

Adult Day Center Participant Tuberculosis (TB) Risk Assessment

- Use this tool to identify asymptomatic adults (persons 18 years and older) who require testing for latent TB infection (LTBI).
- Test for LTBI using a Mantoux tuberculin skin test (TST) or an Interferon-Gamma Release Assay blood test (IGRA) (e.g., QuantiFERON®-TB Gold or T-SPOT®), unless an appropriately documented negative test dated within the past 90 days or appropriately documented positive test result is available.
- IGRAs are preferred for people who have received the bacille Calmette-Guerin (BCG)³ vaccine.
- A negative TST or IGRA does not rule out active TB disease.
- For persons with TB symptoms⁴, abnormal chest x-ray consistent with TB disease, or a positive TST or IGRA: **Evaluate for active TB disease by** obtaining a chest x-ray, symptom screen, performing a physical exam and if indicated,⁵ sputum testing (i.e., AFB smears, cultures and nucleic acid amplification).

Risk Assessment

Check the appropriate risk factor boxes below. LTBI testing is recommended for persons with any of the following risk factors.

Risk Factor	Yes	No
Close contact to someone with infectious TB disease		
Birth, travel, or residence in a country with a high TB rate (e.g., any country other than		
the United States, Canada, Australia, New Zealand, or a country in western or northern		
Europe)		
Immunosuppression, current or planned – includes but is not limited to HIV infection,		
organ transplant recipient, treated with TNF-alpha antagonist (e.g., infliximab,		
etanercept), steroid use equivalent to prednisone ≥15 mg/day for ≥1 month, other		
immunosuppressive medication use, poorly controlled diabetes mellitus, end stage		
renal disease		
Resident or employee of a high-risk congregate setting (e.g., correctional facility,		
health care facility, homeless shelter)		

¹ TST documentation must include the date of the test (i.e., month, day, year), the number of millimeters of induration (if no induration, document "0" mm) and interpretation (i.e., positive or negative).

² IGRA documentation should include the date of the test (i.e., month, day, year), the qualitative results (i.e., positive, negative, indeterminate or borderline) and the quantitative assay (i.e., Nil, TB and Mitogen concentrations or spot counts).

³ BCG vaccination is not a contraindication for TST or IGRA testing; disregard BCG history when interpreting test results.

^{4.} Cough that lasts 3 weeks or longer, chest pain, coughing up blood, weakness or fatigue, weight loss, no appetite, chills, fever, or sweating at night.

⁵Sputum testing is indicated for all patients with chest x-ray findings compatible with TB regardless of TST or IGRA results or certain TB symptoms. Please consult with a TB expert.

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Patient Name:				nt Date of Birth		
have reviewed the above in	nformation,	based on this	information, the p	atient requires	the follo	wing:
		testing indicat		Date		
Clinician Name:						
Clinic Name:			Clinic Phone:			
TB Blood Test (i.e., Interferon-Gamma Rel	lease Assay l	blood test [IG	RA])			
Name of TB blood test	□QuantiF	ERON®-TB		POT®		
Date of blood draw						
Results						
Interpretation of reading	□Positive	* □Negativ	ve □Indeterminat	e \square Borderline	(T-SPOT	® only
Quantitative Result						
performing a physical exam ar	nd if indicated	d sputum testi	3 disease by obtainin ng.	g a cnest x-ray, s	ymptom	screen,
performing a physical exam ar	nd if indicated	d sputum testi	ng.	TST – Second		screen,
performing a physical exam ar	nd if indicated	(TST)	ng.			screen,
Tuberculin skin	testing	(TST)	ng.			SCI EETI,
Tuberculin skin 1	testing	(TST)	ng.			SCI EEII,
Tuberculin skin to Administration Name of person administer	testing	(TST) TST – First S	ng.		Step □R fc	rearm
Tuberculin skin to Administration Name of person administered Date and time administered	testing	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	rearm
Tuberculin skin to Administration Name of person administered Date and time administered Location	testing	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	orearm
Tuberculin skin to Administration Name of person administered Location Tuberculin manufacturer	testing ring test d and lot #	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	orearm
Tuberculin skin the Administration Name of person administered Location Tuberculin manufacturer Tuberculin expiration date Signature of person administered	testing ring test d and lot #	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	orearm
Tuberculin skin to Administration Name of person administered Location Tuberculin manufacturer Tuberculin expiration date Signature of person adminitest	testing ring test d and lot #	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	orearm
Tuberculin skin the Administration Name of person administered Location Tuberculin manufacturer Tuberculin manufacturer Tuberculin expiration date Signature of person adminitest Results (read between 48-	ring test d and lot # stering 72 hours)	(TST) TST – First S	itep R forearm	TST – Second	Step □R fc	orearm

Reader's signature

^{*}Consult grid on <u>Candidates for Treatment of Latent Tuberculosis Infection (LTBI)</u> (https://www.health.state.mn.us/diseases/tb/candidates.pdf).

^{**}For persons with a positive TST: Evaluate for active TB disease by obtaining a chest x-ray, symptom screen, performing a physical exam and if indicated sputum testing.

^{***}If results are negative, perform the second step one to three weeks after First Step.

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Adapted by the Minnesota Department of Health TB Prevention and Control Program from materials produced by the Global TB Institute and the Francis J. Curry National TB Center

Minnesota Department of Health
www.health.state.mn.us/tb
4/16/2024

To obtain this information in a different format, call: 651-201-5414