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The Global Laboratory Initiative Roadmap

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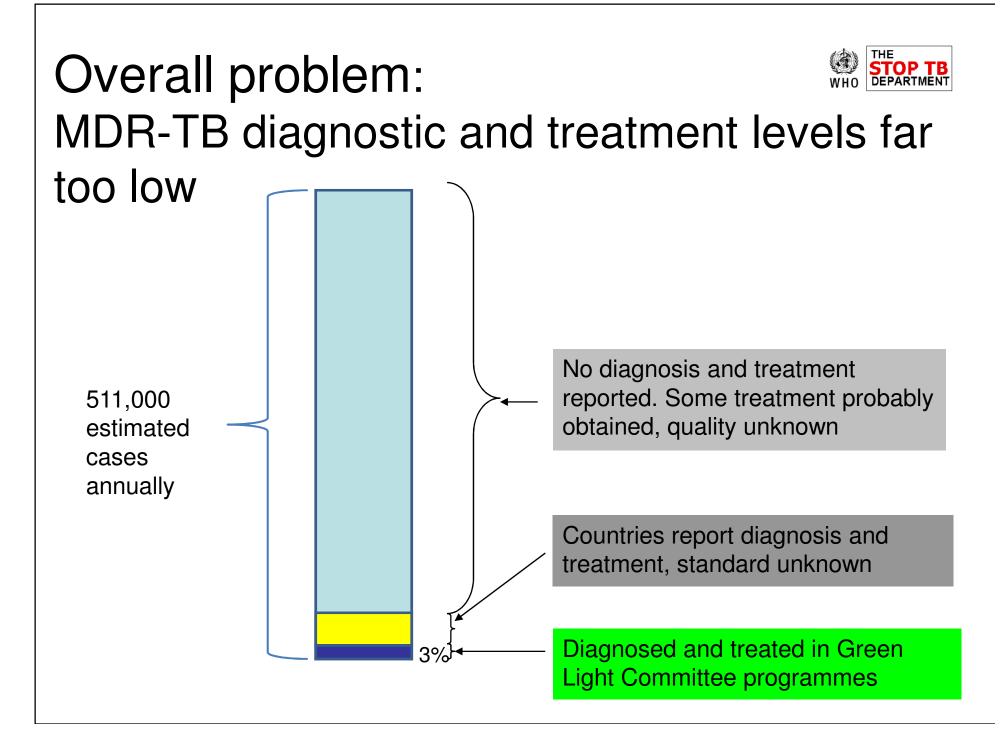


Global TB estimates - 2007

(Updated February 2009)



	Estimated number of cases	Estimated number of deaths
All forms of TB Greatest number of cases in Asia; greatest rates per capita in Africa	9.27 million (139 per 100,000)	1.77 million (27 per 100,000)
Multidrug-resistant TB (MDR-TB)	511,000	150,000
Extensively drug- resistant TB (XDR-TB)	50,000	30,000
HIV-associated TB	1.4 million	456,000



Laboratory scale-up

Driven by

- Case detection moving towards universal access
- HIV- associated and drug resistant TB

Challenged by

- Weak health systems
- Inadequate human resources
- Insufficient programmatic and managerial capacity
- Inadequate infrastructure (biosafety)
- Problems of availability and access
- Slow technology transfer
- Lack of recognition of laboratory importance in TB control, weak communication between NTPs and laboratory services



Acceleration

Recent developments:

- At least 20 new technologies in various stages of development and evaluation
- Distinct target areas for drug-resistant TB being addressed
- WHO policy formulation
 - Liquid culture, rapid speciation and line probe assays endorsed by WHO 2007-2008;*
 - LED microscopy and selected non-commercial culture and drug susceptibility testing methods 2009
- Expanded access to new diagnostics and laboratory strengthening





Why a Roadmap?



Process

- May 08: GLI CG meeting
 - GLI strategic objectives defined
- May 08: 1st annual GLI meeting
 - Consultant findings on stakeholder interviews and country fact finding visits
 - Break-out group discussions to identify gaps and next steps
- Oct 08: Dedicated TBCAP funding
- Oct 08 Jun 09:
 - Conceptual framework defined
 - Country case studies pursued and common themes identified
 - Stakeholder interviews continued
 - WHO policy recommendations incorporated
- Jun 09 Aug 09
 - Intensive revision by Writing Committee, GLI CG and external laboratory experts



Purpose and scope

- <u>Structured framework</u> for TB laboratory strengthening based on WHO-GLI norms and standards, documented best-practices at country level, growing lessons from the field ('learning by doing')
- <u>Generic document</u> encompassing managerial, operational and technical aspects of TB laboratory strengthening within the context of national laboratory strategic plans
- <u>Broad user base</u> including NTP and NRL managers, technical agencies, donor agencies, implementing partners, programme budgeting and planning officers
- <u>Living document</u>, responsive to changes in TB diagnostic landscape and WHO policy frameworks
- Supported by resource list for tools and technical procedures



Core elements

- Laboratory infrastructure and maintenance
- Equipment validation and maintenance
- Specimen referral and transport mechanisms
- Policy framework for implementing new TB diagnostics
- Laboratory commodity and supply chain management
- Laboratory information and data management systems
- Laboratory quality management systems
- Laboratory human resource development



Stepwise approach (1)

Policy change at country level, based on

- Local epidemiology (TB, HIV, MDR-TB)
- NTP priorities for case detection (risk groups)
- Laboratory networks and capacity
- Laboratory staff resources and skills base
- Treatment policies for drug-resistant TB
- Financial resources



Stepwise approach (2)

Expansion of laboratory services based on

- Tiered system (peripheral, intermediate, central)
- Available technologies
- Ancillary laboratory needs related to specialised treatment (eg. ART, second-line anti-tuberculosis drugs)
 - General microbiology, biochemistry, haematology, etc.
- Integrated approach



Stepwise approach (3)

Phase 1: Laboratory preparedness

- Assessment of TB laboratory networks and diagnostic policies
- Upgrade of laboratory infrastructure and biosafety
- Development and implementation of GLP, SOPS, QA, etc.
- Training of core laboratory staff
- Initiation of NTP policy reform on diagnostics

Phase 2: Introduction of new diagnostics

- Integration of new diagnostics into NTP policies and procedures
- Procurement and installation of instruments, reagents, supplies
- Validation of new tools and laboratory performance
- Adjustment of NTP policy based on local data
- Phase 3: Impact assessment
 - Continued mentoring, technical support and oversight
 - Assessment of impact on NTP outcomes



Analytical process

- Quantify or estimate TB, TB-HIV and MDR-TB burden
- Identify and target patient risk groups, eg.
 - Treatment failures
 - Non-converting patients
 - HIV+ individuals
- Quantify or estimate diagnostic need to identify cases
 - Number of suspects to be screened
 - Number and type of laboratories at each service level
- Estimate budget for comprehensive laboratory services
 - All core components
 - Capacity for diagnosis and monitoring
 - Ancillary laboratory tests



Policy considerations

- Current technologies not mutually exclusive
 - Conventional culture capacity required for SM- specimens
 - Conventional DST capacity required to detect XDR-TB
- Liquid culture and line probe assay as gold standards, to be phased in without loss of existing culture and DST capacity
- LED microscopy as alternative for both fluorescence and conventional light microscopy
- Selected non-commercial culture and DST methods not alternatives for gold standards, but may provide interim solution



Issues for SRL discussion

- Definition of SRL
 - Technical expert in all aspects of laboratory strengthening; or
 - Technical expert in particular aspects of laboratory strengthening
- Role of SRL network
 - In overall laboratory capacity development
 - In drug resistance surveillance

- ...

Moving beyond TB



Strengthening TB laboratories

'From unimaginable...to indispensable'



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- <u>With additional input from</u>: Catherine Mundy, Giorgio Roscigno, Akos Somoskovi, Veronique Vincent
- <u>GLI partners interviewed</u>: APHL, ASM, CDC, FIND, GTZ, KNCV, PATH, PEPFAR, PIH, TBCAP, Union, WHO
- And with apologies for any unintended oversight...

