

STANTON TERRITORIAL HEALTH AUTHORITY

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TITLE: Mycobacteria Reporting	Revision Date:	Issue Date:
	07-April-2017	07-April-2015
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Approved by:	Signed by:	
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Yellowknife, Northwest Territories

PURPOSE:

To standardize the reporting of direct smear results using Fluorescence stain. Opening of positive culture tubes for Kinyoun smear is no longer practiced at Stanton due to Containment Level restrictions. Positive cultures are reported out finalized as such and are referred to Provincial Lab in Edmonton.

INTRODUCTION:

The presence of an active and highly infectious Mycobacteria case in clinical samples is determined by the Direct Smear, and confirmed by culture. The presence of latent Mycobacteria infection is determined from direct smear negative but culture positive cases.

The option to send positive Direct Smear concentrates to Prov Lab Edmonton for PCR for MtB identification and Rifampin resistance detection is optional and up to the physician in charge. Historically Rifampicin resistance is not prevalent in the Northwest Territories.

In Direct Smear negative patients, PCR MtB probe is not recommended as a confirmatory method to Direct Smear results. Direct Smear Negative samples, even if they contain Mycobacteria, may contain such low numbers of AFB that even amplification techniques give unreliable results.

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Positive Cultures will be referred out to Provincial Lab. New Mycobacteria guidelines advise to prepare culture smears in a CL-3 Lab, which Stanton Bacteriology is currently not.

SPECIAL SAFETY PRECAUTIONS:

- Handle all patient samples and testing reagent using "Routine Practices"
- Please refer to the Northwest Territories Infection Prevention and Control Manual, March 2012
- Prior to testing all patient are to be identified as per I-0500 Use of Two Patient Identifiers.

DIRECT SMEAR MISCROSCOPY (AUROMINE-RHODAMINE):

- Read the QC slide first. If results are as expected, the patient sample smears can be read. If not, re-stain all smears with a new QC slide.
- Refer to Expected Results chart above in the Quality Control of AFB Smears section.
- Read smears under DRY objectives. Scan using 20 x and move to 40 x objective if suspicious bacilli are seen.
- If AFB are seen in the direct smear, or if Tech is in doubt of AFB presence on smear, perform a Kinyoun smear to confirm for presence/absence of suspicious bacilli. Do not use Kinyoun smear for Direct Smear quantification.

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INTERPRETING DIRECT SMEAR RESULTS:

• Direct Smears are interpreted in a graded format for quantities

lf	Then	Action			
AFB is quant	AFB is quantified and interpreted following the chart below:				
AFB seen	Smear	Confirm direct s morphological o • Kinyour smear o consum Quantify accord	smear with characterist in can be do can be mad ling; must le ding to the I <u>Direct Sn</u> # of AFB	a Kinyoun stain. Kinyo ics of the bacilli. ne directly over the A/ e from the concentrate et new concentrate dry Direct Smear Interpret near Interpretation C Quantity Reported	Coun offers better (R stain or a new e (more time y and fix). cation Chart below. Chart STAFB Keypad code
	positive	Entire smear 10	< 6 1-9	Negative (no AFB)	AFB1
		1	1-9	2+ (few)	AFB6
		1	10-90	3+ (moderate)	AFB7
		1	>90	4+ (numerous)	AFB9
		Record result in	n ink on the	TB worksheet.	
		Log into Soft to result the Direct Smear.			
No AFB		Kinyoun smear not required unless Tech is unsure of direct			
-OR-	Smear	smear results.			
<6 bacilli	negative	Record result in ink on the TB worksheet.			
seen		Log into Soft to result the Direct Smear.			

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RESULTING DIRECT SMEARS IN LIS:

Step	Prompt	Action			
Follow	Follow the steps below to Result AFB direct smears:				
1	Result Entry	Type in order # of sample to access log			
2	Locate <mark>"STAFB"</mark>	Click Ctrl+K to access keypad if it is not already open. • From STAFB keypad chose the option corresponding to Direct Smear chart above.			
4	Status the Test (Finalize	STAFB line).			
5	Click Instant Report button	A SDR rule will automatically fire, asking to send a report to HPU. Click "No" for now. You'll have another option to do so when you save the report.			
6	Preview Instant Report. Stain, Din 23/01/15 Microscopi Direct Sme Acid Fast <u>Culture, J</u>	Example below: <pre>rect AFB Smear & FINAL 23/01/15 14:11 ic Report: bar was examined by Fluorescent stain. Bacilli WERE NOT seen. AFB - PLATED</pre>			
7	Save Culture	SDR violation box automatically pops up again. Clicking "Yes" will CC the report to HPU. This is desired in patient samples, but not TB Weekly (QC) cultures. Do Not click "Yes" for any QC samples.			

File the TB Worksheet in the appropriate slot in the water room.

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CULTURE SMEAR MISCROSCOPY (KINYOUN STAIN) :

- Smears from culture are not performed due to CL restrictions
- Perform Kinyoun stains on Direct Smears that are difficult to interpret or for confirmation of AFB seen in Fluorescent stain
- Kinyoun stain may be done directly over top of the fluorescent smear. A new smear does not need to be made but can be done if desired. Remaking a new smear will require the smear to dry and be fixed again (may take an hour before staining)
- Make a QC slide for Kinyoun stain. After staining, read the QC slide first. If results are as expected, the patient sample smears can be read.
- Refer to Expected Results chart above in the Quality Control of AFB Smears section.
- Scan smears on 50x oil immersion and switch to 100x to confirm morphology.. Read the entire smear. This may take up to 45 mins to an hour in some smears.

RESULTING KINYOUN SMEARS MADE FROM DIRECT SMEARS:

• Kinyoun smears are examined for the presence or absence of AFB. Quantities for reporting should be graded using the results of the Fluorescent stain. Kinyoun should be used a confirmatory tool. Refer to Direct Smear guidelines above.

Step	Prompt	Action
Follov	v the steps below to result	t positive MGIT and LJ cultures:
1	Result Entry	Type in order # of sample to access log
2	Media Comments (plate log)	Locate "MGIT" or "LJ" in plate log below CXAFB. Click Ctrl+K to access keypad if it is not already open (see step below).

RESULTING POSITVE CULTURES IN LIS:

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3	MGIT Media line	Follow the keypad to add the positive growth culture to the
	Or	isolates tab.
	LJ Media line	
		Ensure the Quantitation (weeks to positivity) is entered and
		the Isolate comment is added "Culture will be
	Isolate Tab	referred out for further examination at
		ALBERTA PROVINCIAL LABORATORY OF PUBLIC
		HEALTH IN EDMONTON.
	Finalize the culture.	Click the "^F" button
		This provides the ability to transcribe results from Prov
	Add a referral test code:	Lab. See Miscellaneous Tests Procedures.
	RFFTB	R Add Test Significant T Test Common Med
	NET TB	# Test D Test Comment M + 1 1 STAFB
		REFTB
		A SDR rule will automatically fire, asking to send a report
		to HPU. Click "No" for now. You'll have another option to
		do so when you save the report.
	Click Instant Report	Standard Deviation 2
	button	STAFE - TSTAFE:F /AUTOPR IPRHP1
		Do you want to accept this rule?
		Yes No
	Droview Instant Depart	Culture, AFB should display everything in the Isolates Tab.
	Fleview Instant Report	Referral Test code should say "Plated".
		SDR violation box automatically pops up again.
		Clicking Yes CC the report to HPU. This is desired in
	Save Culture	patient samples, but not for AFB quality control cultures.
		If resulting a patient test, click "Yes".
		If resulting an AFB control, click 'No" (do not want the
		results to be faxed to HPU).

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RESULTING 7 WEEK NEGATIVE CULTURES IN LIS:

Step	Prompt	Action
Follow	w the steps below to result	AFB cultures:
1	Result Entry	Type in order # of sample to access log
2	Media Comments	Locate "MGIT" or "LJ" in plate log below CXAFB.
	(plate log)	Click Ctrl+K to access keypad if it is not already open (see
	(plate log)	step below).
3	MGIT Media line	MGIT – Negative (MGIT will have automatically crossed
	Or	the Negative result over).
	L.I Media line	LJ – Manually result. Follow the keypad to add the
		negative growth comment "No growth after 7 weeks".
		Follow keypad CXTB1 (Urines) or CXTB2 (Resp) to result
	CXAFR Test Comment	negative cultures \rightarrow AFB2 code
	Of an D Foot Comment	"Mycobacteria Cultures were Negative after 7
		weeks incubation".
	Finalize the culture.	Click the " ^F " button
A SDR rule will au		A SDR rule will automatically fire, asking to send a report
		to HPU. Click "No" for now. You'll have another option to
		do so when you save the report.
	Click Instant Report	Standard Deviation Rule Violation
	button	STAFE - TSTAFE:F. /AUTOPR IPRHP1 STAFE - TSTAFE:F. /AUTOPR IPRHP1 Do you want to accept this rule?
	Preview Instant Report	
		SDR violation box automatically pops up again.
	Save Culture	Clicking Yes CC the report to HPU. This is desired in
		patient samples, but not for AFB quality control cultures.

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REFERRAL OF POSITIVE CULTURES TO PROVINCIAL LAB:

Positive MGIT or LJ cultures should be sent out as Category A. See referral manual for instructions. Provincial Lab referred in isolate forms are available on the shared drive or pre-printed by the Bacteriology door.

RELATED DOCUMENTS

- MIC81500 Auramine-Rhodamine Stain
- MIC81600 Kinyoun Stain
- See Section 5 of Specimen Management Manual Referrals

REFERENCES:

- Northwest Territories Health and Social Services. (2014). *NWT Tuberculosis Manual.* Yellowknife: Northwest Territories Health and Social Services.
- Ontario Mycobacteriology Bench Manual, Central Public Health Laboratory, 2003.

REVISION HISTORY:

REVISION	DATE	Description of Change	REQUESTED BY
1.0	3-FEB-2015	Initial Release	L. Driedger
	03Feb2015	Review	S. Webber