

Tuberculosis Disease: Laboratory Results for Nurses

Andrea Liptack, MSN, RN

TB Nurse Consultant, WI TB Program

Shea Rabley, MN, RN

TB Nurse Consultant, MCCT



Agenda

- Definitions, resources, background
- Most common active TB disease labs:
 - AFB smear microscopy
 - TB PCR/ NAAT
 - Culture
 - Susceptibility

- Other related labs you may see:
 - NTM cultures
 - IGRAs
 - Therapeutic drug monitoring (TDM)



Definitions of Abbreviations Used in This Presentation

AFB

Acid Fact Bacillus

MTBC

Mycobacterium tuberculosis complex

DST

Drug Susceptibility
Testing

NTM

Non-tuberculous Mycobacteria

MAC

Mycobacterium avium complex

TDM

Therapeutic drug monitoring

PCR and **NAAT**

Polymerase Chain Reaction

Nucleic Acid Amplification Testing

IGRA

Interferon Gamma Release Assay

Resources

- Sputum collection instructions (video by PHMDC, WI)
- Understanding TB Laboratory Testing for Public
 Health Nurses, APHL
- Xpert®MTB/RIF, Cepheid
- MDDR in MTBC by DNA Sequencing User Guide, CDC.
- MCCT webinar on therapeutic drug monitoring





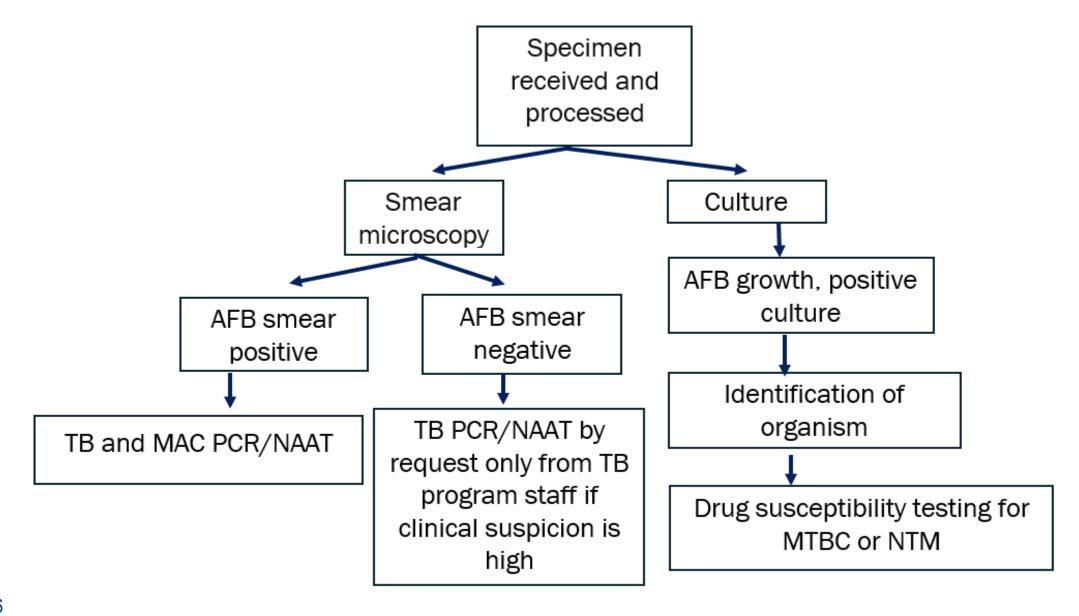
Mycobacterium tuberculosis Complex

- Rod shaped, acid-fast, aerobic, slow-growing, intracellular pathogenic bacteria
- MTBC includes:

M. tuberculosis, M. bovis, M. africanum, M. microti, M. caprae, M.pinnipedii, M. canettii.



Example Testing Algorithm







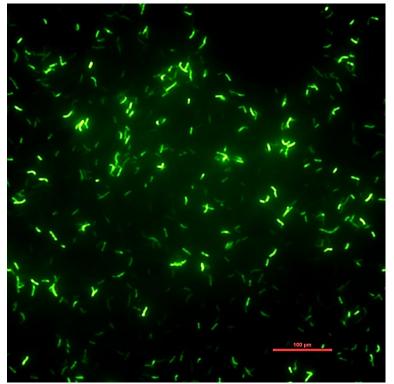
AFB Smear



Positive AFB Smear

This tells us the laboratorian saw AFB through the microscope. The waxy coating prevents the destaining agent from washing the stain off the TB.

What you might see through the microscope



Laboratory photos used with permission from Nate Simon, WSLH



- You can remember this by thinking that the lab worker is "smearing" the specimen onto a glass slide to look at in their microscope.
- Because of this type of bacteria's thick waxy coating, other organisms besides *Mycobacterium* tuberculosis may make an AFB smear positive.
- All this test type tells us is that there are bacteria in the specimen on this slide on the microscope that look like they may be TB.



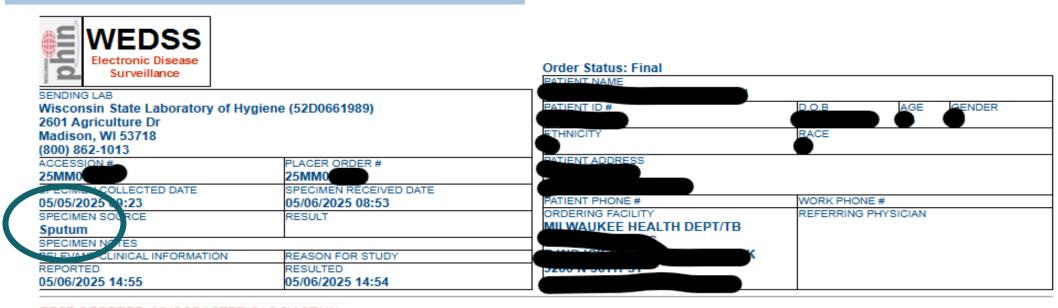
- This test is rapid and inexpensive for fast indication of TB disease but is not confirmatory.
- Most AFB smear positive test results in Wisconsin are for people with an NTM infection or with non-pathogenic (aren't causing illness) NTM in their respiratory tract.



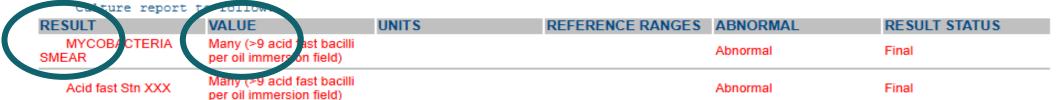
Smear Microscopy, Result Grading

Result	Graded Scale	Qualitative Scale	Interpretation, if infectious MTBC
Negative	Negative	Negative	Potentially infectious
Rare (1-9 AFB per 100 fields)	1+	Positive	Likely low-level infectiousness
Few (1-9 AFB per 10 fields)	2+	Positive	Likely moderately infectious
Moderate (1-9 AFB per field)	3+	Positive	Likely moderately infectious
Many (>9 AFB per field)	4+	Positive	Likely highly infectious

Smear Microscopy Example



TEST ORDERED: MYCOBACTERIOLOGY STAIN



Performing Organization: Wisconsin State Laboratory of Hygiene Performing Organization Address: MADISON, WI





Is this TB?

Maybe. Could be another fluorescing bacilli such as NTM (most common cause of an AFB smear positive test) and some legionella species (limited specificity).







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Is TB in there?

Maybe. Sputum specimen collection is one small sample of one small part of the lung during one moment in time. The person may still have TB, but no TB bacteria were seen on the AFB smear (limited sensitivity).



TB PCR/ NAAT §

Rapid Diagnosis



TB PCR

Detects MTBC

- Rapid identification of TB.
- Is confirmatory that TB DNA is present in the tested sample.
- Not performed serially (once only for rapid diagnosis).



TB PCR

Detects MTBC

Sensitivity for TB:

- Greater than 95% for AFB smear positive, culture confirmed.
- 55–75% for AFB smear negative, culture confirmed.



TB PCR

Detects MTBC

The lab may then perform TB PCR on newly smear positive respiratory specimens.

Newly smear positive (WI definition): no positive AFB smears or cultures in the last 12 months.

• Check with your state or regional consultant, they may need to order or "approve" TB PCR on smear negative specimens for high-risk individuals or order PCR testing on smear positive specimens.



TB PCR Result Interpretations

Example Lab Result	Interpretation
"Mycobacterium tuberculosis complex DNA detected"	Positive, TB identified
"Mycobacterium avium complex DNA detected"	Positive, MAC identified
"No Mycobacterium tuberculosis complex DNA detected"	Negative, TB not identified
"No Mycobacterium avium complex DNA detected"	Negative, MAC not identified
"Inhibitory substances that prevent nucleic acid amplification were detected"	Testing of no diagnostic help



TB PCR, Limitations ₹

Is this TB?

Yes! We have identified the DNA of the organism that is causing the positive smear as being present in the tested sample. TB is confirmed.



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Is TB in there?

If positive, yes; if negative, maybe. We are reliant on capturing a good specimen with lots of TB bacteria, and for this test, TB DNA must also be in the specimen at the time it was collected.



Culture



The Gold Standard of Diagnosis



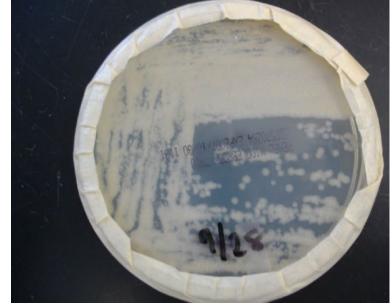
Culture

Positive MTBC Culture

This tells us that TB is alive and dividing in the growth media. Both solid and liquid growth media are available to labs. This is the most sensitive method of detection.

What you would see





Laboratory photos used with permission from Nate Simon, WSLH



Culture

Either liquid (broth) or solid growth media are used for the incubation period.

- Liquid broth: Mycobacteria Growth Indicator Tube (MGIT) is used, it is an automated instrument, it detects oxygen consumption through a florescent pad (glows and alerts lab tech if positive).
- Solid media: Middlebrook 7H11 plates are used for nonrespiratory and specimens from people with known TB. These are visually inspected at lease once per week for 6 weeks.

Culture, Example

Collected Type Source Submitter ID 25MM00 4/20/2025 0620 Sputum Mycobacteria Culture (Final result) Type/Src: Sputum

Mycobacteria Culture Result

Mycobacterium tuberculosis complex (A)

Comments:

The M. tuberculosis complex includes the species M. tuberculosis, M. bovis, M. bovis BCG, M. africanum and M. microti. 98% of the isolates in the M. tuberculosis complex identified at the State Laboratory of Hygiene are M. tuberculosis.

WSLH automatically performs culture-based drug susceptibility testing on an initial isolate from each TB patient.

A laboratory developed real-time PCR assay was used for identification from culture growth. This test was developed and its performance characteristics determined by the Wisconsin State Laboratory of Hygiene, a Clinical Laboratory Improvement Amendments (CLIA) certified, high complexity clinical laboratory. It has not been cleared or approved by the U.S. Food and Drug Administration.

Cultures are incubated for a total of six weeks. Further reports will be issued if additional Mycobacteria species are detected.



Culture, Limitations ©

Is this TB?

Yes! We have identified the organism growing and dividing in the growth media is MTBC. TB is confirmed. Isolation of MTBC almost always signifies disease.



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Yes! Since TB grew in culture, it is alive and viable. If the TB has been exposed to TB drugs for more than a few days, it is likely already experiencing changes that make it harder to grow and less infectious.



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Is TB in there?

Yes! Culture is the gold standard for TB diagnosis. It is the slowest method. It can take up to 42 days to wait to see if it is growing and dividing in the liquid and solid media (not good for rapid diagnosis).



Phenotypic and Molecular DST



DST, Phenotypic and Molecular

- Performed for all newly culture-confirmed specimens.
- Guides treatment plan and regimen such as medication changes and duration of therapy.
- Allows for appropriate treatment of contacts with appropriate regimens (susceptible to TB drugs).

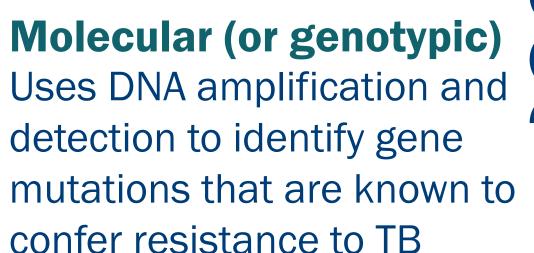


Phenotypic vs. Molecular DST

Phenotypic ()



Incubates a known concentration of MTBC in a known concentration of drug to observe if it grows or is inhibited from growing.

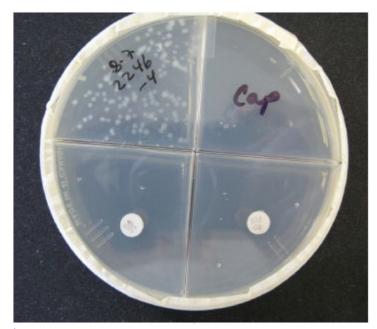


drugs.



Phenotypic vs. Molecular DST

Phenotypic DST, Agar



Laboratory photos used with permission from Nate Simon, WSLH

Molecular DST, GeneXpert ®

Type/Src: Aspirate/Neck

Result Units

MTB complex DNA detected. (A)

Rifampin gene mutation detected. (A)

Possible rifampin resistance.



DST, Phenotypic

- Usually takes 2-4 weeks after a positive culture.
- Phenotypic DST provides results for first line TB drugs: isoniazid (INH), rifampin (RIF), ethambutol (EMB).
- Pyrazinamide (PZA) results likely not available due to nationwide discontinuation of this testing.



DST, Phenotypic Result Interpretation

Result	Interpretation
Susceptible	TB strain is likely to show responsiveness to drug (TB will not grow and divide well or at all).
Resistant	TB strain is unlikely to show responsiveness to drug (TB will probably continue to grow and divide, evading the effect of the drug).



DST, Phenotypic Example

Susceptibility

Mycobacterium tuberculosis complex 1501

Isoniazid (0.) mcg/mL)
Isoniazid (1.0 mcg/mL)

WYCOBAC FRIAL SUSCEPTIBILTY
Susceptible

Susceptible

Results apply only to sample tested. **A**=Abnormal; **AA**=Panic; **H**=High. **L**=Low

Report ID:

Page: 1 of 2

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⊕ 05-12-2025 11:30 AM

UW Madison

⇒ WISCONSIN DIVISION OF PUBLIC HEALTH

pg 3 of 5



Wisconsin State Laboratory of Hygiene Communicable Disease Division 2601 Agriculture Dr; PO Box 7904 Madison, WI 53718 (800) 862-1013 Laboratory Report

James J. Schauer, Ph.D., PE, MBA, Director Errin C. Rider, Ph.D., D(ABMM),M(ASCP)CM, Director of Clinical Laboratory Services

Rifampin (1 mcg/mL)
Ethambutol (5 mcg/mL)
Pyrazinamice (100
mcg/mL)

Susceptible Susceptible Susceptible

Iso1 - Mycobacterium tuberculosis complex:

This is a final report.

Reported results were determined using the MGIT 960 Broth System.

Antibiotic values (mcg/ml) shown are the Equivalent Reference Method concentrations.



DST, Molecular

If TB PCR positive, rapid molecular testing can be performed.

- Cepheid GeneXpert® MTB/RIF testing is performed.
- Predicts if the TB bacteria will be susceptible to rifampin (fastest predictor of MDR-TB).
- Detects mutations in the rpoB genetic loci in 1–2 business days. A mutation in the rpoB gene loci predicts rifampin resistance.



DST, Molecular

Other considerations

- May detect "silent" mutations or mutations of unknown significance— ones that do not confer resistance.
 Phenotypic DST confirms resistance.
- GeneXpert® only predicts resistance to rifampin.
- Performed on primary respiratory specimens— you need a culture to grow to perform this testing on nonrespiratory specimens.



DST, Molecular Result Interpretation

Result	Interpretation
MTB DETECTED; Rif Resistance DETECTED	Likely TB is resistant to rifampin (rifampin likely can not be used as an effective drug in the regimen).
MTB DETECTED; Rif Resistance NOT DETECTED	Likely TB is susceptible to rifampin (rifampin can be used as an effective drug in the regimen).

DST, Molecular Example

TEST ORDERED: CEPHLID GENEXPERT MTB/RIF

Performing Organization: Wisconsin State Lab ratory of Hygiere

Performing Organization Address: MADISON, WI

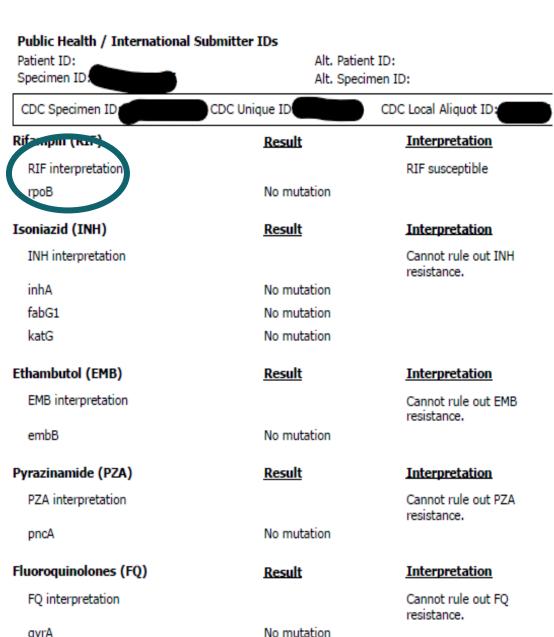
This real-

time PCR assay simultaneously detects the presence of both M. tuberculosis complex DNA and genetic mutations associated with rifampin resist ance from raw (induced or expectorated) or concentrated sputum sediments.

The assay is FDA cleared only for sputum specimens from patients for whom there is clinical suspicion of tuberculosis and who have received no antituberculosis therapy, or less than three days of therapy. It has not been approved for specimens from pediatric patients or to be use d as a test of cure.

RESULT	VALUE	UNITS	REFERENCE RANGES	ABNORMAL	RESULT STATUS
MYCOBACTERIUM TUBERCULOSIS COMPLEX TARGET GENE	MTB compl x DNA detected.			Abnormal	Final
M TB rpo 3 XXX QI PCR	N TB complex DNA detected.			Abnormal	Final
RESULT	ALUE	UNITS	REFERENCE RANGES	ABNORMAL	RESULT STATUS
RPOB GENE MUTATION	No rifampin gene mutation detected				Final
M TB rpoB XXX QI PCR	No rifampin gene nutation detected.				Final
RESULT	VALUE	UNITS	REFERENCE RANGES	ABNORMAL	RESULT STATUS
RIFAMPIN INTERPRETATION	Likely rifampin susceptible.				Final
M TB rpoB XXX QI PCR	Likely rifam in susceptible.				Final





No mutation

DST, Molecular, CDC Expanded Panel

Molecular detection of drug resistance (MDDR) can be done at CDC. This provides an expanded result panel for first- and secondline TB drugs. This testing will likely need to be requested by the TB program, provider, or sending lab. It may be automatically sent out based on first line resistance results depending on your lab.

gyrA gyrB



Other TB Labs

NTM and IGRA



NTM PCR and Cultures

NTM are commonly found in soil and water

Mycobacterium peregrinum (A)
Mycobacterium avium complex (A)

Comments:

SUPPLEMENTAL REPORT: 4/23/2025

Please note that M. avium complex has been isolated in addition to M. peregrinum reported on 4/22/2025.

RESULT	VALUE
MYCOBACTERIA CULTURE RESULT	MYCOBACTERIUM AVIUM COMPLEX
Mycobacterium XXX Cult	Mycobacterium, avium-intracellulare group
Mycobacterium	avium complex

Can cause similar symptoms to TB disease in those with underlying lung disease or immune compromise. Are not contagious from person to person. May require treatment with antibiotics. MAC is most common NTM.

IGRAs

QuantiFERON®-TB Gold (QFT)

Detects CD4 and CD8 T cell interferon-gamma response to MTB antigens.

Four numeric results: positive, negative, indeterminate.

T-SPOT ®.TB

Detects memory T cells that produce interferon-gamma in response to MTB antigens.

Results as number of "spots", positive, negative, or borderline.



QFT, Numeric Vales

Mitogen	Positive Control: Ensures the sample can respond to interferon gamma (baseline immune response).
Nil	Negative control: Adjusts for background interferon gamma.
TB1	Primarily detects CD4 T cell response.
TB2	Optimized for CD4 and CD8 T cell detection.

IGRA, Result Interpretation

IGRA Test Result	QuantiFERON	T-Spot	Notes
Positive	TB-Nil is higher than or equal to 0.35 IU/mL	8 spots or more	Infection is likely in individuals with risk factors. Consider retesting in low- or norisk individuals.
Negative	TB-Nil is lower than 0.35 IU/mL	4 spots or less	Infection is unlikely.
Indeterminate or invalid	High nil value or low mitogen value	High nil value or low mitogen value	Collect another specimen for retesting since these results cannot be interpreted. This occurs if controls do not perform as expected.
Borderline (not clear)	Not applicable	5, 6 or 7 spots	Uncertain likelihood of TB infection. Collect another specimen for retesting.



QFT, Example

TEST ORDERED: QUANTIF		
RESULT	VALUE	UNITS
QUANTIFERON PLUS INTERPRETATION	Positive	
M TB IFN-g Bld-Imp	POSITIVE	
sease(e.g., AFB s	nfection possible. This mear and culture, chest	t x-ray).
	Services for Interpret:	
RESULT	VALUE	UNITS
PNIL	0.15	[IU]/mL
Gamma interferon background Bld IA-aCnc	0.15	[IU]/mL
RESULT	VALUE	UNITS
P TB AG-1 NIL	0.52	[IU]/mL
M TB IFN-g CD4+ bckgrnd cor Bld-aCnc	0.52	[IU]/mL
RESULT	VALUE	UNITS
P TB AG2-NIL	0.54	[IU]/mL
M TB IFN-g CD4+CD8+ bckgrnd car bid aCnc	0.54	[IU]/mL
RESULT	VALUE	UNITS
P MITOGEN-NII	0.28	[IU]/mL
Mitogen TorvF bckgrd cor Bld-aCnc	0.28	[IU]/mL



Resources for IGRAs

- SNTC Webinars: IGRA for Nurses, webinar recording
- MCCT: All About IGRA for Civil Surgeons, webinar recording
- QuantiFERON®-TB Gold Plus, Qiagen
- <u>T-SPOT Technology (ELISPOT)</u>, Revvity



Therapeutic Drug Monitoring

- TDM is the measurement of the level of certain drugs in the blood at designated time intervals to assure that dosing of the medications meets the specific targets.
- TDM is recommended when a person is experiencing a slower than expected response to TB treatment. TDM is not usually recommended as a routine part of monitoring for PWTB.

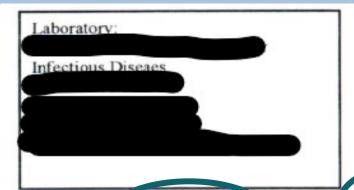


Therapeutic Drug Monitoring

- Blood is drawn two hours and six hours after the patient takes their TB medications (if on standard RIPE therapy).
- When drug levels are lower than expected, medication doses are adjusted.
- Not all laboratories in the U.S. offer TDM. If TDM is needed for a patient, consult with your state TB Program for guidance.



Therapeutic Drug Monitoring, Example



Patient name
Patient's facility ID numbers
Sample draw date: 4/1/2025
Sample draw time: 1013
Date of last dose: 4/1/2025
Time of last dose: 0800
Amount of last dose: 1200 mg
Frequency: Daily

RIFAMPI (RIF) Concentration in mcg / nL): 7.68

If the time of the dose and blood draw were not accurately recorded, accurate interpretation of the concentration is not possible.

The normal range for RIFAMPIN (RIF) serum or plasma concentrations is 8-24 mcg/ml approximately 2 hours after an oral dose. Samples later than 2 hours after the dose may display concentrations below the normal range. Two plus six hour post dose samples help to distinguish between malabsorption versus delayed absorption.

RIF has profound concentration-related activity, with higher concentrations showing better activity in vitro and in animal models. RIF and its desacetyl metabolite are not associated with concentration-related adverse effects, unlike RBN. RIF hepatotoxicity, etc., does not have a clear association with plasma concentrations.

Hepatic dysfunction may produce elevated RIF concentrations. In most patients, renal dysfunction does not affect RIF concentrations. Concurrent medications may have reduced concentrations in the presence of RIF, and also may require monitoring.





Thank you! TB laboratory staff and Nate Simon, PhD, TB Lab Coordinator at Wisconsin State Laboratory of Hygiene for info, lab photos, and review