

5.11 TUBERCULOSIS NOTIFIABLE

RECOMMENDATIONS

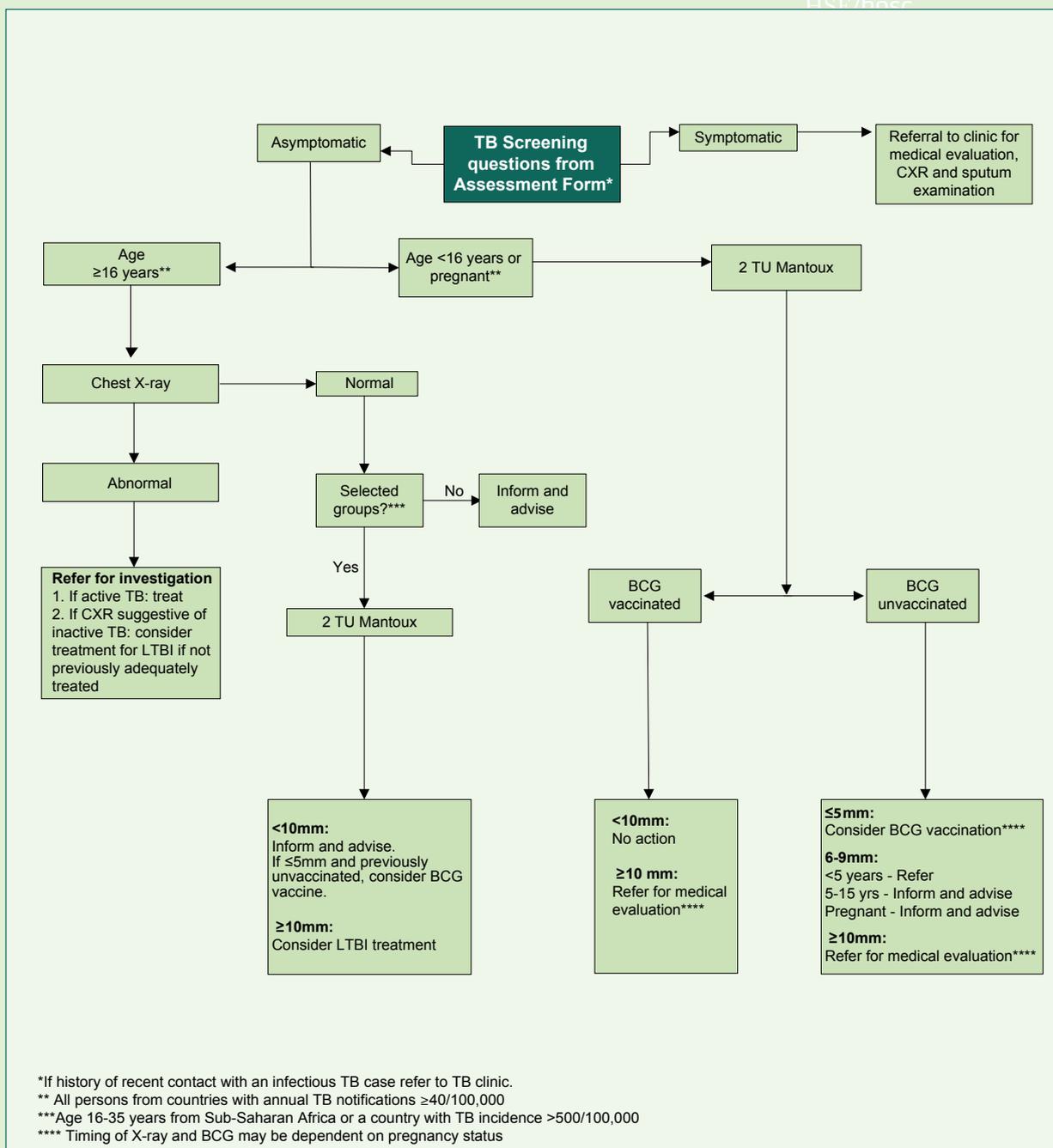
Risk assess:

All migrants from countries where prevalence of TB disease is known to be ≥ 40 cases per 100,000 population, as per the National TB guidelines 2010.

See Appendix F for current list of countries

Algorithm for assessing all migrants from countries with ≥ 40 cases of TB per 100,000 pop.

TB Screening Questions from Assessment Form*



Source: Adapted from HSE HPSC Guidelines on the prevention and control of tuberculosis in Ireland 2010.⁽⁵⁾

Epidemiology

The incidence of TB disease in immigrant groups is high particularly within the initial years following arrival, principally due to the reactivation of latent infection.⁽¹⁾ Immigrants are at increased risk of disease if they originate from countries with a high incidence of TB (see appendix F) and HIV. Overcrowded living conditions, poverty and migration from a war zone also contribute to the spread of TB.⁽¹⁾

Ireland

The following information on the notification rate of tuberculosis (TB) in Ireland is contained in the HPSC Report on the Epidemiology of Tuberculosis in Ireland 2012.⁽²⁾

In Ireland, the notification rate of TB has started to decrease since 1998 (11.7/100,000 population in 1998 and 7.8/100,000 population in 2012). However, during this period the rate in the foreign-born population increased significantly from 8.7/100,000 in 1998 to 20.9/100,000 in 2012, while the rate in the Irish-born population decreased from 11.2/100,000 in 1998 to 5.2/100,000 in 2012. The notification rate in foreign-born cases reached a peak in 2008 (Figure 5.11.1).

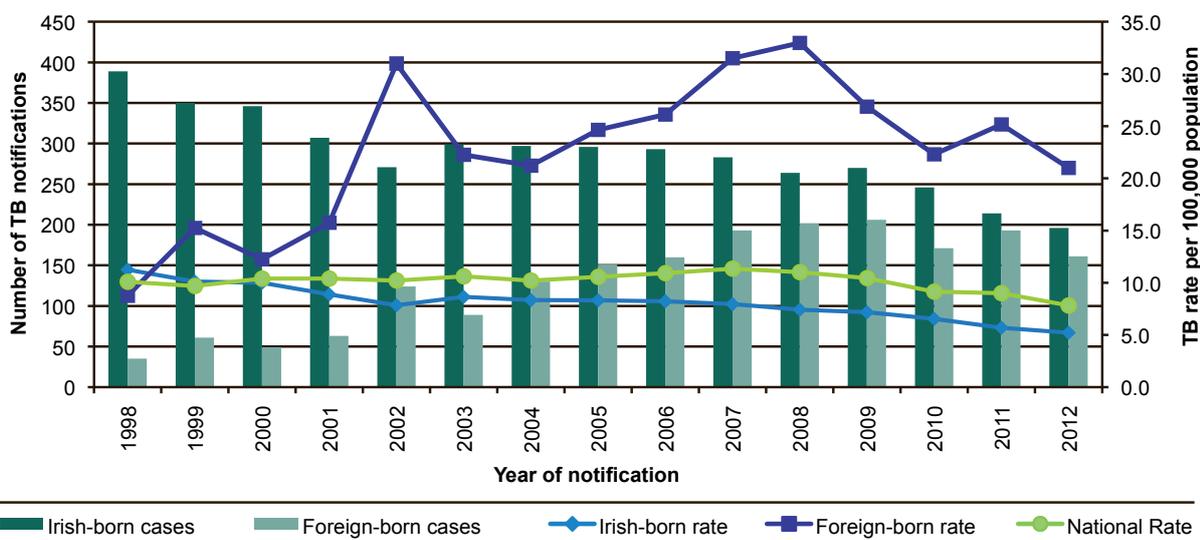


Figure 5.11.1. TB cases and rates per 100,000 population by geographic origin, 1998 to 2012.

Source: Report on the Epidemiology of Tuberculosis in Ireland 2012, HPSC.⁽²⁾

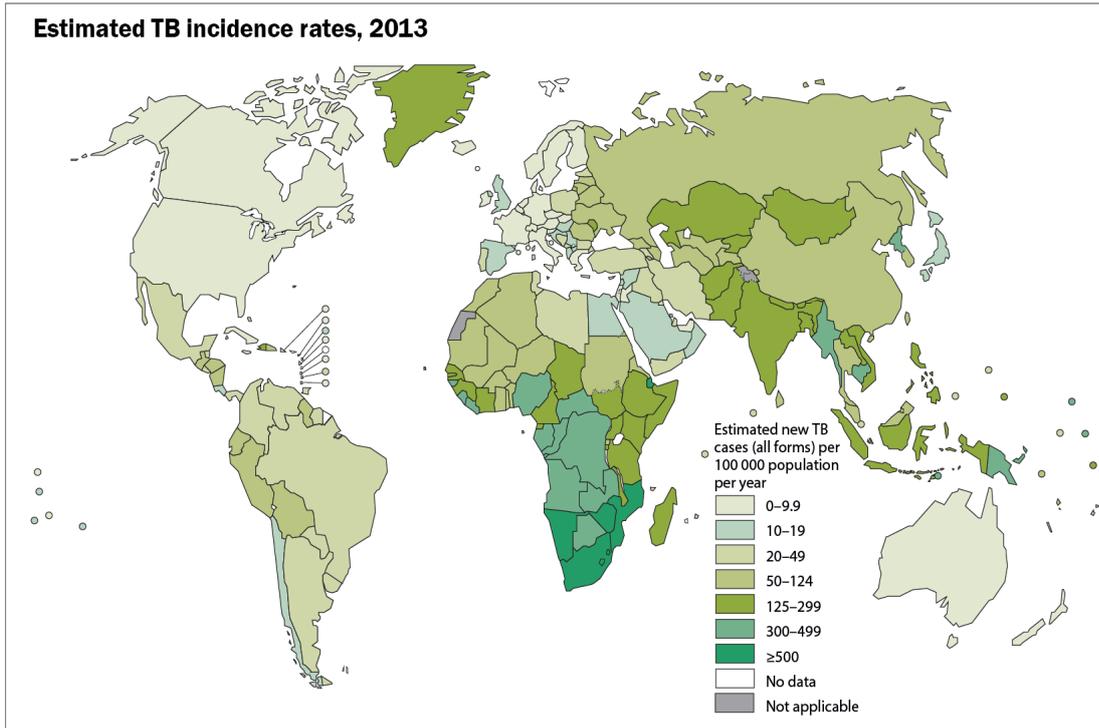
In 2012, the majority (76.1%) of cases born outside Ireland were aged between 15 and 44 years compared to 35.7% of Irish cases occurring in this age range. The median age of foreign-born cases was 33 years compared to 51 years among Irish-born cases.

The majority of foreign-born cases were from Asia, followed by Africa and Europe.

The percentage of drug resistant TB cases is still low in Ireland although it has been increasing in recent years. Between 2002 and 2012 there were 30 multi-drug resistant (MDR) TB cases in Ireland; 24 (80%) were foreign-born and 6 (20%) were Irish-born (Personal communication Dr Joan O'Donnell, HPSC, 2015).

Worldwide

There were an estimated 9 million new TB cases globally in 2013. Approximately 1.5 million people died from TB in 2013.⁽³⁾ The estimated TB incidence rate by country in 2013 is shown in figure 5.11.2. Globally, 13% of new TB cases were co-infected with HIV in 2013 (figure 5.11.3).



The boundaries and names shown and the designations used on this map do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement.

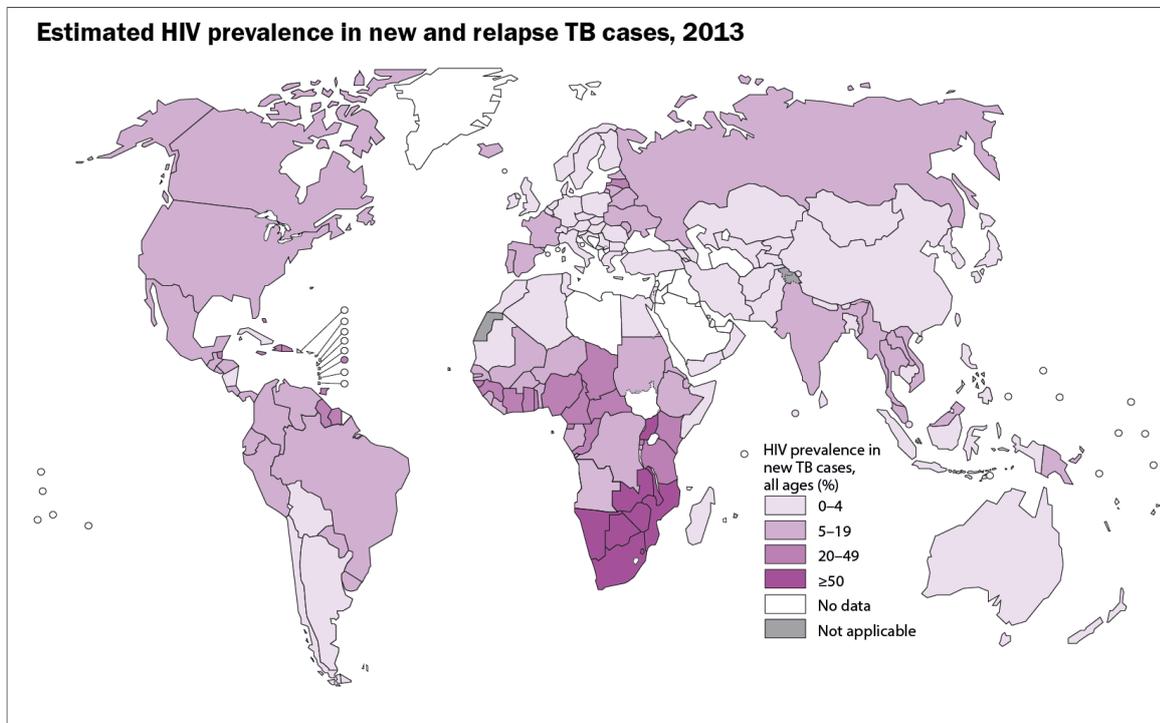
Data Source: *Global Tuberculosis Report 2014*. WHO, 2014.



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Figure 5.11.2 Estimated TB incidence rate by country, 2013

Source: WHO Global TB report 2014 (3)



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Data Source: *Global Tuberculosis Report 2014*. WHO, 2014.



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Figure 5.11.3 Estimated HIV Prevalence in new and relapse TB cases, 2013

Source: WHO Global TB report 2014 (3)

Rationale for assessment

- TB is a significant cause of morbidity and mortality worldwide.
- Many migrants coming to Ireland are from countries with a high incidence of TB.⁽⁴⁾
- TB is a curable disease.⁽⁵⁾
- The tests for diagnosing TB are simple and well established (Mantoux, IGRA, chest X-ray, sputum analysis)
- Effective treatment regimens are available.⁽⁵⁾
- Latent TB (LTBI) is curable.⁽⁵⁾
- Treatment of TB and LTBI reduces the spread of disease.⁽⁵⁾

Assessment

The following guidelines for assessment are derived from the *Guidelines on the Prevention and Control of Tuberculosis in Ireland (2010)*.⁽⁵⁾

- All new migrants to Ireland who originate from a country with a high incidence of TB should be provided with an opportunity to be assessed for TB. Those include people who have recently arrived or returned from a country with an incidence of TB ≥ 40 cases per 100,000 population per year.⁽⁵⁾ Countries with an incidence of ≥ 40 cases per 100,000 population per year are mainly located in Africa, the Eastern Mediterranean, Central and Eastern Europe, South East Asia, the Western Pacific and Central and South America. A list of current high incidence countries can be found in **Appendix F**.
- Every effort should be made to identify candidates either at reception centres for asylum seekers or in other clinical settings.
- TB screening for active disease and LTBI should be encouraged.
- Voluntary assessment for HIV should be offered to those with diagnosed TB as co-infection with HIV is common.⁽⁵⁾

Health assessment

A full history and examination should be undertaken for all new entrants to enquire into past history of TB and BCG status, current symptoms, and recent contact with a TB case. Those with symptoms should be urgently referred to a respiratory clinic for further clinical assessment.

Chest X-ray

Chest X-rays should be offered to all new migrants aged ≥ 16 years who are from a country with a high incidence of TB (provided they are not pregnant). All those with abnormal chest X-ray results suggestive of active disease or of inactive TB should be referred for medical evaluation. Treatment of LTBI should be considered in those with radiological evidence of inactive TB, if not previously treated.

Tuberculin Skin Test (TST) (Mantoux test)

Individuals ≥ 16 years

Asymptomatic individuals with a normal chest X-ray in a selected group i.e. those aged 16 to 35 years from Sub-Saharan Africa or a country with a TB incidence greater than 500 per 100,000* should be offered a TST (2TU Mantoux test) regardless of BCG vaccination status. Pregnant females (no chest X-ray, see above) should also have a TST (2TU Mantoux test), regardless of BCG vaccination status.

*These countries include Botswana, Cambodia, Djibouti, Lesotho, Namibia, Sierra Leone, South Africa, Swaziland, Timor-Lest, Zambia and Zimbabwe.

Those with TST results ≥ 10 mm should be referred for further medical evaluation and considered for LTBI treatment. Individuals with TST results < 10 mm should be informed and advised of the signs and symptoms of TB disease and asked to seek medical care if they experience these symptoms. Consider BCG vaccination for all those aged ≤ 35 years with TST results < 6 mm who are previously unvaccinated (see *TB disease algorithm above*).

While all age groups should be considered for treatment of LTBI, care should be taken when prescribing LTBI therapy for those with co-morbidities which increase the likelihood of hepatotoxicity. The use of Directly Observed Therapy (DOT) should also be considered in this population on a case by case basis if resources allow.

Individuals <16 years

All new entrants aged 0 to 15 years should be screened initially by health questionnaire and TST (2TU Mantoux test).

Unvaccinated (BCG)

All those under 16 years of age with a negative TST result (<6mm) should be offered BCG vaccination after consideration has been given to the individual's HIV status.

Unvaccinated children under five years of age with a Mantoux reading of 6-9 mm should be referred to a TB clinic where treatment for LTBI should be considered if the chest X-ray is normal.

Unvaccinated children aged five to 15 years with a Mantoux reading of 6-9mm and without a history of recent contact with a TB case should be advised of the signs and symptoms of TB.

All unvaccinated children (aged 0 to 15 years) with a Mantoux reading of ≥ 10 mm should be referred to a clinician with experience in the management of LTBI and chemoprophylaxis should be considered if the chest X-ray is normal.

Vaccinated with BCG

Vaccinated children should be referred for a chest X-ray and chemoprophylaxis considered if the Mantoux reading is ≥ 10 mm. If the result is <10mm, no further action is required.

Interferon Gamma Release Assay (IGRA) testing

Foreign-born individuals can have a higher incidence of LTBI and may be more likely to have clinical conditions that increase the likelihood of reactivation of LTBI, such as HIV infection. Evidence from international studies suggests that IGRA tests have a higher specificity than tuberculin skin tests and have less potential for false positive results.⁽⁶⁻⁹⁾

It is recommended that the TST should be used initially to detect LTBI and a person with a positive result should be considered to have LTBI. False negative results are not uncommon in immunodeficient individuals; therefore if a clinician is concerned about the possibility of such a TST result, an IGRA can be conducted. LTBI can be considered if an IGRA test is positive, while indeterminate results should be repeated. Indeterminate results may indicate laboratory error or anergy, therefore a person's history, clinical features and laboratory findings must be taken into account when diagnosing LTBI using an IGRA.⁽⁵⁾

The use of IGRA can be considered:

- As a confirmatory test in those individuals with a positive TST.
- In screening new entrants with concomitant conditions that increase the individual's risk of reactivation of LTBI.

Contacts

Contacts of cases of active TB that are notified to public health will be followed up through Departments of Public Health following notification.

TB/HIV

The management of patients with TB and HIV is complex. Cases of TB/HIV should always be referred to physicians with expertise in treating both TB and HIV. See Chapter 10 of Guidelines on the prevention and control of tuberculosis in Ireland 2010.⁽⁵⁾

References

- (1) Tuberculosis at the end of the 20th century in England and Wales: results of a national survey in 1998. *Thorax* 2001;56:173-9.
- (2) Health Protection Surveillance Centre. Report on the Epidemiology of Tuberculosis in Ireland 2012 [Internet]. [cited August 2015]. Available from: <http://www.hpsc.ie/A-Z/VaccinePreventable/TuberculosisTB/Epidemiology/AnnualReports/2012/File,15063,en.pdf>
- (3) World Health Organization. Global Tuberculosis control 2014. Geneva: WHO; 2014. http://www.who.int/tb/publications/global_report/en/
- (4) CSO Ireland 2011 [cited 2013 Jun 12]; Available from: <http://www.cso.ie/en/census/census2011reports/census2011profile6migrationanddiversity-aprofileofdiversityinireland/>
- (5) Health Protection Surveillance Centre. Guidelines on the prevention and control of tuberculosis in Ireland 2010. Dublin; 2010. <http://www.hpsc.ie/AboutHPSC/ScientificCommittees/Publications/File,4349,en.pdf>
- (6) Public Health Agency of Canada. An advisory committee statement (ACS) by the Canadian Tuberculosis Committee on Interferon Gamma Release Assays for Latent TB Infection. *Canada Communicable Disease Report* 2007;33(10):1-18.
- (7) Tuberculosis: Clinical diagnosis and management of tuberculosis, and measures for its prevention and control. March 2011. Available from: <http://www.nice.org.uk/nicemedia/live/13422/53638/53638.pdf>
- (8) Centers for Disease Control and Prevention. Guidelines for using the Quantiferon-TB test for detecting *Mycobacterium tuberculosis* infection, United States. *Morbidity and Mortality Weekly Report* 2005; 54 (RR15)(December 16):49-55.
- (9) Pai M, Dheda K, Cunningham J, Scano F, O'Brien R. T-cell assays for the diagnosis of latent tuberculosis infection: moving the research agenda forward. *Lancet Infect Dis* 2007;7(6):428-438.

Appendix F. List of countries with a TB incidence of $\geq 40/100,000$ population

| Country | WHO region | Estimated rate of TB per 100,000 population |
|---------------------------------------|-----------------------|---|
| Swaziland | Africa | 1349 |
| South Africa | Africa | 1003 |
| Sierra Leone | Africa | 674 |
| Namibia | Africa | 655 |
| Lesotho | Africa | 630 |
| Djibouti | Eastern Mediterranean | 620 |
| Marshall Islands | Western Pacific | 572 |
| Zimbabwe | Africa | 562 |
| Mozambique | Africa | 552 |
| Timor Leste | South-East Asia | 498 |
| Kiribati | Western Pacific | 429 |
| Gabon | Africa | 428 |
| Zambia | Africa | 427 |
| Cambodia | Western Pacific | 411 |
| Democratic People's Republic of Korea | South-East Asia | 409 |
| Botswana | Africa | 408 |
| Congo | Africa | 381 |
| Myanmar (Burma) | South-East Asia | 377 |
| Central African Republic | Africa | 367 |
| Mauritania | Africa | 350 |
| Papua New Guinea | Western Pacific | 348 |
| Democratic Republic of Congo | Africa | 327 |
| Angola | Africa | 316 |
| Liberia | Africa | 304 |
| Somalia | Africa | 286 |
| Gambia | Africa | 284 |
| Kenya | Africa | 272 |
| Philippines | Western Pacific | 265 |
| Ethiopia | Africa | 247 |
| Guinea-Bissau | Africa | 242 |
| Tuvalu | Western Pacific | 241 |
| Cameroon | Africa | 238 |
| Madagascar | Africa | 234 |
| Pakistan | Eastern Mediterranean | 231 |
| Bangladesh | South-East Asia | 225 |
| Mongolia | Western Pacific | 223 |
| Haiti | The Americas | 213 |
| Lao People's Democratic Republic | Western Pacific | 204 |
| Micronesia (Federated States of) | Western Pacific | 194 |
| Afghanistan | Eastern Mediterranean | 189 |
| Indonesia | South-East Asia | 185 |
| Bhutan | South-East Asia | 180 |
| Uganda | Africa | 179 |
| Guinea | Africa | 178 |
| India | South-East Asia | 176 |
| Cote d'Ivoire | Africa | 172 |
| Greenland | Europe | 170 |
| United Republic of Tanzania | Africa | 165 |
| Malawi | Africa | 163 |
| Nepal | South-East Asia | 163 |
| Republic of Moldova | Europe | 160 |
| Chad | Africa | 151 |
| Vietnam | Western Pacific | 147 |
| South Sudan | Eastern Mediterranean | 146 |
| Cape Verde | Africa | 144 |
| Kyrgyzstan | Europe | 141 |
| Equatorial Guinea | Africa | 139 |
| Kazakhstan | Europe | 137 |
| Senegal | Africa | 137 |

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|---------------------------|-----------------------|-----|
| Burundi | Africa | 130 |
| Bolivia | The Americas | 127 |
| Thailand | South-East Asia | 119 |
| Georgia | Europe | 116 |
| Sudan | Eastern Mediterranean | 114 |
| Guyana | The Americas | 109 |
| Nigeria | Africa | 108 |
| Republic of Korea | Western Pacific | 108 |
| Tajikistan | Europe | 108 |
| Niger | Africa | 104 |
| Morocco | Eastern Mediterranean | 103 |
| Solomon Islands | Western Pacific | 97 |
| Azerbaijan | Europe | 95 |
| Peru | The Americas | 95 |
| Romania | Europe | 94 |
| Eritrea | Africa | 93 |
| Sao Tome and Principe | Africa | 93 |
| Ukraine | Europe | 93 |
| Russian Federation | Europe | 91 |
| Algeria | Africa | 89 |
| Rwanda | Africa | 86 |
| China, Macao SAR | Western Pacific | 83 |
| Malaysia | Western Pacific | 80 |
| Uzbekistan | Europe | 78 |
| China, Hong Kong SAR | Western Pacific | 77 |
| Turkmenistan | Europe | 75 |
| China | Western Pacific | 73 |
| Togo | Africa | 73 |
| Ghana | Africa | 72 |
| Belarus | Europe | 70 |
| Benin | Africa | 70 |
| Northern Marianna Islands | Western Pacific | 69 |
| Brunei | Western Pacific | 68 |
| Lithuania | Europe | 66 |
| Sri Lanka | South-East Asia | 66 |
| Vanuatu | Western Pacific | 65 |
| Wallis and Futuna Islands | Western Pacific | 65 |
| Dominican Republic | The Americas | 62 |
| Guatemala | The Americas | 60 |
| Mali | Africa | 60 |
| Equador | The Americas | 59 |
| Burkina Faso | Africa | 54 |
| Honduras | The Americas | 54 |
| Nauru | Western Pacific | 54 |
| Latvia | Europe | 53 |
| Armenia | Europe | 52 |
| Singapore | Western Pacific | 50 |
| Bosnia Herzegovina | Europe | 49 |
| Yemen | Eastern Mediterranean | 49 |
| Guam | Western Pacific | 48 |
| Panama | The Americas | 48 |
| Brazil | The Americas | 46 |
| Iraq | Eastern Mediterranean | 45 |
| Paraguay | The Americas | 45 |
| Maldives | South-East Asia | 41 |
| Qatar | Eastern Mediterranean | 41 |
| Suriname | The Americas | 40 |
| Belize | The Americas | 40 |
| Libya | Eastern Mediterranean | 40 |

Source: Public Health England.

Available from: <http://www.hpa.org.uk/Topics/InfectiousDiseases/InfectionsAZ/Tuberculosis/TBWorldwideSurveillanceData/>.

Accessed 26/01/2014.