Next Review Date: 14/05/2026

Type: Laboratory Services Program SOP

Policy Number: 15-160-V1 Date Approved: 14/05/2024

PROGRAM Standard Operating Procedure – Laboratory Services			
Title: MIC20300 – Gram stain reporting in LIS-Respiratory Specimens	Policy Number: 15-160-V1		
Program Name: Laboratory Services			
Applicable Domain: Lab, DI and Pharmacy Services			
Additional Domain(s): NA			
Effective Date: 14/05/2024	Next Review Date: 14/05/2026		
Issuing Authority: Director, Laboratory and Diagnostic Imaging Services	Date Approved: 14/05/2024		
Accreditation Canada Applicable Standard: NA			

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GUIDING PRINCIPLE:

The culture of poorly collected respiratory specimens is a wasteful use of laboratory resources and can lead to erroneous reporting and treatment of patients. These specimens need to be scored for acceptability using the Q-score method.

PURPOSE/RATIONALE:

This standard operating procedure describes how to report the gram stain results of respiratory specimens in the LIS in a consistent manner.

SCOPE/APPLICABILITY:

This standard operating procedure applies to Medical Laboratory Technologists (MLTs) reporting the gram stain of respiratory specimens in the LIS.

SAMPLE INFORMATION:

Type •	Sputum, endotracheal aspirates (ETT) and auger suction specimens are Q-scored for quality Bronchial aspirates (washings), bronchoalveolar lavage (BAL) specimens, specimens collected from sterile catheter down ETT and specimens from cystic fibrosis patients are NOT Q-scored for quality
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REAGENTS and/or MEDIA:

- Methanol
- Gram Crystal Violet
- Gram Iodine (Stabilized)
- Gram Decolorizer
- Gram Safranin

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SUPPLIES:

Glass microscope slide

QC slide

Immersion oil

Slide storage tray

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EQUIPMENT

- Hot plate
- Microscope

SPECIAL SAFETY PRECAUTIONS:

Containment Level 2 facilities, equipment, and operational practices for work involving infectious or potentially infectious materials or cultures:

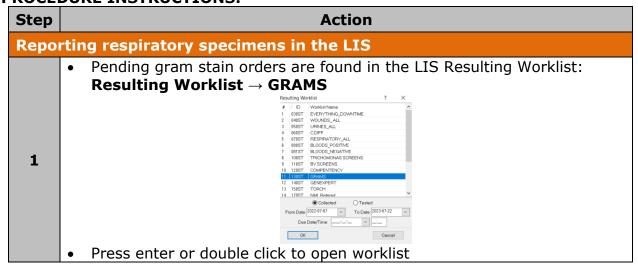
- Ensure that appropriate hand hygiene practices be used
- Lab gown must be worn when performing activities with potential pathogens
- Gloves must be worn when direct skin contact with infected materials is unavoidable
- Eye protection must be used when there is a known or potential risk of exposure of splashes
- All procedures that may produce aerosols, or involve high concentrations or large volumes should be conducted in a biological safety cabinet (BSC)
- The use of needles, syringes and other sharp objects should be strictly limited

All patient specimens are assumed to be potentially infectious. Routine Practices must be followed. Since viable micro-organisms are used, all cultures must be handled with appropriate precautions. All equipment in contact with cultures should be decontaminated by appropriate methods.

QUALITY CONTROL:

- Quality control is performed daily
- A TQC order is automatically generated daily to record the QC results
- Refer to MIC60060-Microbiology Stain Quality Control

PROCEDURE INSTRUCTIONS:



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Enter the accession number on the slide and select enter to mark the

Select enter again to open Result Entry or double click on accession number to open

Under low power (X10, LPF): screen slide to locate good specimen areas to obtain an overall impression of cell types present.

- Observe slide for stain crystals:
 - If an excess of precipitated stain is observed, prepare another
 - > If precipitate continues, use freshly filtered crystal violet
- Determine if slide has been properly decolorized:
 - > Depending on the source of the specimen, the background should be generally clear or gram negative
 - > If white blood cells are present, they should appear completely gram negative
 - > If slide is over decolorized, prepare another smear
- Determine if thickness of smear is appropriate:
 - > For proper interpretation, areas must be no more than one cell thick, with no overlapping of cells. Prepare a new slide if unreadable
- Examine for evidence of inflammation:
 - > Determine areas representative of inflammation and areas of contamination with squamous epithelial cells

Under low power (X10, LPF): average the number of epithelial cells and white blood cells:

4

2

3

None seen	No cells seen
1+	< 1 cell seen
2+	1 - 9 cells seen
3+	10 - 25 cells seen
4+	> 25 cells seen

Calculate the Q-score of the specimen. The Q-score is calculated by assessing the quantity of epithelial cells and neutrophils. Examine 20 to 40 fields and interpret as follows:

5

Q-score Table				
Eni celle /I DE	White blood cells /LPF			
Epi cells/LPF	0	1-9	10-25	>25
0	Q 0	Q 1	Q 2	Q 3
1-9	Q-1	Q 0	Q 1	Q 2
10-25	Q-2	Q-1	Q 0	Q 1
>25	Q-3	Q-2	Q-1	Q 0

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			chial aspirates (washir	
6	bronchoalveolar lavage (BAL), specimens collected from sterile catheter down ETT and specimens from cystic fibrosis patients.			
	If the Q-score indicates the sample is of good quality (Q-score 1-3) or if the specimen/sample type is bronchial aspirate (washings),			
			sterile catheter down	, -
7	cystic fibrosis pat	ient, add one drop	of immersion oil to the	e slide. In a
		•	nce of inflammation or ine 20 to 40 fields to 6	
	morphology and		ine 20 to 40 heids to t	observe cen
	If the Q-score inc	dicates the sample is	s not of good quality,	
8			nchoalveolar lavage (E cystic fibrosis patient,	
		the slide to observe	•	, do not add
			antitate epithelial cells,	white blood
	cells, red blood ce	ells and bacteria as f	Ollows:	
		None seen	No cells seen	
		1+	< 1 cell seen	
		2+	1 - 9 cells seen	
9		3+	10 - 25 cells seen	
		4+	> 25 cells seen	
	NOTE: Bacteria are not reported if the Q-score indicates specimen is			
	unsatisfactory for culture and the sample is not a bronchial aspirate			
		_	e (BALs), from sterile	catheter down
	NOTE: Only report "None seen" for white blood cells and bacteria. If no			
	epithelial cells or red blood cells are seen, do not report this Under the test code: STGM2 , use the STGM2 keypad to report the			
		-		-
10	quantity of epithelial cells, white blood cells, red blood cells and bacteria if indicated by Q-score. Report cells in this order to maintain consistency			
	with reporting. Reporting Mixed oropharyngeal flora in respiratory gram stain:			
			d neither are predomi	
	intracellular, r	nixed oropharyngea	al flora can be reporte	d
11	 If smear has ≥2 morphotypes and one or more are predominant or intracellular, the predominant or intracellular morphotypes are reported 			
	individually and other morphotypes are reported as mixed			
	oropharyngea	l flora		

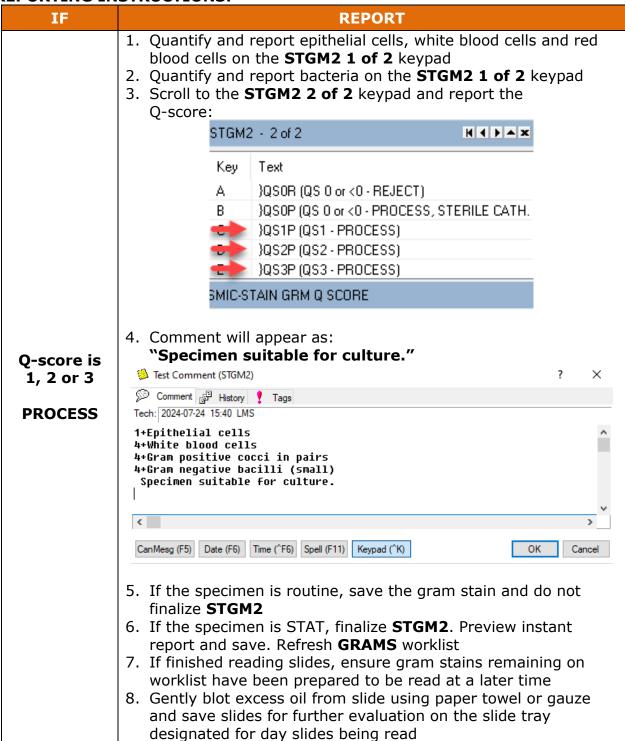
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REPORTING INSTRUCTIONS:



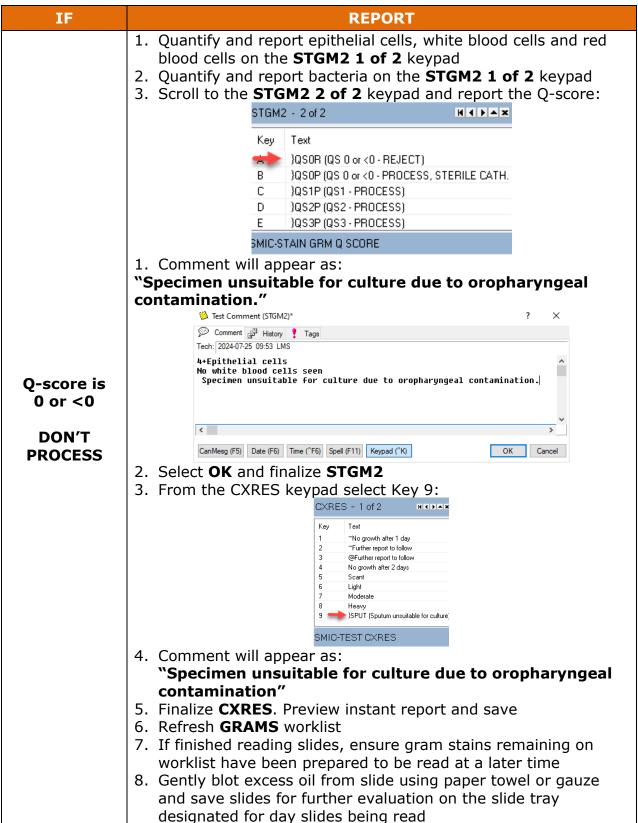
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te blood cells and red
ad
GM2 1 of 2 keypad

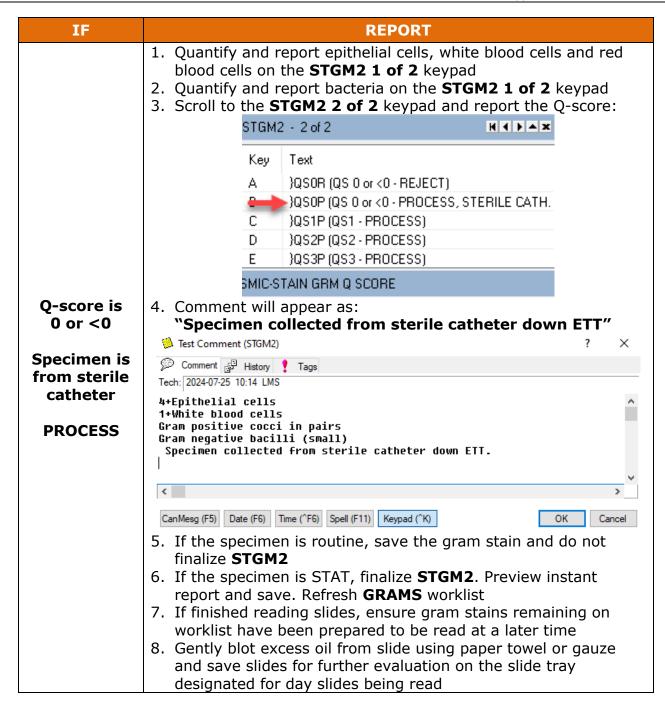


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IF	REPORT
Q-score is not performed	 Quantify and report epithelial cells, white blood cells and red blood cells on the STGM2 1 of 2 keypad Quantify and report bacteria on the STGM2 1 of 2 keypad
Sample is bronchial aspirate (washings) OR bronchoalveolar lavage (BALs)	 If the specimen is routine, save the gram stain and do not finalize STGM2 If the specimen is STAT, finalize STGM2. Preview instant report and save. Refresh GRAMS worklist If finished reading slides, ensure gram stains remaining on worklist have been prepared to be read at a later time Gently blot excess oil from slide using paper towel or gauze and save slides for further evaluation on the slide tray designated for day slides being read

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LIMITATIONS:

- 1. Use results of gram stains in conjunction with other clinical and laboratory findings. Use additional procedures (e.g., inclusion of selective media, etc.) to confirm findings suggested by gram stained smears.
- Careful adherence to procedure and interpretive criteria is required for accurate results. Accuracy is highly dependent on the training and skill of microscopists.
- 3. Gram stain positive, culture negative specimens may be the result of contamination of reagents and other supplies, presence of antimicrobial agents, or failure of organisms to grow under usual culture conditions (medium, atmosphere, etc.).
- 4. False gram stain results may be related to inadequately collected specimens or delays in transit.
- 5. Prior treatment with antimicrobial drugs may cause gram positive organisms to appear gram negative.

CROSS-REFERENCES:

- MIC10100-Microbiology Specimen Processing
- MIC60060-Microbiology Stain Quality Control

REFERENCES:

1. Leber, A. (2016). *Clinical microbiology procedures handbook.* (4thed.) Washington, D.C.: ASM Press

APPROVAL:

May 14, 2024
Date

Director, Laboratory and Diagnostic Imaging Services

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REVISION HISTORY:

REVISION	DATE	Description of Change	REQUESTED BY
1.0	07 Feb 19	Initial Release	L. Steven
2.0	31 Mar 22	Procedure reviewed and added to NTHSSA policy template	L. Steven
3.0	19 Feb 24	Procedure reviewed	L. Steven
4.0	25 July 24	Procedure updated to reflect new Q score reporting and not cancelling specimens	L. Steven
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