

Tuberculosis Screening and Testing

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Who Should be Tested for TB Infection?

Targeted Testing for TB Infection

The simplified version:

- Persons who are at increased risk for M. tuberculosis infection
- Persons at increased risk for progression to active disease if infected with M. tuberculosis (even if not at increased exposure risk)



- Persons tested for administrative reasons (e.g., mandatory employment testing)
- Persons with symptoms of active TB disease (fever, night sweats, cough, and weight loss)



Who Should be Tested for TB Infection?

Targeted Testing for TB Infection



- Contacts of persons with active TB
- HIV positive individuals
- Immigants from high prevalence countries
- Injection Drug Users
- Residents and Employees of high risk congregate settings:
 - Correctional facilities and Homeless Shelters
 - Hospitals, Clinics, Nursing Homes, Substance Abuse Facilities
- Newest Category:
 - Patients considering treatment with TNF- α Antagonists
- Children exposed to high-risk adults or environments

Contacts of Individuals with Active TB



- Among close contacts to a TB Case:
 - 30% have TB Infection
 - 1-3% have active TB disease
- Without TB Infection treatment:
 - 10% with TB Infection with develop Active TB
 - Approximately 5% of contacts with newly acquired TB Infection progress to TB disease within 2 years
 - The other 5% activate > 2 years after acquisition
- Examination of contacts is one of the most effective strategies for TB Infection diagnosis and TB control!

Percent Risk of Disease by Age



Age at Infection	Risk of Active TB
Birth – 1 year*	43%
1 – 5 years*	24%
6 – 10 years*	2%
11 – 15 years*	16%
Healthy Adults	5-10% lifetime risk
HIV Infected Adults ⁺	30-50% lifetime

TB Infection Diagnostics



• TB Skin Test (TST)

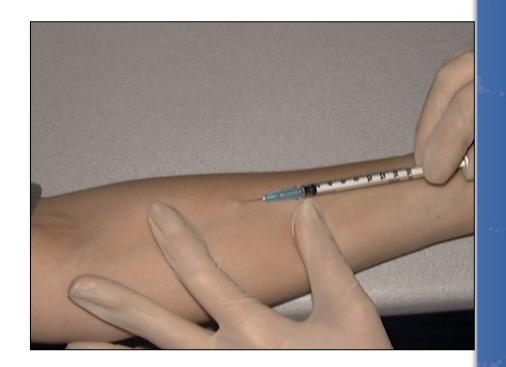
Interferon Gamma Release Assays (IGRA)

The Tuberculin Skin Test (TST)



0.1 ml of 5 TU PPD tuberculin injected intradermally

• **Induration** in millimeters read 48-72 hours after injection



Reading the TB Skin Test



Measure **induration**, not erythema!!!





TB Skin Test (TST)

• Pros:

- Inexpensive
- Simple to perform (if you know what you are doing.... and know how to herd cats......)

• Cons:

- Must return in 48-72 hrs
- Interpretation is somewhat subjective
- False Negatives:
 - Elderly
 - Immunosuppressed
- False Positives:
 - Low risk populations
 - Non-tuberculous mycobacteria
 - BCG vaccination



Classifying the Tuberculin Reaction

5 mm is classified as positive in



- Recent contacts of TB case
- Persons with fibrotic changes on chest radiograph consistent with old healed TB
- Patients with organ transplants and other immunosuppressed patients



Classifying the Tuberculin Reaction

10 mm is classified as positive in

- Recent arrivals from high-prevalence countries
- Injection drug users
- Residents and employees of high-risk congregate settings
- Mycobacteriology laboratory personnel
- Persons with clinical conditions that place them at high risk
- Children <4 years of age, or children and adolescents exposed to adults in high-risk categories



Classifying the Tuberculin Reaction



15 mm is classified as positive in

Persons with no known risk factors for TB

 Targeted skin testing programs should only be conducted among high-risk groups

Let's talk about IGRAs



Antigens for Newer Generation IGRAs

Negative control or nil (e.g., saline, heparin)



• Positive control or mitogen: non-specific immune response stimulator (e.g., phytohemagglutinin)

- *M. tuberculosis*-specific antigens
 - Unlike PPD used in TST, do not cross-react with BCG or NTM (some exceptions)
 - ESAT-6, CFP-10, TB 7.7 (actually simulated using overlapping peptides)

Antigens for Gamma-Release Assays



Tuberculosis	Antig	jens			
complex	ECAT	CED	Environmental	Antigens	
	ESAT	CFP	strains	ESAT	CFP
M tuberculosis	+	+	M abcessus		-
M africanum	+	+	M avium	-	2
	100703		M branderi	-	-
M bovis	+	+	M celatum	-	8
BCG substrain			M chelonae	-	2
gothenburg	-	1.2	M fortuitum	-	=
moreau			M gordonii	-	-
			M intracellulare	-	-
tice		7	M kansasii	+	+
tokyo		-	M malmoense		-
danish	-	_	M marinum	+	+
glaxo	322	16	M oenavense	-	-
	11.77		M scrofulaceum	-	2
montreal	+	•	M smegmatis	-	-
pasteur	3	-	M szulgai	+	+
			M terrae	-	-
			M xenopi	-	-

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FDA Approved IGRAs

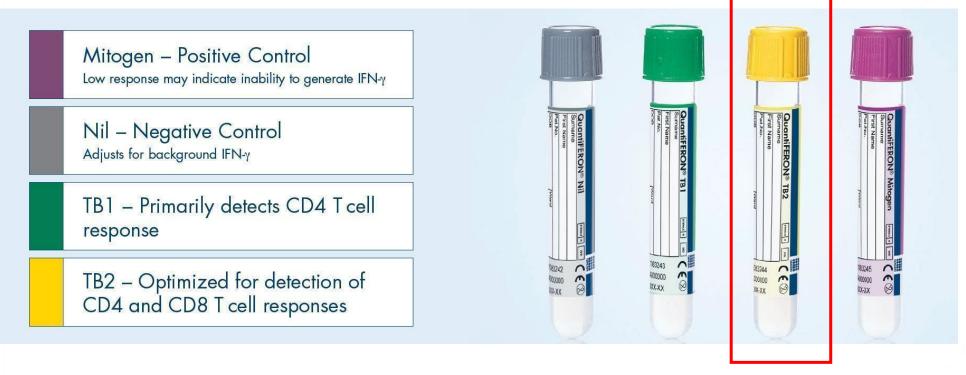


- QuantiFERON®-TB Gold Plus (QFT-Plus)
 - FDA approved 2017

- T-Spot[®].*TB* (T-Spot)
 - FDA approved July 2008

QuantiFERON®-TB Gold Plus





- > Essentially 2 tests in one blood draw
- > TB1 and TB2 should be close in value

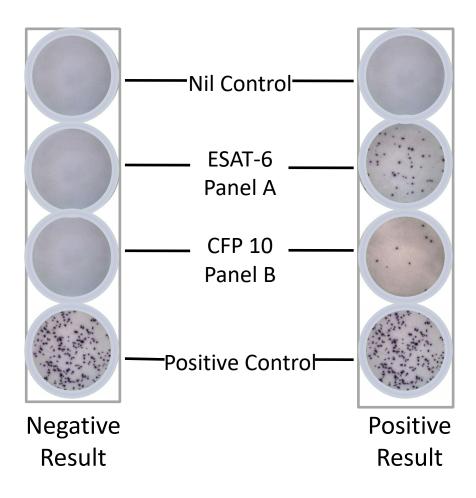
Interpretation Criteria for the QFT-GIT Test

Nil (IU/mL)	TB Antigen minus Nil (IU/mL)	QFT-GIT (IU/mL)	Mitogen	Interpretation
≤ 8.0	\leq 0.35 or $<$ 25% of Nil value	Negative	≥ 5.0	M. tuberculosis infection unlikely
≤ 8.0	\geq 0.35 and \geq 25% of Nil value	Positive	ANY	M. tuberculosis infection likely
≥ 8.0	ANY	Indeterminate	ANY	Indeterminate
≤ 8.0	≤ 0.35 and or < 25% of Nil value	Indeterminate	< 5.0	Indeterminate



Interpretation of Results





Interpretation Criteria for the T-Spot.TB

F	Result	Nil*	Response# Mitogen++		Interpretation+	
Р	ositive	≤ 10 spots	≥ 8 spots	Any	M.tuberculosis infection likely	
Во	orderline	≤ 10 spots	5, 6, or 7 spots	Any	Uncertain likelihood of <i>M. tuberculosis</i> infection	
N	egative	≤ 10spots	≤ 4 spots		M Tb infection unlikely	
Inde	terminate	> 10 ≤ 10	Any < 5 spots	Any < 20 spots	Uncertain likelihood of M. tuberculosis infection	



Indeterminate and Borderline Results



Indeterminate

- Negative control result is too high
 - High background production of IFN- γ
- Positive control result is too low
 - Immunocompromised patients may not respond to mitogen

- Borderline (T-Spot only)
 - Falls within borderline zone close to negative/positive cut point

(New) ATS/CDC/IDSA Guidelines



IDSA GUIDELINE







Official American Thoracic Society/Infectious Diseases Society of America/Centers for Disease Control and Prevention Clinical Practice Guidelines: Diagnosis of Tuberculosis in Adults and Children

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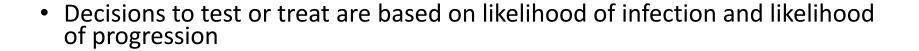
	Groups with Increased Likeli- hood of Infection with Mtb	Benefit of Therapy		LTBI Testing Strategy		
1	Household contact or recent expo- sure of an active case	Yes	Likely to be Infected Low to Intermediate Risk of Progression (TST ≥ 10mM)		Likely to be Infected High Risk of Pro- gression (TST ≥ 5mM)	
uomaa	Mycobacteriology laboratory personnel	Not demonstrated				
KISK OI INTECTION	Immigrants from high burden countries (>20 / 100,000)	Not demonstrated				
2	Residents and employees of high risk congregate settings	Yes				
ı	None	Not demonstrated	Unlikely to be Infe (TST > 15mM)	cted		
			Risk of Developing Tuberculosis if Infected			
			Low	Intermediate (RR 1.3 -3)	High (RR 3-10)	
			No risk factors	Clinical predisposition Diabetes Chronic renal failure Intravenous drug use	Children age less than 5 HIV infection Immunosuppres- sive therapy Abnormal CXR consistent with prior TB Silicosis	
			Benefit of Therapy			

In developing a diagnostic approach for the evaluation of those with suspected LTBI, we recommend the clinician weigh the likelihood of infection, the likelihood of progression to TB if infected, and the benefit of therapy (Horsburgh, C.R., Jr., and E.J. Rubin. 2011. Clinical practice. Latent tuberculosis infection in the United States. The New England journal of medicine 364:1441-1448). Recommendations were formulated for each of the three groups illustrated above. These groups are concordant with current recommendations for the interpretation of the TST (2000. Targeted tuberculin testing and treatment of latent tuberculosis infection. American Thoracic Society. MMWR Recomm Rep 49:1-51).

Not demonstrated

Yes

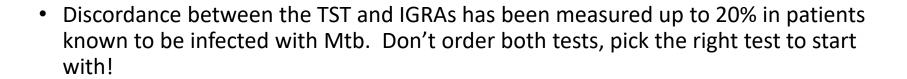
New in the Diagnosis Guidelines

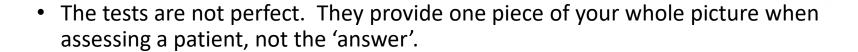




- IGRAs are recommended for testing for TB infection in individuals ≥ 5 years old with low or moderate risk if infection or progression
 - Note: IGRAs are a 'better' choice
 - When TST administration is questionable
 - In BCG vaccinated populations (increased specificity)
 - In populations with a poor rate of return
- Testing in low-risk populations is still not recommended. When it is necessary, such as required HCW screenings, use an IGRA
- In populations at high risk for infection or progression, either a TST or IGRA is appropriate

Pearls for TST vs. IGRAs





No test (TST or IGRA) overrides clinical, epidemiologic or historical data



