U.S. Dep

## U.S. Department of State

OMB No. 1405-0113
EXPIRATION DATE: XX/XX/XXXX
ESTIMATED BURDEN: 15 MINUTES
(See Page 2 - Back of Form)

			TUBERCULOSIS WORKSHEET							
	Photo	Surnames	Surnames Given Nam				Age			
Birth Date (mm-a			Document Type	Doo	cument Numb	per	Case or Alien Number			
		Mediated Immunity to Tub losis Technical Instructions,	erculosis when required, perform one t	type only, and attac	ch results					
IC		ned, mark which test: uantiFERON	QuantiFERON (indi	icate optimal densi	ty value	T-Spot ( <i>indicate</i>	spot count fo	or each)		
	□ т-	-Spot	Nil Control:			Nil Control: _				
Date	drawn <i>(mn</i>	n-dd-yyyy)	TB Antigen1:			Panel A:				
H	Positive Negative Indetermine	nate, Borderline, or Equivoca	TB Antigen2:			Panel B:				
2. CI		Indication (Mark all that app								
	Chest X- Age <u>&gt;</u> 15	Ray not indicated years	Known HIV infection Extrapulmonary tubero	culosis	History o	tory of Tuberculosis  Date Chest X-Ray				
الم	· -	symptoms of tuberculosis	IGRA positive				(mm-dd-yyyy)			
3. CI	nest X-Hay Normal F	<u> </u>	o complete all of Section 3) onormal Findings (Indicate ca		marking all t	that annly in the	tables below	()		
	INOIIIIai i		Need Smears and Cultures			es Not Need S				
	Infiltrate	or consolidation	Miliary findings	Miliary findings Discrete linear opacity Discrete nodule(s) without calcification Volume loss or retraction			Do Not Mark as Class B Other on DS-2054  Smooth pleural thickening (if at CPA, must confirm is not effusion [do lateral or decubitus radiograph or ultrasound]) Diaphragmatic tenting			
		markings suggestive of fibro								
닏	Cavitary		=							
Ш		<ul> <li>or mass with poorly defined (such as tuberculoma)</li> </ul>	=							
Pleural effusion				Irregular thick pleural reaction Other			Single or scattered calcified pulmonary nodule(s)			
	Hilar/med	diastinal adenopathy	Guilei					node(s)		
Radi	ologist's Re	emarks								
-		Radiologist's Name (Printed)	Badio	logist's Signature (i	Required)	Date I	nterpreted (m	nm-dd-vvvv)		
4. Sr		ears and Cultures Decision		iogioto oignaturo (i	- ioquii ou)	Dato	norprotod (m	iiii da yyyy		
	1		gns or symptoms of TB, no k	nown HIV infection	, and:					
	X-ray	Normal or 'No specimens re	equired' and test for cell-med	iated immunity to T	B negative (if	performed)				
	X-ray	Normal or 'No specimens re	equired' and test for cell-med	iated immunity to T	B positive (if	performed)				
П	Yes, are	indicated - Applicant has (Ma	ark all that apply):							
	·	s or symptoms of TB		Extrapulmonary	ТВ					
		st X-ray suggests TB	ੂ	End of treatment						
	Knov	vn HIV Infection	_							
5. Sp	outum Sme	ears and Cultures Results								
Γ		Date specimen obtained	Date smear result rep	ported	Posit	ive Negative	$\Box$			
	Sputum	(mm-dd-yyyy)	(mm-dd-yyyy)		FUSIL	ive negative	<u>,                                     </u>			
	Smear Results	1.								
	ricounto	2.								
		3.								
	Sputum	Date specimen obtained (mm-dd-yyyy)	Date culture result repo			ive Negative	NTM	Contaminated		
	Culture Results	1.								
Cult	Locuito	2.								
		3.								

Ар	perculosis Classification plicants can be both Class B1 and Class B3, or Class B2 and Class B3. However, other combinations of tuberculosis classifications are not rmitted.
	No TB Classification CXR not suggestive of tuberculosis, no tuberculosis signs or symptoms, no known HIV infection, TST or IGRA negative (if performed), not a contact
	Class A Applicant has tuberculosis disease
	Class B0, TB, Pulmonary Diagnosed with tuberculosis by the panel physician or presented to the panel physician while on tuberculosis treatment and successfully completed DGMQ-defined DOT
	Class B1 TB, Pulmonary CXR suggests tuberculosis, or tuberculosis 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 Started but did not finish extrapulmonary 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 If treated, LTBI treatment: If treated, mark LTBI regimen:  Current LTBI treatment LTBI treatment by panel physician Completed LTBI treatment LTBI treatment by non-panel physician Started but did not finish LTBI treatment Dates of treatment to 3HP Other
	Class B3 TB, Contact Evaluation Applicants who are a recent contact of a known tuberculosis case.
	No preventive treatment Window Prophylaxis Isoniazid Rifampin 3HP Other Dates of treatment to
	Source Case:
	Name
	Case or Alien Number, if known
	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  DST results not yet available
Rema	rks

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	previously diagnos	ed with (	Class A TB by the panel	ohysician	adical evaminat	ion		
			·	_		ion		
was the diagr	osis made:	ositive la	boratory tests	Clinical di	agnosis			
agnostic Ches	Radiograph							
Facility perfo	rming chest radiogr	aph:				_		
Date Radiog	raph obtained (mm-	-dd-yyyy)	):					
ndings Present								
Normal	or no findings sugg	estive of	tuberculosis		Hilar/mediastin	al adenopat	hy	
Infiltrate	or consolidation				Miliary findings			
Reticula	ar marking suggesti	ve of fibr	osis		Discrete linear	opacity		
_				_			calcification	า
_		orly defin	ed margine (euch ac	H	Volume loss or		Ja. 5.1. 15 at. 15 1	
		ony denin	ed margins (such as	님				
				_		oleural reacti	on	
(mm-dd-yyyy) (mm-dd-yyyy) (mm-dd-yyyy) (mm-dd-yyyy) (mm-dd-yyyy) (mm-dd-yyyy) 1.  2.  3.  Drug Susceptibility Test Results		Ц_	Other					
						1		
		Date results reported		Positiv	ve Negative			
(111111-0	ia-yyyy)		(mm-dd-yyyy)		Negative	_		
						-		
						-		
outum Culture I	Result at Diagnosis							
Date specimen obtained			Date results reported					
(mm-dd-yyyy)			(mm-dd-yyyy)		ve Negative	NTM	Contaminated	
rug Susceptibil	ty Test Results				l			
M	Date specimen obtained (mm-dd-yyyy)  utum Culture Result at Diagnosis  Date specimen obtained (mm-dd-yyyy)  g Susceptibility Test Results  Method of DST:  MGIT Agar LJ  Dr						d	
П мен Г			(111111 dd ))))	7	(111	n da yyyy		
IVIGIT	Agai L	J						
		Drug			Susceptible	Res	adenopathy  pacity s) without calcification paraction ural reaction  NTM Contam  The Conta	
3t-11116 D3 I				e specimen obtained Date DST reported (mm-dd-yyyy)				
Required for	Amikacin							
multidrug-								
multidrug- resistant	Capreomycin	ic acid (F	PAS)					
multidrug- resistant	Capreomycin Para-aminosalicyl		PAS)					
multidrug- resistant	Capreomycin Para-aminosalicyl Fluoroquinolone,		PAS)					
multidrug- resistant	Capreomycin Para-aminosalicyl Fluoroquinolone,		PAS)					
multidrug- resistant	Capreomycin Para-aminosalicyl Fluoroquinolone,		PAS)					
multidrug- resistant	Capreomycin Para-aminosalicyl Fluoroquinolone,		PAS)					
Dete this section Applicant was Applicant was Applicant was Applicant was was the diagnostic Chest Facility performance and the property of th	Capreomycin Para-aminosalicyl Fluoroquinolone,		PAS)					

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History of Class A TB, Continue	ed								
Were molecular tests used in add	ition to the	required s	putum sm	nears, cultu	ıres, and [	DST:			
No									
Yes (mark all that apply):									
	Ťuber	acterium culosis	Resis	mpin stance	Isoniazid Resistance				
Molecular Test	Positive	Negative	Positive	Negative	Positive	Negative			
Hain Line Probe Assay								Performed, attach results	
GeneXpert								Performed, attach results	
Other								Performed, attach results	
Tuberculosis Treatment									
Treating physician or institution									
Approved DOT site:									
☐ Unapproved TB treatment	site:								
Drug	Dosa	ıge			Start Date	e <i>(mm-dd-</i>	vyyy)	End Date (mm-dd-yyyy)	
Isoniazid									
Rifampin									
Ethambutol									
Pyrazinamide									
Other, specify:									

## PAPERWORK REDUCTION ACT AND CONFIDENTIALITY STATEMENTS

## PAPERWORK REDUCTION ACT STATEMENT

Public reporting burden for this collection of information is estimated to average 15 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

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