvia

Outcomes of "transfer-in" patients are matched against "transfer-out" patients at national level to eliminate unknown outcomes because of transfer. Remaining unknowns are given the outcome "default".

#### **Serbia and Montenegro**

Reporting of cases were by site of disease were not available for the entire country. Re-treatment outcomes are from Kosovo only.

#### Spain

The NTP notes that notification data are provisional. "Pulmonary" refers to respiratory cases, while "extrapulmonary" include only meningeal cases.

#### **Sweden**

DOTS coverage was not reported, but assumed to be same as in the previous report.

#### **Tajikistan**

The NTP notes that age and sex data are incomplete.

#### **Turkey**

NTP reported to WHO at the time this report was going to press. Data are therefore not shown in the annexes, and not reflected in results. DOTS coverage in 2003 was reported as 2%. The country had 18 590 TB cases in 2003, and 17 923 WHO notifications (among which, 446 cases were reported under DOTS). Among 4245 non-DOTS new smear-positive cases registered in 2002, 79% were successfully treated.

#### **United Kingdom**

The NTP notes that notification and treatment outcome data are provisional. WHO notifications were revised from 6400 to 6697 cases at the time this report was going to press. Treatment outcome data do not include Scotland.

**EUR** 

**Africa** 

**The Americas** 

**Eastern Mediterranean** 

**Europe** 

**South-East Asia** 

**Western Pacific** 

### **SOUTH-EAST ASIA: SUMMARY OF TB CONTROL POLICIES**

	STATUS <sup>a</sup>	MANUAL	MICROSCOPY	SCC <sup>d</sup>	DOTe	MONITORING OUTCOME <sup>f</sup>
BANGLADESH	DOTS	YES				
BHUTAN	DOTS	YES				
DPR KOREA	DOTS	YES				
INDIA	DOTS	YES				
INDONESIA	DOTS	YES				
MALDIVES	DOTS	YES				
MYANMAR	DOTS	YES				
NEPAL	DOTS	YES				
SRI LANKA	DOTS	YES				
THAILAND	DOTS	YES				
TIMOR-LESTE	DOTS					

SEAR



- a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)
- b Manual: national TB control manual (recommended)
- c Microscopy: use of smear microscopy for diagnosis (core component of DOTS)
- d SCC: short course chemotherapy (core component of DOTS)
- e DOT: directly observed treatment (core component of DOTS)
- $f\quad \hbox{Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)}$

Country data for South-East Asia: estimated burden of TB

	Ė	Incidence, 1990	1990	-	Pre	valence	Prevalence, 1990		Deat	Death, 1990			lnci	Incidence, 2003	003		Preval	Prevalence, 2003			Death, 2003	2003	
	All cases incl. HIV+ New ss+ incl. HIV+	HIV+	lew ss+ incl. F		All cases incl. HIV+ All cases excl.	√ +∧II		HIV+ AI	Il cases incl. HIV+	+ All case	All cases excl. HIV	₹	cases incl. HIV+		New ss+ incl. HIV+	H	All cases incl HIV+	₹	cases excl HIV+	All cases incl HIV+	+AIH i	All cases excl l	+AIH
	number	rate	number	rate	number	rate	number	rate	number rate		number rat	rate	number r	rate	number rate		number rate	e unmper	er rate	number	rate	number	rate
Bangladesh	269 193	246	121 126	111	810 699	741	810 699	741	71342 65		71 342 6	65 3	360 767 2	246	162 331 111	_	719411 490	0 719339	39 490	83 533	22	83 467	22
Bhutan	3 526	208	1 586	94	10 618	929	10618	979	934 55	10	934 6	22	2 492	110	1 121 50	_	4 3 7 8 1 9 4	4 4 3 7 7	77 194	467	21	466	7
DPR Korea	35 493	178	15 972	80	106 891	536	106 891	536	9 4 0 6		9 406 4	47	40 277	178	18 124 80		42 429 187	7 42 428	28 187	3 549	16	3 549	16
India	1 420 446	168	634 211	75	4 263 735	504	4 260 935	503	377 611 45	5 374	964	1 7	788 043 1	168	798 338 75	5 3 085 876	5 876 290	0 3054470	70 287	352 085	33	329 915	31
Indonesia	519 763	285	233 706	128	1 565 313	860	1 565 313 8	860	137 748 76	. 137	748	9 92	627 047 2	285	281 946 128	_	483 735 675	5 1 482 607	07 674	143 178	65	142 168	9
Maldives	388	180	175	81	1 169	542	1 169	542	103 48	3	103 4	48	142	45	64 20	0	125 39		125 39	7	2	7	7
Myanmar	69 260	171	30 847	9/	208 022	514	207 910	513	18 402 45		18 296 4	45	84 546 1	171	37 655 76		92 429 187	7 90 475	75 183	12 410	25	11 748	24
Nepal	39 361	211	17 636	92	118 501	929	118 493 (	989	10 435 56		10 427 5	99	53 139 2	211	23 809 95		80 074 318	8 79 556	56 316	7 399	59	7 138	78
Sri Lanka	10 186	61	4 582	27	30 677	182	30 677	182	2 7 0 1 6		2 700 1	16	11 530	09	5 187 27		16956 89	9 16 948	48 89	1 685	6	1 680	6
Thailand	77 404	142	34 377	63	226 398	416	225 061	414	21 069 39	9 19	805	36	89 351 1	142	39 683 63		30 4 18 208	8 127 792	92 203	12 152	19	10 853	17
Timor-Leste	4 115	556	1 850	250	12 393 1	1 674	12 393 1 6	674	1 091 147	7	1 091 14	147	4 323 5	556	1 944 250		5 8 7 2 7 5 4	4 5 863	63 753	747	96	742	95
Region	2 449 136	190	1 096 068	82	2449136 190 1096068 85 7354416 570 7350160	220		569	650 840 50		646 817 5	50 3 0	3 061 657 1	190 1 3	1 370 201 85	85 5 661 702	702 351	1 5 623 979	79 348	617 211	38	591 734	37

See Explanatory notes, page 151.

I ' I I I I I						ă	DOTS			non-DOTS	STC
Pop   All Cases   Pop   All Cases   Smear-positive   Poin   Smear-positive   Poin   Smear-positive   Point	Re-treatm	Re-treatment cases	Other	Detection rate	%	Notifi	Notifications		% of	Notifications	% of
Country	a- Relapse After failure After defau	After default Other	A	All cases New ss+	ğ	All cases	New ss+	DDR	mlnd	All cases New ss+	wlnd +ss
thousands number rule rate number rate number rules number number rules number rules number rules number number rules number number number rules number numb	nary										
146 736 88 156 88 156 60 53 618 37 56 123 24 9 13 2 2 6 1 2 2 6 1 10 2 6 1 10 2 6 1 2 6	ber number number	number number	number	% %	dod	number rate	number rate	%	cases ss+	number nu	number cases ss+
2 257         1 026         45         380         16         443         294           2 264         51 280         418 10         18         17 32         77         18112           1 085 462         188 754         10 73 065         10         43 27 1         41         419 668         459 47         13           2 19 883         178 260         178 260         81         92 566         42         77 561         77 561           3 18         137         137         43         68         21         0         26           4 945         78 95         75 744         55         5         26006         1           25 164         38 31         30 925         12         14 348         57         8 94           19 065         9477         8 986         47         4 321         23         4764         2 650           778         3247         2560         45         1756         355         1756         34	120 2 505			24 33	66	88 156 60	53 618 3	7 33	89		
22 664         51 280         44 810         184         17 382         77         18112           1 066 462         118 754         1073 065         510         433 271         41         419 688         4594 24         13           219 883         17 8260         17 2500         81         92 566         42         7561         7561           49 845         78 195         77 448         55         26         26         26           49 845         78 3831         30 925         123         44 348         57         894         2650         84           19 055         9477         8 986         47         4 321         23         4 764         2 650         86           68 33         45 04         54 64         356         17 586         17 586         884           778         32 77         7 760         356         177         13 20         17 586	344 38 19	#		41 32	100	1 026 45	360	16 32	99		
1005 462   1188 754   1073 065   101 433 271   41 419 688   459 424   13	606 1 700 1 105	833	7 532	104 96	8	39 396 174	16 445	73 91	49	2 414	947 46
219 883 178 260 178 260 81 92 566 42 77 561 77 561 318 137 137 43 68 21 0 26 26 42 77 561 26 42 77 561 77 561 77 561 77 561 77 561 77 561 77 561 77 574 153 27 484 55 26 006 11 70 065 9477 8 998 47 4321 23 4764 2 650 77 8 77 73 750 750 750 756 756 71 77 77 750 756 756 71 77 77 750 756 756 71 77 77 750 756 756 756 756 756 756 756 756 756 756	253 48 117 12 206	61 295 42 288		60 54	29	836 768 79	372 088	35 47	55	236 297 6	61 183 28
4318         137         137         43         68         21         0         26           4945         78 1945         75 1448         55         26 006         1           25 164         33 831         30 925         12 348         57         894           19 065         9477         8 998         47         4 321         23         4 764         2 650           683         45 64         54 849         45         17 596           778         3247         7 760         35         1077         1320	047 4 086			28 33	86	178 260 81	92 566	42 33	54		
49 465 78 199 75 744 153 27 448 55 26 006 1 26 64 33 831 30 92 5 123 14 348 57 88 94 19 065 9477 8 99 8 47 4 32 1 23 4 764 2 650 62 833 54 504 8 54 89 45 47 560 778 73 750 356 1077 356 120	40 3 0	0		96 106	100	137 43	89	21 106	72		
25 164 33 831 30 925 123 14 348 57 8 894 19 065 9477 8 998 47 4 321 23 4 764 2 650 62 833 54 504 54 504 87 28 459 45 17 596 778 2 277 2 760 355 1077 1320	796 4 494 964	1 487		90 73	92	75 744 153	27 448	55 73	51		
19.065 9.477 8.998 47 4.321 23 4.764 2.650	619 2 064 300	412	2 194	28 60	8	30 925 123	14 348	22 60	62		
62 833 54 504 67 28 459 45 17596 778 3 27 2 760 355 1 107 132 1 240	811 216 47	161	271	78 83	74	7 307 38	3 652	19 70	64	1 69 1	669 53
778 3.217 2.760 355 1.027 13.2	756 1 693			61 72	100	54 504 87	28 459	45 72	62		
25 25 25 25 25 25 25 25 25 25 25 25 25 2	473 20 8	18	431	64 53	78	2 760 355	1 027 132	2 53	45		
Region 1614 648 1 686 837 1 555 385 96 672 878 42 480 998 636 706 180 865	865 64 936 14 649	64 217 42 288	10 428	51 49	12	77 1 314 983 81	610 079	38 45	26	240 402 63	62 799 29

69 55 81 72 72 76 75 75 60 60 22 not eval 12 5 6 6 7 89 4 360 85 1 097 84 078 3 731 8 036 2 663 379 1 990 106 423 28 23 80 0 not eval % trans-ferred default 30 9 5 28 New smear-positive cases - non-DOTS failed died 4 0 7 % compl-eted 9 7 10 17 70 4 88 42 % of notif regist'd 32 88 00 100 100 Number of cases notified regist'd 1 355 43 784 407 654 1 070 4 286 150 698 157 115 407 654 82 Country data for South-East Asia: treatment outcomes for cases registered in 2002 not eval default 9 2 2 0 New smear-positive cases – DOTS

6 % % %

out complstd cured eted died failed det ε 0 ε - 5 81 85 86 86 77 71 71 79 69 81 % of notif regist'd 100 Number of cases

notified regist of a cases

notified regist of a cases

364 380

14 280 14 280

24 15 20 76 280

76 230 76 230

24 162 23 922

13 307 13 307

3 643 3 643

25 559 22 555

10 90 1090 451 162 Bangladesh Bhutan DPR Korea India Indonesia Maldives Myanmar Nepal Sri Lanka Thailand Region

See Explanatory notes, page 151.

Country data for South-East Asia: re-treatment outcomes for cases registered in 2002	ontn-East	Asia:	re-tre	atmer	outc	omes	. TOT 6	ISes re	gister	ed In Zu	70																
				Relap	Relapse - DOTS	.s							After fail	After failure - DOTS	TS						After	After default - DOTS	- DOTS				
	number	%	%	%	%	%	%	%	%	number	%	%	%	%	%	6 %	% %	number	er %	%	%	%	%	%	%	%	
			-ldmoo				trans-	not				compl-			tra	trans- n	not			compl	÷			trans-	not		
	regist'd	cured	eted	died	eted died failed default ferred	default	ferred	eval sı	saccess	regist'd	cured	eted	died f	failed de	default fer	ferred ev	eval success	ess regist'd	d cured	ed eted	died	failed	default	ferred	eval	saccess	
Bangladesh	2485	99	9	4	-	7	2	16	69	365	69	2	4	3	14	4	7 7	72 1510	0 67	7	4	8	15	ю	9	69	
Bhutan	37	65	14	2	11		2		78	19	56	21		2	2	,	42 4	47 29	9 17	14		3	က		62	31	
DPR Korea	638	78	2	2	9	4	က		83	268	69	9	œ	œ	4	4	_	76 191	11 76	3 7	9	3	4	က		83	
India	34 317	73	3	7	9	12	-		75	8 727	22	က	∞	15	16	_	9	60 41 034	4 67	4	7	5	16	-	0	71	
Indonesia																											
Maldives	က	100							100	-				100													
Myanmar	3 762	29	6	8	3	6	3		9/																		
Nepal	1 898	83	2	7	က	က	3	0	84	300	71	~	7	14	2	2	0 7	71 465	5 75	3	8	2	00	3	0	79	
Sri Lanka	208	62	2	6	2	20	2		29	30	43	10	13	13	17		3	53 141	.1 45	9	6	_	38	~	-	20	
Thailand	1 416	28	7	17	9	8	2		64	574	49	9	17	6	12	7	4)	22									
imor-Leste									1																		
00000	44 764	44 764 72 3 7 E 11	۲,	7	Ľ	-	-	-	75	10 284	8	~	o	14	7	-	•	61 43 370	67	7	_	Ľ	16	-	•	7.4	

See Explanatory notes for previous table, page 151.

Country data for South-East Asia: trends in DOTS treatment success and detection rates, 1994–2003

		DOT	S new sn	near-pos	DOTS new smear-positive treatment success	ment suc	(%) sseo:				DOT	S new sr	near-pos	itive cas	DOTS new smear-positive case detection rate		(%)	
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
Bangladesh	73	71	72	78	80	8	83	8	84	7	14	18	23	23	23	25	53	33
Bhutan	71	26	96	82	06	82	06	93	98	28	24	23	22	26	29	31	32	32
DPR Korea					91	8	91	91	88					2	56	53	79	91
India	83	79	6/	82	84	82	84	82	87	0	1	_	2	7	12	24	31	47
Indonesia	94	91	81	72	28	20	87	98	98	~	2	7	12	18	19	20	27	33
Maldives	98	26	93	8	8	8	26	26	92	96	97	94	94	101	81	79	87	106
Myanmar		99	62	82	82	81	82	81	81		56	26	29	32	48	99	65	73
Nepal			82	87	88	87	98	88	98		9		16	44	26	99	22	09
Sri Lanka	77	79	80	9/	92	8	77	80	81	63	61	71	75	75	89	73	71	70
Thailand			78	62	89	77	69	75	74		0	2	21	39	46	73	9	72
Timor-Leste								73	81								29	23
Region	80	74	77	72	72	73	83	84	82	2	4	9	00	41	18	27	33	45

SEAR

Country data for South-East Asia: age and sex distribution of sme	h-East A	sia: age	and se	x distrik	ution o	r smear-	positive	er-positive cases in DOTS areas, 2003 (absolute numbers)	DOTS	areas, 2	003 (ab	solute n	umbers	_							
				MALE						۳	FEMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+69	0-14	15-24	25-34	35-44	45-54	55-64	+59
Bangladesh	320	5 166	7 275	8 0 2 8	6 947	5 501	4 142	544	4 298	4 282	3 2 5 8	2 086	1 150	591	864	9 4 6 4	11 557	11 316	9 033	6 651	4 733
Bhutan	6	62	20	20	25	20	13	14	24	39	17	13	15	9	23	119	88	37	38	35	19
DPR Korea	81	1 101	2 173	2 541	2 340	1 327	299	06	792	1 542	1 531	1316	723	326	171	1 893	3715	4 072	3 656	2 050	888
India	1 890	42 830	54 948	56 283	47 204	30 256	16 242	4 120	31 332	31895	19 662	11 520	6 903	3 379	6 0 1 0	74 162	86 843	75 945	58 724	37 159	19 621
Indonesia	532	9 570	12 647	10 925	9 558	6 720	3 615	809	8 734	10 127	7 889	6 085	3 907	1 649	1 140	18 304	22 774	18 814	15 643	10 627	5 264
Maldives	_	14	7	4	6	6	4	0	00	2	_	2	_	0	_	22	12	2	14	10	4
Myanmar	107	2 536	4 408	4 427	3 269	1 974	1 296	154	1 781	2 442	2 003	1 491	943	617	261	4 3 1 7	6 850	6 430	4 760	2 917	1 913
Nepal	122	2 039	1 658	1619	1 769	1 639	735	189	1 283	1 107	873	609	486	220	311	3 322	2 7 6 5	2 492	2 378	2 125	922
Sri Lanka	1	286	399	609	999	421	315	12	250	181	148	149	103	103	23	536	580	757	814	524	418
Thailand	41	1 636	4 6 1 5	4 259	3 497	2 740	3 241	49	944	1 678	1 350	1 279	1 264	1 866	06	2 580	6 293	2 609	4 776	4 004	5 107
Timor-Leste	2	130	135	107	86	99	41	13	86	116	92	92	43	17	18	228	251	183	174	109	28
Region	3 119	65 370	88 315	88 852	75 381	50 673	30 206	5 793	49 577	53 414	36 808	24 629	15 538	8 774	8 912	114 947	141 729	125 660	100 010	66 211	38 980

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

Country data for South-East Asia: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers)	h-East As	sia: age	and sex	distrib	ıtion of	smear-p	ositive	cases in	non-DC	JTS area	ıs, 2003	(absolu	te numb	ers)							
				MALE						뿐	FEMALE						_	ALL			
	0-14	0-14 15-24	25-34 35-44	35-44	45-54	55-64	+69	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14 15-24	15-24	25-34 35-44		45-54	55-64	+59
Bangladesh Bhutan																					
DPR Korea	2	53	106	137	129	85	72	3	31	81	92	79	46	44	8	84	187	213	208	131	116
India	521	4 421	6 810	7 304	5 661	3 483	1 776	625	3 179	4 422	3 658	2 535	1 419	909	1 146 7 600	2 600	11 232	10 962	8 196	4 902	2 382
Indonesia	_																				
Maldives																					
Myanmar																					
Nepal	_																				
Sri Lanka	-	25	68	85	126	74	74	2	52	37	38	38	29	17	3	80	105	123	164	103	91
Thailand																					
Timor-Leste																					
Region	527	4 499	6 984	7 526	5 916	3 642	1 922	630	3 265	3 265 4 540 3 772 2 652 1 494	3 772	2 652	1 494	299	1157 7764 11524 11298 8568	7 764	11 524	11 298		5 136	2 589

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

SEAR

Country data for South-East Asia: smear-po	outh-Ea	st Asia	: smea		tive no	tificati	on rat	ositive notification rates (per 100 000 population)	100 00	od oc	ulation	) by aç	by age and sex,		2003						
			_	MALE						ш	FEMALE							ALL			
	0-14	15-24	0-14 15-24 25-34 35-44	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	+59
Bangladesh	-	34	62	93	123	183	178	2	30	39	40	39	37	24	2	32	51	89	82	109	100
Bhutan	2	27	33	19	33	38	59	က	25	26	17	17	27	12	2	56	30	18	25	32	19
DPR Korea	က	64	115	148	227	135	100	က	48	85	92	129	89	43	ო	26	100	120	178	100	29
India	-	45	71	91	106	108	70	3	36	46	36	30	56	14	2	41	29	65	69	29	40
Indonesia	2	44	29	74	94	108	71	2	4	54	25	09	26	56	2	43	09	64	77	80	46
Maldives	-	41	32	22	90	152	69	0	54	24	7	49	19	0	-	33	28	17	69	88	36
Myanmar	1	52	106	147	151	149	123	2	37	28	64	9	99	49	2	45	82	105	107	106	83
Nepal	2	80	06	122	198	276	169	4	72	63	29	89	78	44	က	29	77	92	133	174	102
Sri Lanka	_	17	30	47	89	63	28	~	18	15	14	19	21	9	_	17	22	31	46	44	38
Thailand	-	29	83	93	107	137	194	-	17	59	27	36	29	88	1	23	26	29	20	96	135
Timor-Leste	က	131	265	224	300	331	363	6	111	319	168	248	211	142	9	121	287	197	275	270	250
Region	-	44	72	91	109	117	85	က	35	47	4	38	35	22	2	40	09	29	75	92	51

Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available.

Country data for South-East Asia: number of TB cases notified, 1980-2003	or South-	East As	ia: num	ber of 1	B case	s notifie	d, 1980.	-2003																
	1980	1981	1982	1983	1984	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003
Bangladesh	39 774	42 644	49 870	52 961	45 679	41 802	45 599	45 355	44 280	45 191	48 673	56 052	31 400	54 001	48 276	56 437	63 471	63 420	72 256	79 339	75 557	76 302	81 963	88 156
Bhutan	1 539	2 657	720	1 017	904	1 073	1 582	809	1 126	1 525	1 154	966	140	108	1 159	1 299	1 271	1 211	1 292	1 174	1140	1 037	1 089	1 026
DPR Korea	0								0									11 050	1 152	12 287	34 131	29 284	40 159	41 810
India	705 600	705 600 769 540	923 095 1 075 098 1 109 310 1 168 804 1 279 536	1 075 098	1 109 310 7	168 804 1	1 279 536 1	403 122 1	1 457 288 1	1 510 500 1 519 182		555 353 1	1121 120 1 081 279 1 114 374 1 218 183	181 279 1	114 374 1		1 290 343 1 1	132 859 1 1	102 002 1	1 218 743 1	1157181	085 075 1	060 951 1	073 065
Indonesia	25 235	32 461	33 000	31 809	32 432	17 681	16 750		97 505	105 516	74 470	808 09	98 458	996 29	49 647	35 529	24 647	22 184	40 497	69 064	84 591	92 792	155 188	178 260
Maldives	73	112	111	143	123	91	111	115	82	203	152	123	92	175	249	231	212	173	176	153	132	139	125	137
Myanmar	12 744	12 461	12 744 12 461 12 069	11 012 11 045		10 506	10 840	11 986	9 348	10 940	12 416	14 905	17 000	19 009	15 583	18 229	22 201	17 122	14 756	19626	30 840	42 838	57 012	75 744
Nepal	1 020	337	1 459	200	190	52	252	1 012	1 603	11 003	10 142	8 983		13 161	15 572	19 804	22 970	24 158	24 135	27 356	29 519	29 519	30 329	30 925
Sri Lanka	6 212	6 288	7 334	9999	6 376	5 889	9659	6 411	6 092	6 4 2 9	9999	6 174	6 802	6089	6 132	5 710	5 366	6 542	6 925	7 157	8 413	7 499	8 939	8 998
Thailand	45 704	49 452	48 553	65 413	69 240	77 611	52 152	51 835	50 021	44 553	46 510	43 858	47 697	49 668	47 767	45 428	39 871	30 262	15 850	29413	34 187	49 656	49 581	54 504
Timor-Leste																							2 760	2 760
Region	837 901	915 952	837 901 915 952 1 076 211 1 244 819 1 275 299 1 323 509 1 413 418 1	1 244 819 1	1 275 299 1	323 509 1	1 413 418 1	520 444 1	667 348 1	735860 1	520 444 1 667 348 1 735 860 1 719 365 1 747 252 1 322 709 1 287 176 1 288 759 1 400 850 1 470 352 1 308 981 1 279 041 1 464 312 1 414 228 1 414 141 1 485 366	747 252 1	322 709 1	287 176 1 2	298 759 1	100 850 1	170 352 1 3	108 981 1	279 041 1	464 312 1	414 228 1	414 141 1	485 366 1	1 552 625
number reporting	0	00	80	80	80	80	00	7	6	00	80	80	7	80	00	00	80	6	6	6	6	6	10	7
	007	0	0	0	c	0	0	1	007	0	0	6	1	0	0	0	0	00	000	000	007	007	007	007

SEAR

Country data for South-East Asia: case notific	South-	East A	sia: c	ase no		ation rates (per 100 000 population), 1980–2003	ed) se	r 100 (	000 pc	pulati	on), 1	980–2(	003											
	1980	1981	1982	1983	1984 19	1985	1986	1987	1988	1989	1990	1991	1992	1993	1994	1995 ′	1996	1997	1998	1999	2000	2001	2002	2003
Bangladesh	47	49	99	28	48	43	46	45	43	42	44	20	27	46	40	46	20	49	22	29	22	25	22	09
Bhutan	117	197	52	72	62	72	104	39	20	92	89	28	80	9	9	72	69	49	99	29	22	49	20	45
DPR Korea									0									51	2	99	153	131	178	184
India	102	109	129	147	148	153	164	176	179	182	179	180	127	121	122	131	136	117	112	122	110	105	101	101
Indonesia	17	21	21	20	20	1	10		22	29	4	33	25	33	56	18	12	7	20	33	40	43	71	81
Maldives	46	69	99	83	69	20	29	29	42	6	20	22	40	74	102	95	82	92	64	54	45	46	40	43
Myanmar	38	36	34	31	30	28	59	31	24	27	31	36	41	45	36	41	20	38	32	42	65	68	117	153
Nepal	7	2	6	4	~	0	_	9	6	09	24	47		99	9/	92	107	110	107	119	126	123	123	123
Sri Lanka	43	43	49	4	41	38	42	40	37	39	40	36	33	39	35	32	30	36	38	39	45	40	47	47
Thailand	66	105	101	134	139	153	101	66	92	83	98	80	82	88	84	79	89	51	27	49	99	81	80	87
Timor-Leste																							374	355
Region	80	85	98	11	11	113	119	125	13	137	133	133	66	94	93	66	102	88	98	96	92	06	94	96

Country data for South-East Asia: new smear-positive cases, 1993-2003

					Nun	Number of cases	Se								œ	ate (per 10\	Rate (per 100 000 population)	lation)				
	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003
Bangladesh	18 993	1 710	20 524	29 674	33 117	37 737	37 821	38 484	40 777	46 811	53 618	16	-	17	23	56	29	28	28	59	33	37
Bhutan		352	367	308	284	270	315	347	359	364	360		20	20	17	15	14	16	17	17	17	16
DPR Korea					3 980	403	5 073	16 440	14 429	18 576	17 392					18	2	23	74	64	82	77
India	225 256	226 543	264 515	290 953	274 877	278 275	345 150	349 374	384 827	395 833	433 271	25	25	28	31	28	28	35	34	37	38	41
Indonesia	62 966	49 647	31 768	11 790	19 492	32 280	49 172	52 338	53 965	76 230	92 266	33	56	16	9	10	16	24	25	25	35	42
Maldives	126	125	114	106	96	88	88	65	29	09	89	53	51	46	41	36	32	31	22	20	19	21
Myanmar			8 681	9 7 1 6	9 695	10 089	11 458	17 254	21 161	24 162	27 448			20	22	21	22	24	36	44	49	22
Nepal	6 6 6 7 9	10 442	8 591	10 365	11 323	11 306	13 410	13 683	13 683	13 714	14 348	33	51	41	48	52	20	28	28	22	26	22
Sri Lanka	3 335	3 405	3 049	2 958	3 506	3 761	3 911	4 314	4 316	4 297	4 321	19	19	17	16	19	21	21	23	23	23	23
Thailand		20 260	20 273	16 997	13 214	7 962	14 934	17 754	28 363	25 593	28 459		35	35	29	22	13	25	29	46	41	45
Timor-Leste										1 090	1 027										148	132
Region	317 355	312 484	357 882	372 867	369 583	317 355 312 484 357 882 372 867 369 583 382 171 481	481 332	510 053	561 939	606 730	672 878	23	22	25	56	25	26	32	33	36	38	42

## **Notes**

#### **Bangladesh**

There is a discrepancy between the population estimate used by the national TB programme (137 059 519) and that used by the UN (146 736 131). Using the country's population estimate, there would be 336 976 estimated TB cases (of which, 162 331 smear-positive cases), and DOTS detection of smear-positive cases would be 35% (instead of 33%).

#### **Bhutan**

Estimates of the population vary widely, from 800 000 to over 2 million. The UN population estimate is 2.2 million.

#### India

DOTS coverage reached 73% by the end of 2003, but it was 67% calculated on a quarter-by-quarter basis (starting from 60% in the first quarter). Cases notified under DOTS included 31 341 cases (of which 13 603 were new smear-positive) in patients receiving non-rifampicin regimens.

#### **Myanmar**

DOTS coverage was 95% in 2003, but reached 100% by end of the year.

#### Nepal

Data refer to a mid-July to mid-July calendar.

#### **Thailand**

Data refer to an October-September calendar.

#### **Timor-Leste**

DOTS coverage was not reported, but assumed to be same as in the previous report.

**Africa** 

**The Americas** 

**Eastern Mediterranean** 

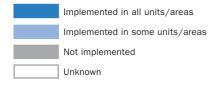
**Europe** 

**South-East Asia** 

**Western Pacific** 

## **WESTERN PACIFIC: SUMMARY OF TB CONTROL POLICIES**

						MONITORING
AMEDICANI CAMOA	STATUS <sup>a</sup>	MANUAL <sup>b</sup> YES	MICROSCOPY°	SCC <sup>d</sup>	DOTe	OUTCOME
AMERICAN SAMOA	DOTS	YES				
AUSTRALIA	DOTS	YES				
BRUNEI DARUSSALAM	DOTS					
CAMBODIA	DOTS	YES				
CHINA	DOTS	YES				
CHINA, HONG KONG SAR	DOTS	YES				
CHINA, MACAO SAR	DOTS	NO				
COOK ISLANDS	DOTS	YES				
FIJI	DOTS	YES				
FRENCH POLYNESIA	DOTS	YES				
GUAM	DOTS	YES				
JAPAN	DOTS	YES				
KIRIBATI	DOTS	YES				
LAO PDR	DOTS	YES				
MALAYSIA	DOTS	YES				
MARSHALL ISLANDS	DOTS	YES				
MICRONESIA	DOTS	YES				
MONGOLIA	DOTS	YES				
NAURU	DOTS	NO				
NEW CALEDONIA	DOTS	YES				
NEW ZEALAND	DOTS	YES				
NIUE	DOTS	YES				
NORTHERN MARIANA IS	DOTS	YES				
PALAU	DOTS	YES				
PAPUA NEW GUINEA	DOTS	YES				
PHILIPPINES	DOTS	YES				
REP. KOREA	DOTS	YES				
SAMOA	DOTS	YES				
SINGAPORE	DOTS	YES				
SOLOMON ISLANDS	DOTS	YES				
TOKELAU	NON-DOTS	NO				
TONGA	DOTS	YES				
TUVALU	NON-DOTS	NO				
VANUATU	DOTS	YES				
VIET NAM	DOTS	NO				
WALLIS & FUTUNA IS	DOTS	YES				



a Status: DOTS status (bold indicates DOTS introduced in 2003. Blank indicates no report received)

b Manual: national TB control manual (recommended)

c Microscopy: use of smear microscopy for diagnosis (core component of DOTS)

d SCC: short course chemotherapy (core component of DOTS)

e DOT: directly observed treatment (core component of DOTS)

f Outcome monitoring: monitoring of treatment outcomes by cohort analysis (core component of DOTS)

Country data for the Western Pacific: estimated burden of TB

,	-	Incidence, 1990	3. 1990	r	ď	revaler	Prevalence, 1990			Death. 1990	1990	F	lncic	ncidence. 2	2003		Prevalence.	ance, 2003		De	Death, 2003		Г
	All cases incl.	+AIIA	All cases incl. HIV+ New ss+ incl. HIV+		All cases incl.	+ HIV+	All cases incl. HIV+ All cases excl	HIV+	All cases incl.	HIV+	All cases excl.	+ / / / /	All cases incl. HI	HIV+ Ne	New ss+ incl. HIV+	'+ All cases	.⊑	- All cases excl	cl HIV+	All cases incl	HIV+ All ca	All cases excl H	+   
	number	rate	number	rate	number	rate	number	rate	number	rate	number	rate	number ra	rate	number rate		number rate	number	rate	number rat	e	number r	rate
American Samoa	24	25	1	23	89	145	89	145	5	11	5	7		30	8 1.	3	34 54	34	54	4	9	4	9
Australia	1 067	9	478	က	1 195	7	1 195	7	171	_	171	_	1 128	9				1132	9	113	_	111	_
Brunei Darussalam	159	62	72	28	443	172	443	172	33	13	33	13	195	24	88	2	217 61	217	61	17	2	17	2
Cambodia	56 202	277	24 848	255		1 584	153 874	$\overline{}$	12 009	123	11 593	119	830	208	31 758 225		107 836 762		742		95	11 469	81
China	1 345 828	116	605 046	25	3 748 967	325	3 748 967	325	282 443	24	282 443	24	1 334 066 1	05	758	က	203 059 246	3 200 204	245	235 864		233 918	9
China, Hong Kong SAR	6 326	111	2 845	20	17 611	309	17 608	309	1 329	23	1 327	23	5 439	77	2 446 35		5 554 79	5 548	79	468	7	467	7
China, Macao SAR	348	93	156	42	696	260	696	260	73	20	73	20	382	82	172 3		414 89	413	88	45	10	4	10
Cook Islands	10	25	4	23	27	145	27	145	2	7	2	7	2	30	2 1.	3	11 59	1	29	-	9	_	9
III	376	25	169	23	1 047	145	1 047	145	62	7	79	7	249	30	112 13	3	318 38	317	38	38	4	37	4
French Polynesia	102	25	46	23	283	145	283		21	11	21	11	73	30	33 13	3	94 39	94	38	11	2	11	2
Guam	189	141	85	63	525	392	525	392	40	30	40	30		09	44 27	_	171 105		105	19	12	19	12
Japan	64 322	52	28 927	23	71 939	28	71 895		10 323	8	10 313	80	39 927	31	17 956 14			53 154	42	4 973	4	4 960	4
Kiribati	101	141	45	63	282	392	282		21	30	21	30		09			53 60		09		4	4	4
Lao PDR	7 376	178	3 318	80	20 547	497	20 547	•	1 548	37	1 548	37		157	3 999 71					1 482	26	1 475	56
Malaysia	21 380	120	9 591	25	59 540	334	59 536	334	4 488	25	4 485	52	25 785 1	90	11 567 47		33 128 136	32			17	3 942	16
Marshall Islands	63	141	28	63	174	392	174		13		13	30		09				32	09		4	2	4
Micronesia	136	141	61	63	378	392	378	392	28	30	28	30	99	09	29 27	_	68 62			9	9	9	9
Mongolia	4 875	220	2 194	66	13 581	613	13 581	613	1 023		1 023	46	5 025 1	94			6 149 237	9		834	32	833	32
Nauru	5	25	2	23	14	145		145	-		1	11		30		3			36	0.5	4	0.5	4
New Caledonia	241	141	108	63	671	392			51	30	51	30	137	09	61 27	_	236 103	235	103	56	7	56	7
New Zealand	356	11	160	2	398	12	398		22	2	22	2		11		2			11	41	1	41	-
Niue	1	25	0.5	23	3	145			0.3	11	0.3	11	9.0	30	0.3	3		1	29	0.1	9	0.1	9
Northern Mariana Is	62	141	28	63	172	392			13	30	13	30		09	21 27	_	53 67	. 23	29	9	80	9	∞
Palau	21	141	10	63	09	392	09	392	4	30	4	30	12	09	6 27				76	2	8	2	œ
Papua New Guinea	10 992	267	4 925	120	30 621	744	30 621		2 307	99	2 307	26		235	6 020 105		30 240 529	30 108		2 782	49	2 700	47
Philippines	205 495	336	92 457	151	572 431	937	572 431	-	43 126	71	43 126	71	885	96							49	38 811	49
Rep. Korea	36 762	98	16 539	39	102 406	239	102 406	``	7715	18	7 715	18		87			56 522 118	26 4		4 715	10	4 690	10
Samoa	83	25	37	23	232	145	232	_	17	7	17	<del>-</del>		30	24 13		79 44		44	6	2	တ	2
Singapore	1 763	28	789	56	1 967	65	1 963		283	<b>о</b>	282	6	1 749	41	783 18			_	42	198	2	193	2
Solomon Islands	449	141	202	63	1 250	392	1 250	ľ	94	30	94	30	286	09		_			09	21	4	21	4
Tokelau	8.0	52	0.4	23	2	145	2		0.2	11	0.2	7	0.5	30	0.2	3	0.9 59		29	0.1	9	0.1	9
Tonga	52	25	23	23	144	145	144	`	1	7	1	=	31	30	14 13	3	45 44	45	44	2	2	2	2
Tuvalu	2	25	2	23	13	145	13	1	1.0	11	1.0	7	က	30	1	3	6 55		59	9.0	9	9.0	9
Vanuatu	210	141	92	63	586	392	586		4	30	4	30		09	57 27				71	18	00	18	ω
Viet Nam	133 660	202	59 924	91	372 326	563	372 326	263	28 051	42	28 051	42	144 942 1	178	64 982 80	194	970 2	193 762	238	18 750	23	18 116	72
Wallis & Futuna Is	7	25	က	23	20	145	20	145	-	1	-	=	4	30	2 1.	3	4 30	4	30	0.3	2	0.3	7
Region	1 899 047	125	853 229	26	5 175 246	341	5 174 709	341	395 426	26	394 995	26	1 933 054 1	112	868 388 50	4	081 006 236	4 073 485	235	326 862	6	321 957	6
				3		,				2				!		1							2

See Explanatory notes, page 151.

								Whole country	untry							L		DOTS					non-DOTS	
						Ne	New cases			ک ا	Re-treatment cases	ases	Other		Detection rate	┝		Notifications	ons		% of		Notifications	yo %
	Pop	All	All cases	Smea	Smear-positive		Pulm Smear-negative		Extra- Re	Relapse After	After failure After default	default Other	 	i	All cases New ss+	`₽	All cases	ses	New ss+	DDR	A pulm	All cases	New ss+	mlnd
	apacanoq+	Country	WHO	oter oter	roter	confirmed	or	ď	pulmonary	rodmin	roden	redmina redmina	hor	%	76	Š	dania	40	rodania	% ofcr	100 00000	- to		too oooo rodmin
	COCCOMING	i di														Т	2	1		1	١,			00000
American Samoa	62	8 00	e 6	2 4	24 5	m +	0 8	0 0	104	ć		Ω		~ ∂			E 0		2 9					
Australia Brunei Danissalam	378	247				- 25	36 136	310	4 CA	27	c			ء 2 ج								228	60	n n
Ombodio	330	20 206		4	1			4 30 7		76.4	2 2	- 00				+	ac							
China	1 304 106	20 300		C						70 as as	0		8 27 700	18			4							
China. Hong Kong SAR	7 049	6 035		•		25 23 399			594	319	0	1 6/	549	103	3 73	100	.,	63	1 426	20 58	38 22	1 163	353	36
China, Macao SAR	464	382			138 3					32	0	9				-		1		1		L		
Cook Islands	18	0	0	0	0	0	0		0	0	0	0		J										
ifi	839	179	179	21	70	8		49	55	2				7'5			179				63 59	6		
French Polynesia	244	51		20	21		36		2	2				1 6								3		
Guam	163	61		14	0	0	39	15	7	0	0	0		2										
Japan	127 654	31 638	31 638		10843	173		13 621	6 160	1 014				7								4 10 288	3 631	46
Kiribati	88	278	284 33	324	99 11	113		7.1	110	4				54		H						3		
Lao PDR	5 657	2 824	2 780			33 2 277	77	495	317	86	2	39		è								6		
Malaysia	24 425	15 912	15671	64 7.8	7 989 3	33		5 811	1 465	406	20	175	4	46 61								3		
Marshall Islands	53	09			20 3		20	22	18	0	0	0		18							10 48	3		
Micronesia	109	109		91		25	41		21	0	0	က		7 15.								2		
Mongolia	2 594	4 007	3 918 1		1541 5	59 1 541	41	812	1 419	146	58	31		₹		_						2		
Nauru	13	3				8	_		-	0	0	0		7.										
New Caledonia	228	36	36	16	4	9	19		12	_				7						6 23	23 61	_		
New Zealand	3 875	424					181	125	136	19			e)	38 94		$\dashv$		- 1		- 1		2		
Niue	2	0		0			0		0	0	0	0		_										
Northern Mariana Is	79	45		22	16 2	20	30		10	0	0	0		ര്			45				75 46	2		
Palau	20	6							2	0	0	0				+								
Papua New Guinea	5 711								6 769	1 367			724								12	12 798	3 2 3 1 0	32
Philippines	/9 999	134 375	134 375 16	74 726	0/97/	91 /26/0		55 942	1 693	4 070	C	7										7	0	7
Nep. Noted	4, 700	40.300				2			1312	3 20 1	900	2	ň			+						1		
Singaporo	4 253	1 500		2 00		- 4	047	0 12	173	2 6	o ¢	o 4		n à								2 B D B D B D B D B D B D B D B D B D B	346	7
Solomon Jelande	4 233	303					130	107	2 5	<u> </u>	<u> </u>	2 0		9. 0						•				
Tokelau	2	0					9 0		2			0				+	0							
Tonda	1 401	•	, 4	. 4	. 1	, 1	. 1	) e <sup>e</sup>	0 0	0 0				ć				, <del>(</del>		11.8	80 79			
Tuvalu	=======================================	32		283			. 0	17	ı ∞	OI (	· <del>-</del>	0		1 95%	0							30	0	0
Vanuatu	212	106	104	49	40 1	19	40	45	18	1	-	0		1 82								7		
Viet Nam	81 377	93 421			55 937 6			16 791	14 564	5 449	491	189			4 86	100	92 741	114	55 937	98 69	27 27	_		
Wallis & Futuna Is	15	18	15 10	102	7 4	48	11	7	0	-	0	-		2 343		$\dashv$	15					C		
Region	1 732 104 1 045 572	1 045 572	987 927	57 454 732		26 401 765		376 679 7	70 506	85 982	866	1 208 67 709	7 008	38 51	1 52	06	879 827	7 51	431 396	25 50	50 58	108 100	23 336	27
					l													l	l	ı		1		

See Explanatory notes, page 151.

Country data for the Western Pacific: treatment outcomes for cases registered in 2002

				1	141		E				-					1		S.F.					1		-140	44	1	E		
_			- 1	w smea	New smear-positive cases - DOLS	e cases	200				1			New Sh	New smear-positive cases - non-DOLS	TIVE Cast	-uou - sa	200		1			ő	smear-positive re-treatment cases - DOLS	SITIVE re	-treatme	nt cases	200-	1	1
			, %	%	, %	%	%	%	%	° .	%			%	%	%	%	%	% .		%		%	%	%	%	%	° .	% .	%
•	Number of cases notified regist	r cases regist'd	of notif	cured	compl- eted	died	failed	trans- failed default ferred		not eval su	success	notified regist	Į,	of notif regist'd cu	cor cured et	compl- eted died	ed failed	d defa	trans- default ferred	eval	snccess	Number regist'd	cured	compl-	died	failed	default	ferred		snccess
American Samoa	-	-	100		100						100																			
Australia	127	180	142	21	22	10		_	9	4	78	83	139	167	6	68 1	1		12		77	17	24	47	18			12		7
Brunei Darussalam	112	22	69	09	25	13		3			84											10			20		10			20
Cambodia	17 258	17 396	101	68	က	4	0	2	-		92											875			9	-	က	-		68
China	180 239	180 239	100	06	က	_	_	~	2	က	93		13 681	93	82	7	_	_		2	92	46 932	83	2	က	4	2	_	2	88
China, Hong Kong SAR	1 501	1 529	102	71	00	2	œ	က	2		62	389	363	93	0	_	_	0	0 98		-	246			7	12	6	4		89
China, Macao SAR	135	138	102	87	2	2		-	2	2	68	12	12	100		ľ	75	25				47	62	26	4		4	4		87
Cook Islands	_	_	100	100							100																			
ii.	75	73	46	82		2		80	_		85											2	100							100
French Polynesia	28	28	100		82	7		11			82											80		75	52					75
Guam	31	28	06	89		25			7		89											8	100							100
Japan	6 172	6 602	107	25	24	12	4	2	9		92	4 635	2 985	64	17	16	2	4 0	09		8	743		24	7	7	က	00		7.1
Kiribati	82	82	100	87	7	4	2				94											e	33							100
Lao PDR	1 829	1 738	92	29	7	7	0	7	4		78											117	72		6	7	6	က	9	99
Malaysia	7 958	7 424	93		9/	œ		1	2		92																			
Marshall Islands	18	20	111	06	10						100																			
Micronesia	22	22	100	91					6		91											3			33		33		33	
Mongolia	1 670	1 671	100	83	4	3	2	3	3	0	87											226	22	15	8	11	7	2	0	69
Nauru	2	2	100	20		20					20																			
New Caledonia	21	20	92	45	40	2		10			82											_	100							100
New Zealand	88	93	106		09	2			3	31	09											3		29					33	67
Niue	-	2	200	20	20						100																			
Northern Mariana Is	21	21	100	71		2			24		71																			
Palau	6	8	88	38		25				38	38																			
Papua New Guinea	926	930	100	39	14	က	-	16	2	22	53		1 345							100		82	56	26	12	6	16	12		21
Philippines	65 148	59 453	91	1	= 1	ი .	- 1	2	ი :	0	88	!													•	•		1		
Kep. Korea		4 / 43		200	7	-	7	20	10	-	83	11 345										9/G L	90	7	7	7	٥	20	7	88
Samoa	19	19	100	8		7			2		84																			
Singapore	311	449	144		87	7		9	0		87	238	449	189		65 2	29	_	0 9		65	87		20	=	_	17			20
Solomon Islands	108	108	100	71	19	4		2	2		90											2	100							100
Tokelau																														
Tonga	23	24	104	75	∞	17					83											_	100							100
Tuvalu													8			25 2	25	25		25	25									
Vanuatu	31	88	123	63	16	7	2	က	က		62	7										2								100
Viet Nam Wallis & Futuna Is	56 698 1	56 590 5	100	8 Q	2	m	-	<del>-</del>	2		100											6 0 79	83	9	2	2	7	2	17	8 8
Region	340 666	339 754	100	8	9	2	-	2	2	2	9	31 442	18 982	09	64	10	,	2	12	6	74	57 071	25	9	က	4	2	2	2	87

See Explanatory notes, page 151.

100 % not eval % trans-ferred died failed default After default – DOTS % % % % compl-eted cured 100 regist'd saccess % not eval default After failure - DOTS died failed compl-eted cured regist'd 77 07 19 87 80 % not eval % trans-ferred default ∞ Relapse – DOTS failed died 20 compl-eted œ 67 12 cured 81 80 50 88 54 17 10 807 5 446 9 149 number regist'd 3 China, Hong Kong SAR China, Macao SAR Cook Islands Northern Mariana Is Papua New Guinea Brunei Darussalam Wallis & Futuna Is Japan Kiribati Lao PDR Malaysia Marshall Islands Singapore Solomon Islands American Samoa French Polynesia Nauru New Caledonia New Zealand Philippines Rep. Korea Micronesia Cambodia Mongolia Tuvalu Vanuatu Viet Nam Australia okelan Fonga Guam

Country data for the Western Pacific: re-treatment outcomes for cases registered in 2002

See Explanatory notes for previous table, page 151

Country data for the Western Pacific: trends in DOTS treatment success and detection rates, 1994–2003

,		100		1	1		1707		-			H			1	0/ -1-		
		2	S new sn	DO I S new smear-positive treatment success (%)	live treatr	nent succ	(%) ssac					DOIS ne	w smear-	positive	case dete	DOIS new smear-positive case detection rate (%)	_	
	1994	1995	1996	1997	1998	1999	2000	2001	2002	1995	1996	1997	1998	1999	2000	2001	2002	2003
American Samoa		100			20	100	100	100	100			64		34	23	23	12	24
Australia				99	75	84	74	99	78				23	30	24	20	25	6
Brunei Darussalam					85	9/	63	26	84					123	100	111	129	138
Cambodia	84	91	94	91	92	93	91	92	92	40	34	44	47	53	49	47	55	09
China	94	96	96	96	26	96	92	96	93	15	28	32	32	59	31	31	30	43
China, Hong Kong SAR					82	78	9/	78	62					09	29	22	09	28
China, Macao SAR	75			81		78	88	98	68	98	155	194	163		93	91	79	9/
Cook Islands				20		80		100	100				32			75	39	
Fiji	06	98		87	06	92	85	82	85	47	49	49	26	51	20	61	65	63
French Polynesia		29	98	100	74	85	26	80	82		94	107	91	91	82		84	99
Guam						94	93	71	89						82	26	29	
Japan						9/	20	75	92						22	59	33	40
Kiribati					83	88	91	98	94			34	170	203	196	244	329	419
Lao PDR		70	55	62	75	84	82	77	78		24	33	40	45	40	40	46	47
Malaysia		69				06	78	6/	92	64	69				73	73	69	69
Marshall Islands					83	82	91	98	100				29	96	99	8	119	140
Micronesia	64	80				92	93	100	91	18	30				43	24	71	95
Mongolia	29	78	78	86	84	98	87	87	87	7	31	31	25	29	62	73	74	89
Nauru						20	25	100	90						216	110	112	25
New Caledonia	62	75			20	77	88	8	85	24	31			30	28	28	33	23
New Zealand							30	6	09						41	37	48	22
Niue									100								364	
Northern Mariana Is						80	81	74	71						117	82	96	75
Palau	64	29	75					100	38	115	53	97					156	06
Papua New Guinea		09		93	72	99	63	29	53		4	τ-	7	4	7	80	16	15
Philippines	80		82	83	8	87	88	88	88	0	0	က	10	20	48	25	62	89
Rep. Korea	71	92	71	82					83	34	65	26	28					23
Samoa	20	80	100		98	94	92	77	84	48	30	48		63	49	43	77	51
Singapore	88	98				92	85	88	87	22	26				13	22	39	4
Solomon Islands		65	73	92	92		81	88	06		22	71	91	63	92	85	81	107
Tokelau																		
Tonga	88	75	82	75	94	80	93	95	83	48	78	64	26	63	86	75	161	80
Vanuatu						88	88	88	79					36	40	77	52	70
Viet Nam	91	91	06	85	93	92	92	93	92	30	09	79	83	84	83	28	88	98
Wallis & Futuna Is								100	100								49	356
Region	06	9	93	93	92	94	92	93	16	16	28	32	33	33	37	38	39	20
100001	:	;		;		;	;	:	-	!	ì	;	;	;	;	;	:	;

				MALE						Ē	FEMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34		45-54	55-64	+59	0-14	15-24	25-34	35-44	45-54	55-64	65+
American Samoa																					
Australia	0	4	က	_	9	_	12	0	9	7	2	2	7	7	0	10	10	က	∞	က	4
Brunei Darussalam	0	2	25	17		8	6	0	6	14	11	4	2	9	0	14	39	28	12	13	15
Cambodia	37	805	1 514	2 183	1 848		1 487	46	691	1 287	1 975	2 208	1 857	1 256	83	1 496	2 801	4 158	4 056	3 586	2 743
China	1 059	24 199	31 471	30 210	31370		31 210	1 350	18 143	18414	14 147 1	11 578	8 661	9 145	2 4 0 9	42 342	49 885	44 357	42 948	34 991	40 355
China, Hong Kong SAR	3	98	78	119	167	150	337	80	75	113	86	22	37	104	11	169	191	205	222	187	44
China, Macao SAR	0	8	8	16	26	6	23	0	7	7	10	7	4	2	0	15	15	26	33	13	28
Cook Islands																					
Fiji	2	6	7	9	6	2	9		2	9	4	4	7		2	14	13	10	13	12	9
French Polynesia		2	2	-	2	4	3		3	-	-	-	0	1		5	က	2	က	4	4
Guam	0	2	_	က	4	7	2	<del>-</del>	3	_	4	2	-	2	_	2	2	7	9	80	10
Japan	_	130	335	368	713	926	2 502	2	133	272	153	177	199	1 271	က	263	209	521	890	1 155	3 773
Kiribati	2	13	2	6	9	9	0	2	20	4	12	7	က	4	10	33	6	21	13	6	4
Lao PDR	9	98	186	240	233	202	200	7	78	105	160	161	115	96	13	172	291	400	394	317	295
Malaysia	216	1211	2 0 1 0	2 073	1 798	1 438	1 601	196	696	1044	857	699	584	929	412	2 180	3 054	2 930	2 467	2 022	2 227
Marshall Islands	9	4	2	7	7	2	2	4	6	2	4	9	_	4	10	13	4	11	13	3	9
Micronesia	0	က	2	2	0	2	_	4	4	4	_	-	2	_	4	7	9	က	-	4	2
Mongolia	10	206	217	171	93	22	39	19	254	233	148	45	32	19	29	460	450	319	138	87	28
Nauru	0	0	0	0	-	0	0								0	0	0	0	1	0	0
New Caledonia	0	<del>-</del>	<del>-</del>	_	<del>-</del>	_	က	0	0	2	2	0	0	က	0	<del>-</del>	က	က	_	<del>-</del>	9
New Zealand	5	6	10	9	9	8	6	7	18	80	_	10	4	2	12	27	18	7	16	12	14
Niue																					
Northern Mariana Is	0	2	2	2	~	0	2	_	က	0	2	_	0	0	_	2	2	4	2	0	2
Palan	0	0	_	-	_	-	0	-	0	0	-	0	_	2	-	0	_	2	_	2	2
Papua New Guinea	15	164	132	83	26	28	9	24	167	148	51	25	17	2	39	331	280	134	81	45	11
Philippines	356	9 3 8 0	9 302		10 713	6 445	3 648	300	3 2 1 8	4 551	4 761	4 000	2 858	2 018	929	9 2 2 8	13 853	16 219	14 713	9 303	999 9
Rep. Korea	10	401	564	222	493	377	481	13	285	291	174	133	150	450	23	989	855	731	929	527	931
Samoa		2				-			2	2	2		2	-		4	2	2		3	-
Singapore	0	10	15	44	20	26	65	0	4	13	13	13	œ	30	0	4	28	22	83	64	92
Solomon Islands	4	14	6	12	14	8	0	6	14	14	16	13	10	-	13	28	23	28	27	18	_
Tokelau																					
Tonga	0	τ-	<del>-</del>	-	τ-	0	2	0	<del>-</del>	0	<del>-</del>	-	2	0	0	2	-	7	2	2	2
Tuvalu																					
Vanuatu	_	2	4	7	2	2	က	0	4	4	က	2	_	7	_	9	∞	10	7	က	2
Viet Nam	49	3 475	7 036	8 486	2 965	990 9	7 793	99	1 659	2 262	2 327	2 574	2 283	4 896	115	5 134	9 298	10 813	10 539	7 349	12 689
Wallis & Futuna Is	0	0	2	2	2	0	0	0	0	-	0	0	0	0	0	0	က	2	2	0	0
Region	1 785	37 230	52 945	56 086	55 619	42 897	49 449	2 063	25 784	28 810 3	24 929 2	21 699	16 846	19 957	3 848	63 014	81 755	81 015	77 318	59 743	69 406

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

1137 5 243 **62**+ 1 924 1866 97 1240 55-64 546 846 37 45-54 1 738 427 946 8 3 196 ALL 35-44 2 030 275 4 1 035 2 123 25-34 13 1 146 309 26 1 594 19 6 2 518 15-24 13 150 727 0-14 0 131 0 174 31 Country data for the Western Pacific: age and sex distribution of smear-positive cases in non-DOTS areas, 2003 (absolute numbers) 320 56 0 1 979 65+ 969 875 28 55-64 114 13 691 231 77 232 850 17 15 93 1 091 327 25-34 9 834 20 0 13 1 498 123 502 15-24 20 7 1151 396 0-14 57 0 0 19 0 78 **65**+ 9 1 229 3 264 817 136 4 69 991 55-64 2 015 432 0 913 27 0 615 24 2 346 350 714 27 1 394 2 331 182 708 24 25-34 7 186 644 13 15-24 10 8 331 5 0 0-14 0 12 China China, Hong Kong SAR China, Macao SAR Northern Mariana Is Brunei Darussalam Cambodia Palau Papua New Guinea American Samoa Australia Samoa Singapore Solomon Islands Tokelau Fiji French Polynesia Wallis & Futuna Is Malaysia Marshall Islands Nauru New Caledonia New Zealand Niue Cook Islands Micronesia Philippines Rep. Korea Japan Kiribati Lao PDR Mongolia Viet Nam Tonga Tuvalu Guam

Note: the sum of cases notified by age is less than the number of new smear-positive cases notified for some countries.

1367

96

3 650

Country data for the Western Pacific: smear-positive notification rates (per 100 000 population) by age and sex, 2003

							f							-							
_				MALE							FEMALE							ALL			
	0-14	15-24	25-34	35-44	45-54	55-64	+59	0-14	15-24	25-34	35-44 4	45-54 5	55-64 6	+59	0-14 1	15-24 2	25-34	35-44	45-54	55-64	+59
American Samoa																					
Australia	0	~	_	0	_	0	က	0	~	_	0	0		_	0	_	_	0	_	0	<del>-</del>
Brunei Darussalam	0	15	72	64	39	96	165	0	28	38	43	27		108	0	21	22	54	34	96	136
Cambodia	_	20	185	321	425	902	1077	5	44	154	253	405	523	466	_	47	170	285	414	298	673
China	-	23	28	58	39	99	73	τ-	19	17	14	16		18	~	21	22	22	28	38	4
China, Hong Kong SAR																					
China, Macao SAR																					
	•	7	7	7	c	ç	72		ď	5	1	5	00		*	0	5	c	9	40	5
Franch Dolymosia		- a	=	- 4	15	- 2	77		5	2 4	ی ا	2 0		17	-	2 5	2 0	0	5 5	200	35
Tiericii Polyliesia	(	0 ,	2 9	0 6	2 6	8 8	4 6	,	2 3	0 0	٥ ٥	0 6		- 3	c	= 8	0 0	0 0	7 7	7 0	2 2
Guam	0	15	ю I	22	68	120	103	4	24	თ ·	98	22	50	101	7	70	တ ၊	78	31	73	102
Japan	0	3	2	7	12	16	37	0	3	4	3	3		14	0	က	2	2	7	10	24
Kiribati																					
Lao PDR	_	16	47	87	128	190	212	<del>-</del>	14	56	22	84	93	87	-	15	36	71	105	138	145
Malaysia	2	53	105	126	147	209	323	2	44	99	23	26		109	2	49	81	06	102	149	208
Marshall Islands																					
Micronesia	0	25	28	35	0	102	22	20	34	53	17	23	96	46	6	59	41	56	11	66	51
Mongolia	2	72	96	66	102	106	92	2	06	104	8	48	26	35	က	8	100	91	75	82	29
Nauru																					
New Caledonia	0	2	2	9	œ	12	49	0	0	7	12	0	0	44	0	က	00	6	4	9	46
New Zealand	_	3	4	2	2	4	4	2	7	3	0	4	2	2	_	2	4	1	3	3	3
Niue																					
Northern Mariana Is																					
Palau																					
Papua New Guinea	_	28	30	25	27	25	∞	5	32	33	16	13		∞	7	30	32	21	21	21	00
Philippines	2	77	147	250	339	344	276	2	40	74	40	124	147	121	2	29	111	177	230	243	190
Rep. Korea	0	19	28	30	39	20	94	-	19	19	12	12		22	-	19	24	21	56	34	72
Samoa		10				53			<del>_</del>	17	30		51	22		7	7	13		41	13
Singapore	0	9	6	17	28	43	88	0	2	6	7	9		32	0	4	6	12	17	27	24
Solomon Islands	4	28	24	22	6	84	0	6	30	40	71	91		16	9	59	32	63	94	86	80
Tokelau																					
Tonga Tuvalu	0	თ	13	19	29	0	72	0	10	0	19	25	69	0	0	თ	7	19	27	37	¥
Vanuatu	2	6	28	63	65	44	98	0	19	27	26	28		09	-	14	27	44	47	34	73
Viet Nam	0	4	103	158	236	297	385	~	20	33	42	74	125	207	0	31	89	66	153	208	289
Wallis & Futuna Is																					
Region	-	27	36	43	22	89	86	-	70	21	20	22	27	59	-	23	29	31	39	48	22

Note: rates are missing where data for smear-positive cases are missing, or where age- and sex-specific population data are not available.

15 897 107 133 37 268 37 268 229 292 0 0 12 16 175 90 728 **454** 33 92 1997 1996 1995 14 603 515 764 0 402 203 991 160 142 218 6 537 6 537 7 8 7 8 7 7 8 4 8 4 61 1 2 285 1 2 283 1 2 285 1 2 285 1 2 285 1 2 285 1 2 285 1 2 285 1 2 285 1 2 285 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 2 2 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 893 1 8 25 7 451 178 134 46 999 1 830 367 2 8 2 8 111 114 52 994 3 401 207 371 57 864 44 1 841 309 20 20 20 230 59 784 1991 | 1988 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 | 136 Country data for the Western Pacific: number of TB cases notified, 1980–2003 8 276 276 7 577 7 507 7 301 1 650 1 1 630 6 1 386 285 285 1 980 1 , Hong Kong SAR American Samoa Australia Brunei Darussalam orthem Mariana Is pua New Guinea Japan Kiribati Lao PDR Malaysia Marshall Islands Micronesia Mongolia reporting reporting vauru New Caledonia New Zealand hilippines ep. Korea

Country data for the Western Pacific: case notific	e Weste	rn Pac	ific: c	ase n		ation rates (per 100 000 population), 1980–2003	ates (p	er 100	000 b	opulat	ion), 1	980-2	003											
	1980	1981	1982	1983	1984	1985	1986	1987	1988 1	1989 1	1990 1	1991 1	1992 1	1993 1	1994 1	1995 19	1996 1	1997 1	1998	1999	2000	2001	2002	2003
American Samoa	9	18	17	22	32	13	20	21	29	1	19	9	2	80	80		0	7	2	7	2	2	က	2
Australia	10	0	00	00	80	7	9	9	9	9	9	9	9	9	9	9		9	2	9	2	2	2	2
Brunei Darussalam	102	143	120	131	118	107	95	80	52	51	99		99	22				51		83	95	63	99	28
Cambodia	39	59	116	103	132	125	122	104	118	84	29	108	155	123	136	ľ	126	129	136	150	144	142	178	199
China			10	11	4	21	24	23	27	27	33	29	27	29	30		41	38	36	36	36	37	36	47
China, Hong Kong SAR	160	150	144	137	145	138	135	131	125	119	114	109	112	110	104	0	103	110	117	112	111	105	68	80
China, Macao SAR	437	226	87	163	229	186	131	117	92	9/	92	98	75	72			136	134	107		100	102	8	80
Cook Islands	207	26	108	165	114	204	96	06	111	9	2	43	64	32	21		0	0	2	16	=	1	2	0
Fiji	33	28	24	27	24	32	28	24	23	30	31	34	33	24	37	56	26	22	21	24	18	22	18	21
French Polynesia	20	42	41	47	47	45	48	44	34	38	30	25	41	38	42		39	41	46	41	27	56	27	20
Guam	52	38	44	42	46	31	40	27	32	22			43	20	99						35	40	32	4
Japan	61	26	24	52	51	48	47	46	44	43	42	41	39	39	36	34	33	33	35	32	31	28	56	25
Kiribati	252	317	321	207	177	161	197	164	303	172	92	125	135	131	330		414	578	339	308	300	222	227	324
Lao PDR	238		141	137	185	118	41	91	186	73	44	46	23	47	25	18	30	39	43	47	42	45	47	49
Malaysia	82	78	82	78	69	29	29	29	65	61	99	09	61	25	29		61	63	49	99	65	63	09	64
Marshall Islands	20	22	36	43	33	40	94	62	56	16		22	113	131			122		86	81	29	108	26	113
Micronesia			98	90	06	77	89	109	84	72	381	354	110	146	164		117	66	115		85	26	117	91
Mongolia	70	64	92	84	88	157	143	120	121	103	7.1	71	65	61	73	116	143	123	119	135	124	139	150	151
Nauru	0	56	104	0	0	0	96	70	06	0	74				38					17	33	24	33	23
New Caledonia	9/	88	81	114	8	29	62	46	89	9/	84	80	78	22	51		53	44	4	37	44	28	59	16
New Zealand	15	14	14	13	13	1	10	6	6	6	10	10	6	8	10		10	6	10	12	6	10	6	10
Niue	29	0	64	101	35	0	192	0	123		0		06	46	92		92	0	0	49	0	0	203	0
Northern Mariana Is		140	370	331	233	232	52	164	72	69	64		137		87	87	88	153	152	66	107	79	20	22
Palau	140	81	135	108	151	191	93	566	117	20		39	25	153	245		28	83		170			22	44
Papua New Guinea	78	92	81	85	86	95	77	29	109	85	61	81	29	167	117		106	161	222	251	227	291	92	322
Philippines	234	237	207	205	287	278	276	288	314	364	519	332	369	272	269		237	274	223	196	158	139	151	168
Rep. Korea	236	255	257	230	212	214	215	210	177	165	149	134	110	106	98		87	72	75	69	47	79	74	71
Samoa	38	32	28	56	54	27	41	18	18	23	27	27	16	30	27		19	19	13	18	25	13	18	15
Singapore	112	86	98	80	81	72	64	22	28	22	23	29	99	26	20		54	53	26	46	43	37	36	38
Solomon Islands	116	132	132	119	128	139	104	115	124	158	120	94	107	105	95		77	80	72	89	69	65	22	61
Tokelau	0	64	0	0	0	121	0	546	61	0	62	63	64		0	131	0			0	0	0		0
Tonga	99	20	46	21	26	20	36	25	14	36	23	20	58	33	23	20	22	21	30	22	24	12	28	15
Tuvalu	441	235	154	291	112	393	327	262	281	299	260	334	329	303	202	378			181	139	157	155	124	283
Vanuatu	152	77	141	156	146	8	97	65	83	66	94	150	122	70	91		71	101	92	63	77	87	49	49
Viet Nam	81	80	93	92	9/	79	79	06	83	81	9/	88	82	75	72		101	104	115	115	115	115	118	114
Wallis & Futuna Is	208	509	42	139	112	109		256	7	220		159	29	79	78		57	98					130	102
Region	27	27	34	34	39	4	46	45	49	20	29	49	48	46	45	51	54	53	20	49	47	47	47	22

Rate (per 100 000 population) 1997 1998 1999 454 732 371 577 251 1482 204 762 1940 Country data for the Western Pacific: new smear-positive cases, 1993–2003 12 867 144 886 7 271 12 14 769 11 058 104 729 1993 1 557 68 88 88 84 898 2 429 108 4 4 4 6 954 China, Hong Kong SAR China, Macao SAR Niue Northern Mariana Is Papua New Guinea Brunei Darussalam Singapore Solomon Islands Tokelau Vallis & Futuna Is Malaysia Marshall Islands New Caledonia New Zealand Philippines Rep. Korea Samoa **Aicronesia** Kiribati Lao PDR **Aongolia** 

## **Notes**

#### **Brunei Darussalam**

Treatment outcomes are for laboratory-confirmed cases, and for nationals only. Non-nationals are deported to their country of origin after two weeks of treatment or conversion to smear-negative status.

#### Cambodia

The latest estimate of the incidence of TB does not take into account the results from a recent disease prevalence survey.

#### **China**

The NTP notes that age and sex data are incomplete. Relapse notifications include both smear-positive cases and smear-negative/-unknown cases.

#### **China, Hong Kong SAR**

Patients' outcomes are assessed at 12 months after diagnosis.

#### Guam

Of 9 smear-negative, culture-negative cases, and 3 cases with smear and culture not done, all 12 completed treatment.

#### Malaysia

Age and sex data are for all new cases.

#### **Palau**

DOTS coverage was not reported, but assumed to be same as in the previous report.

#### **Singapore**

All data refer to TB cases notified among resident population.

#### **Tokelau**

DOTS coverage was not reported, but assumed to be same as in the previous report.

**WPR** 

# The World Health Organization monitors the tuberculosis epidemic and evaluates surveillance, planning, and financial data in support of national TB control programmes.

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