U.S. Department of State

TIDEDCIII OCIC WODKCI

OMB No. 1405-0113
EXPIRATION DATE: xx/xx/xxxx
ESTIMATED BURDEN: 20 MINUTES
(See Page 2 – Back of Form)

| | | | | | | | | MATED BURDEN: 20 MIN Page 2 – Back of Form) | NUTES |
|----|---|--|--|---|-------------|---|-------------------|--|------------|
| | | Name (Last, First, MI) | | | | | | Age | |
| | | Birth Date (mm-dd-yyy | у) | Passport Nun | nber | Al | ien (Case | e) Number | |
| 1. | | Mediated Immunity to Tuber pplicants 2 through 14 years (| | ed TB rate ≥20 | per 100,000 | and contacts | ; perform | one type only. | |
| | | te applied (mm-dd-yyyy) | | • | | e: TB minus r | | | |
| | 1 D | ate drawn <i>(mm-dd-yyyy)</i> Positive Negative ndeterminant, Borderline, or E | <u> </u> | ☐ T-Spot | TB Respons | Number of cells se: Higher of Panel B minus | | · | _ |
| | ☐ Chest X-Ra | mptoms of tuberculosis | Mown HIV infection TST ≥10 mm or IGRA p Contact: TST ≥ 5 mm or | | | _ Date Chest X- | Ray Take | en <i>(mm-dd-yyyy)</i> | |
| | Normal Fin | ndings | Abnormal Findings (Inc | dicate category | and finding | _checking all | that annly | vin the tables below | v) |
| | Infiltrate or Cavitary le Nodule or I margins (SI | Suggest Tuberculosis (Need consolidation sion mass with poorly defined uch as tuberculoma) usion (perform lateral or radiograph or ultrasound, if | nopathy Y thout calcificat | Other on DS 2054 Cardiac Musculoskeletal Other, specify in Remarks Other on DS 20 Pleural thicl Diaphragma Calcified pu | | | not mark as Class | ing y | |
| Re | marks | | | | | | | | |
| | Radiolo | ogist's Name (<i>Printed</i>) | Radiologist's S | ignature <i>(Req</i> | uired) | | Date Inter | preted (mm-dd-yyy | y) |
| 4. | • | rs and Cultures Decision | | | | | | | |
| l | X-ray X-ray Yes, are in Signs Chest Know | licated - Applicant has no sign Normal or 'No specimens req Normal or 'No specimens req dicated - Applicant has (<i>Mark</i> or symptoms of TB : X-ray suggests TB n HIV infection of treatment cultures | uired' and test for cell-media uired' and test for cell-media | ated immunity t | o TB negati | | | | |
| 5. | Sputum Smea | rs and Cultures Results | | | | | | | |
| | Sputum Smear Results | Date specimen obtained (mm-dd-yyyy) 1. 2. | Date specimen reported (mm-dd-yyyy) | Positive | Negative | | | | |
| | Troduito | 3. | | | | | | | |
| | Sputum Culture | Date specimen obtained (mm-dd-yyyy) | Date specimen reported (*Date of exam on D | | Positive | Negative | NTM | Contaminated | |
| | | 2. | | | | | | | |

| 6. Tuberculosis Classification Applicants may have more than one TB Classification. However, they cannot be classified as both Class B1 TB and Class B2 TB. In addition, applicants cannot be classified as Class B3 TB, Contact Evaluation if they are Class A or Class B1 TB, Extrapulmonary. | | | | | | | | |
|--|--|--|--|--|--|--|--|--|
| | No TB Classification CXR not suggestive of tuberculosis, no signs or symptoms, no known HIV infection, TST or IGRA negative (if performed), not a contact | | | | | | | |
| | Class A Applicant has tuberculosis disease | | | | | | | |
| | Class B1 TB, Pulmonary CXR suggests tuberculosis, or signs and symptoms, or known HIV infection and sputum smears and cultures are negative and not a clinically diagnosed case. | | | | | | | |
| | Class B1 TB, Extrapulmonary Applicants with evidence of extrapulmonary tuberculosis. The anatomic site of infection should be documented. | | | | | | | |
| | Anatomic Site of Disease No treatment Current treatment Completed treatment | | | | | | | |
| | Class B2 TB, LTBI Evaluation Applicants who have a tuberculin skin test ≥10 mm or positive IGRA but otherwise have a negative evaluation for tuberculosis. Contacts with TST ≥5 mm or positive IGRA should receive this classification (if they are not already Class B1 TB, Pulmonary). | | | | | | | |
| | □ No LTBI treatment □ Current LTBI treatment (Indicate medications in Part 7) □ Completed LTBI treatment (Indicate medications in Part 7) | | | | | | | |
| | Class B3 TB, Contact Evaluation Applicants who are a recent contact of a known tuberculosis case. | | | | | | | |
| | □ No preventive treatment □ Current preventive treatment (Indicate medications in Part 7) □ Completed preventive treatment (Indicate medications in Part 7) | | | | | | | |
| | Source Case: | | | | | | | |
| | Name | | | | | | | |
| | Alien Number | | | | | | | |
| | Relationship to Contact | | | | | | | |
| | Date Contact Ended (mm-dd-yyyy) | | | | | | | |
| | Type of Source Case TB (Mark only one and attach DST results) Pansusceptible TB MDR TB (resistant to at least INH and rifampin) Drug-resistant TB other than MDR TB Culture negative Culture results not available | | | | | | | |
| Remarl | KS | | | | | | | |
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| ☐ Applicant was on tubercu | with tuberculosis disea osis treatment at the ti | ase by the pan | |) : | | | ysician | | |
|--|---|----------------|-----------------|--|-----------|-----------------|---------|--|--|
| How was the diagnosis made | : Positive labora | tory tests [| ☐ Clinical diag | nosis | | | | | |
| Diagnostic Chest Radiograp | h | | | | | | | | |
| Facility performing chest i | adiograph. | | | | | | | | |
| r domey performing offeet | adiograpii. | | | | | | _ | | |
| Date Radiograph obtained | d (mm-dd-yyyy): | | | | | | | | |
| Findings Present | | | | | | | | | |
| Infiltrate or consolida | ation | | П мііі | ary findings | | | | | |
| Cavitary lesion | | | _ | | | | | | |
| | | , . | | crete linear op | acity | | | | |
| Nodule or mass with tuberculoma) | poorly defined margir | is (such as | ☐ Dis | crete nodule(s | s) withou | t calcification | | | |
| Hilar/mediastinal add | enonathy | | ☐ Vol | ume loss or re | etraction | | | | |
| _ | | | Oth | ier | | | | | |
| Pleural effusion | | | □ Noi | Normal or no findings suggestive of tuberculosis | | | | | |
| | | | | | 3 3 | 3 | | | |
| Sputum Smear Results Date specimen obtained | Date specimen | renorted | | | | | | | |
| (mm-dd-yyyy) | (mm-dd-y) | | Positive | Negative | | | | | |
| 1. 2. | | | | | | | | | |
| 3. | | | | | | | | | |
| Sputum Culture Results | | | | | | | | | |
| Date specimen obtained | | | | | | | | | |
| (mm-dd-yyyy) | (mm-dd-) | vyyy) | Positive | Negative | NTM | Contaminated | | | |
| <u>1.</u> 2. | | | | | | | | | |
| 3. | | | | | | | | | |
| Drug Susceptibility Test Res | ults. Attach with DS F | orms. | | | | | | | |
| Method of | net· | Date specin | men obtained | tained Date DST reported y) (mm-dd-yyyy) | | | | | |
| ☐ MGIT ☐ Agar | LJ | (mm-dd-yyyy) | | (111111 C | iu yyyy) | | | | |
| | | | | | | | | | |
| | Drug | | | Susceptibl | e Res | sistant | | | |
| Required for Rifampin | | | | | | | | | |
| first-line DST Ethambutol | | | | | | | | | |
| | Pyrazinamide | | | | | | | | |
| Required for Amikacin | e | + | | | | | | | |
| multidrug- Capreomyo | in | | | | | | | | |
| resistant Para-amino | salycilic acid (PAS) | | | | | | | | |
| | olone, specify: | | | | | | | | |
| Other, spec | | | | | | | | | |
| | | | | | | | | | |
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7. Previous Tuberculosis Diagnosis and Treatment History for Applicants Diagnosed or Treated Through Panel Physician, Continued

Were molecular tests used in addition to the required sputum smears, cultures, and DST:

No

Yes (mark all that apply):

| | , | acterium culosis | | mpin stance | Isoniazid Resistance | | |
|-----------------------|----------|---------------------|----------|----------------|-------------------------|----------|---------------------------|
| Molecular Test | Positive | Negative | Positive | Negative | Positive | Negative | Second-Line Test |
| Hain Line Probe Assay | | | | | | | Performed, attach results |
| GeneXpert | | | | | | | |

| Treating physician or institution DGMQ-Designated DOT site: Non-DGMQ-Designated DOT site: Drug Dosage Start Date (mm-dd-yyyy) End Date (mm-dd-yyyy) Isoniazid Rifampin Ethambutol Pyrazinamide Other, specify: | Tuberculosis Treatment | | | | | | | | |
|---|-----------------------------------|-----------------|-------------------------|-----------------------|--|--|--|--|--|
| Non-DGMQ-Designated DOT site: Drug Dosage Start Date (mm-dd-yyyy) End Date (mm-dd-yyyy) Isoniazid Rifampin Ethambutol Pyrazinamide | Treating physician or institution | | | | | | | | |
| Drug Dosage Start Date (mm-dd-yyyy) End Date (mm-dd-yyyy) Isoniazid Rifampin Ethambutol Pyrazinamide | DGMQ-Designated DOT site: | | | | | | | | |
| Isoniazid Rifampin Ethambutol Pyrazinamide | Non-DGMQ-Design | nated DOT site: | | | | | | | |
| Rifampin Ethambutol Pyrazinamide | Drug | Dosage | Start Date (mm-dd-yyyy) | End Date (mm-dd-yyyy) | | | | | |
| Ethambutol Pyrazinamide | Isoniazid | | | | | | | | |
| Pyrazinamide | Rifampin | | | | | | | | |
| | Ethambutol | | | | | | | | |
| Other, specify: | Pyrazinamide | | | | | | | | |
| | Other, specify: | | | | | | | | |
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PAPERWORK REDUCTION ACT AND CONFIDENTIALITY STATEMENTS

PAPERWORK REDUCTION ACT STATEMENT

Public reporting burden for this collection of information is estimated to average 20 minutes per response, including time required for searching existing data sources, gathering the necessary documentation, providing the information and/or documents required, and reviewing the final collection. You do not have to supply this information unless this collection displays a currently valid OMB control number. If you have comments on the accuracy of this burden estimate and/or recommendations for reducing it, please send them to PRA_BurdenComments@state.gov

CONFIDENTIALITY STATEMENT

AUTHORITIES: The information asked for on this form is requested pursuant to Section 212(a) and 221(d) and as required by Section 222 of the Immigration and Nationality Act. Section 222(f) provides that the records of the Department of State and of diplomatic and consular offices of the United States pertaining to the issuance and refusal of visas or permits to enter the United States shall be considered confidential and shall be used only for the formulation, amendment, administration, or enforcement of the immigration, nationality, and other laws of the United States. Certified copies of such records may, in the discretion of the Secretary of State, be made available to a court provided the court certifies that the information contained in such records is needed in a case pending before the court.

PURPOSE: The U.S. Department of State uses the facts you provide on this form primarily to determine your classification and eligibility for a U.S. immigrant visa. Individuals who fail to submit this form or who do not provide all the requested information may be denied a U.S. immigrant visa. Although furnishing this information is voluntary, failure to provide this information may delay or prevent the processing of your case.

ROUTINE USES: If you are issued an immigrant visa and are subsequently admitted to the United States as an immigrant, the Department of Homeland Security will use the information on this form to issue you a Permanent Resident Card, and, if you so indicate, the Social Security Administration will use the information to issue a social security number. The information provided may also be released to federal agencies for law enforcement, counterterrorism and homeland security purposes; to Congress and courts within their sphere of jurisdiction; and to other federal agencies who may need the information to administer or enforce U.S. laws. More information on the Routine Uses for this collection can be found in the System of Records Notice State-24, Medical Records.