PROGRAM Standard Operating Procedure – Laboratory Services			
Title: MIC33200 – Genital Culture- Policy Number:			
Lower Genital Tract			
Program Name: Laboratory Services			
Applicable Domain: Lab, DI and Pharmacy Services			
Additional Domain(s): NA			
Effective Date:	Next Review Date:		
Issuing Authority:	Date Approved:		
Director, Laboratory and Diagnostic Imaging Services			
Accreditation Canada Applicable Standard: NA			

## **Uncontrolled When Printed**

## **GUIDING PRINCIPLE:**

Organisms which are associated with infection or disease of the genital tract include *N.gonorrhoeae* (GC), *Chlamydia trachomatis* (CT), *Trichomonas vaginalis* (TV) and Yeasts. Isolation of other organisms such as *S.pyogenes*, *S.agalactiae*, *S.aureus*, and others may be associated with certain specific clinical syndromes or risk of infection in the neonate.

### **PURPOSE/RATIONALE:**

This standard operating procedure describes how to determine the significance of growth in lower genital tract specimens.

#### **SCOPE/APPLICABILITY:**

This standard operating procedure applies to Medical Laboratory Technologists (MLTs) processing specimens for lower genital tract culture.

Туре	<ul><li>Swab</li><li>Amie's with or without charcoal</li><li>Charcoal swabs are recommended</li></ul>		
Source	• Labia • Penis • Vagina • Vulva		
Stability	<ul> <li>If the sample is received in the laboratory and processed greater than 24 hours from collection:</li> <li>Add specimen quality comment: "Delayed transport may adversely affect pathogen recovery"</li> </ul>		
Storage Requirements	Room temperature		

### SAMPLE INFORMATION:

	1. Unlabeled/mislabeled swabs
	2. Specimen container label does not match patient
	identification on requisition
Criteria for	3. Do not accept vaginal swabs from women >13 years of
rejection	age for genital culture unless significant clinical
	information is provided
	4. Do not process vaginal swabs for yeast culture unless
	significant clinical information is provided

**NOTE:** Genital culture is performed on vaginal specimens from patients  $\leq$ 13 years of age

### **REAGENTS and/or MEDIA:**

- Blood agar (BA) Chocolate agar (CHO), Columbia Naladixic Acid agar (CNA), Thayer Martin agar (TM) and MacConkey agar (MAC)
- Identification reagents: catalase, oxidase, Staph latex test, Strep latex test, etc.

### SUPPLIES:

- Disposable inoculation needles
- Microscope slides
- Wooden sticks

### **EQUIPMENT:**

- Biosafety cabinet
- 35° ambient air and 35° CO<sub>2</sub> incubators
- VITEK 2 and supplies

### **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:**

#### Refer to Test Manual for reagent quality control procedures

#### **PROCEDURE INSTRUCTIONS:**

Step	Action		
Proce	Processing specimens for lower genital tract culture		
1	<ul> <li>In the biosafety cabinet:</li> <li>Inoculate BA, CHO, CNA, TM, and MAC with the swab</li> <li>Ensure all surfaces of the swab make contact with the agar</li> <li>Streak for isolated growth using a disposable inoculation needle</li> <li>Prepare smear by rolling the swab gently across the slide to avoid destruction of cellular elements and disruption of bacterial arrangements</li> </ul>		
2	<ul> <li>Incubate all media:</li> <li>Place BA, CHO, CNA and TM in the CO<sub>2</sub> incubator</li> <li>Place MAC in the O<sub>2</sub> incubator</li> </ul>		
3	Allow smear to dry and perform gram stain. Gram stain must be read before culture plates. Refer to MIC20115-Gram Stain Procedure.		
4	<ul> <li>This procedure is divided into 2 sections that include the 4 sources of lower genital tract culture specimens:</li> <li>Vaginal and vulva culture</li> <li>Labia and penis culture</li> </ul>		

### 1. Vaginal Culture and Vulva Culture:

Probable Pathogens <sup>^</sup>				
<u>GNB Aerobic</u> :	<u>GNDC Aerobic</u> :			
<ul> <li>Salmonella spp.(≤13 yrs. old)</li> <li>Shigella spp. (≤13 yrs. old)</li> </ul>	Neisseria gonorrhoeae			
	GPB Aerobic:			
<u>GPC Aerobic</u> :	Listeria monocytogenes			
Staphylococcus aureus				
Streptococcus agalactiae	<u>Yeast</u> :			
Streptococcus pyogenes	Candida spp.			
Potential	Pathogens <sup>^</sup>			
<ul> <li>Gardnerella vaginalis</li> <li>Gram negative bacilli other than Enterobacteriaceae</li> <li>Haemophilus influenzae</li> </ul>	<ul> <li>Neisseria meningitidis</li> <li>Pseudomonas spp.</li> <li>Streptococcus pneumoniae</li> </ul>			
Comme	Commensal Flora			
<ul> <li>Anaerobes</li> <li>Coagulase-negative Staphylococci</li> <li>Corynebacterium spp.</li> <li>Enteric Gram-negative bacilli not listed above</li> </ul>	<ul> <li>Enterococcus spp.</li> <li>Lactobacillus spp.</li> <li>Non-pathogenic Neisseria spp.</li> <li>viridans Streptococcus grp.</li> </ul>			

<sup>^</sup> For organisms not listed, consult the Microbiology Technical Supervisor or refer to the *Manual of Clinical Microbiology* 

#### **INTERPRETATION OF RESULTS:**

Step	Action				
Step					
1	<ul> <li>Ensure growth on culture media correlates with gram stain results. If discordant results are found between the gram stain and growth:</li> <li>Re-examine smear and culture plates</li> <li>Check for anaerobic growth</li> <li>Re-incubate media to resolve</li> <li>Consider re-smearing or re-planting specimen</li> </ul>				
2	<ul> <li>Observe BA, CHO, CNA and TM plates at 24 hrs, 48 hrs, and 72 hrs</li> <li>Observe MAC plate at 24 hours</li> </ul>				
	Single morphology growing on plates:				
3	<ul> <li><u>If organism is a probable pathogen</u>:         <ul> <li>Perform and report full identification</li> <li>Perform and report susceptibility testing as per ASTM</li> </ul> </li> <li><u>If organism is a potential pathogen</u>:         <ul> <li>Perform and report full identification</li> <li>Perform and report susceptibility testing as per ASTM if ANY of the following are true:                 <ul></ul></li></ul></li></ul>				
	Multiple morphologies growing on plates:				
4	<ul> <li>If organism is a probable pathogen:         <ul> <li>Perform and report full identification</li> <li>Perform and report susceptibility testing as per ASTM</li> </ul> </li> <li>If organism is a potential pathogen:         <ul> <li>Perform and report full identification if BOTH are true:                 <ul> <li>Growth is heavy and growth is predominant</li> </ul> </li> <li>Perform and report susceptibility testing as per ASTM if ANY of the following are true:                     <ul> <li>Growth is heavy and growth is predominant</li> <li>Perform and report susceptibility testing as per ASTM if ANY of the following are true:</li></ul></li></ul></li></ul>				
5	not reported individually and not reported as mixed Perform a flood oxidase test on both chocolate and TM agar on day 3. Sub any oxidase positive organisms to chocolate agar immediately.				

### **REPORTING INSTRUCTIONS FOR VAGINAL AND VULVA CULTURE:**

IF	REPORT
No growth after 3 days	<ul> <li>Report: "No growth after 3 days"</li> </ul>
Growth of probable pathogen, NOT Neisseria gonorrhoeae	<ul> <li>Report organism full identification</li> <li>List quantitation</li> <li>Report susceptibility results as per ASTM</li> </ul>
Neisseria gonorrhoeae isolated	<ul> <li>Add organism: "Neisseria gonorrhoeae"</li> <li>List quantification as: "Isolated"</li> <li>Report susceptibility results as per ASTM</li> </ul>
NOTE: N.gonorrhoeae is a critical result if isolated on a child ≤13	<ul> <li>Add isolate comment &amp;REF6</li> <li>In order entry, copy report to OCPHO (HPU1)</li> <li>Refer isolate to APL for susceptibility testing</li> <li>Freeze isolate and log into stored isolates log</li> </ul>
Growth of potential pathogen where full identification is required	<ul> <li>Report organism full identification</li> <li>List quantitation</li> <li>If indicated by procedure, perform and report susceptibility testing as per ASTM</li> </ul>
Growth of potential pathogen where minimal identification is required	<ul> <li>Report "Commensal flora"</li> <li>List quantitation</li> </ul>
Growth of commensal flora	<ul> <li>Report: "Commensal flora"</li> <li>List quantitation</li> </ul>
Mix of enteric Gram-negative bacilli	<ul> <li>Report: "Mixture of coliform organisms"</li> <li>List quantitation</li> </ul>

#### NOTE:

If clinical history states query STI, sexual assault or requests *Neisseria gonorrhoeae* screen and *Neisseria gonorrhoeae* is NOT isolated:

### > Report "No Neisseria gonorrhoeae isolated"

- Add culture comment {GENP
- If growth of yeast is present, add culture comment {GCY

2. Labia Culture and Penis Culture:			
Probable Pathogens <sup>^</sup>			
<ul> <li>Brucella spp.</li> <li>Chromobacterium spp.</li> <li>Eikenella corrodens</li> <li>Pasteurella multocida</li> <li>Pseudomonas aeruginosa</li> <li>Salmonella spp.</li> <li>Shigella spp.</li> <li>Sphingobacterium spp.</li> <li>Vibrio spp.</li> <li>Yersinia spp.</li> <li>GNB Apaerobic</li> <li>Haemainfluer</li> <li>Kingel</li> <li>Morax</li> <li>Neisse</li> <li>GPC Aero</li> <li>β-hem Strept</li> <li>Strept</li> <li>Strept</li> <li>anging</li> </ul>	Sella tularensis*+ ophilus nzaeGPB AeroDiC: • Bacillus anthracis*+ • Bacillus cereus • Bacillus cereus • Bacillus cereus • Listeria spp. • Nocardia spp. • Nocardia spp. 		
Potential Pathogens <sup>^</sup>	Commensal Flora		
<ul> <li>Anaerobes not listed above</li> <li>Enteric GNB not listed above</li> <li>Non-enteric GNB not listed above</li> <li>Enterococcus spp.</li> <li>Staphylococcus intermedius</li> <li>Staphylococcus lugdunensis</li> <li>Yeasts not listed above</li> </ul>	<ul> <li>Bacillus spp. not listed above</li> <li>Coagulase-negative Staphylococci</li> <li>Corynebacterium spp.</li> <li>Micrococcus spp.</li> <li>Non-pathogenic Neisseria spp.</li> <li>viridans Streptococcus grp.</li> </ul>		

\* Risk group 3 organism. If suspected, refer to MIC40100-Suspect High Risk Organism Workup

+ All work-up should be performed in the BSC

<sup>^</sup> For organisms not listed, consult the Microbiology Technical Supervisor or refer to the *Manual of Clinical Microbiology* 

### **INTERPRETATION OF RESULTS:**

Step	Action		
1	<ul> <li>Ensure growth on culture media correlates with gram stain results. If discordant results are found between the gram stain and growth:</li> <li>Re-examine smear and culture plates</li> <li>Check for anaerobic growth</li> <li>Re-incubate media to resolve</li> <li>Consider re-smearing or re-planting specimen</li> </ul>		
2	• Observe BA, CHO, CNA and TM plates at 24 hrs, 48 hrs, and 72 hrs		
	Observe MAC plate at 24 hours		

	Single morphology growing on plates:
	If organism is a probable pathogen:
	Perform and report full identification
	Perform and report susceptibility testing as per ASTM
	If organism is a potential pathogen or commensal flora:
	Perform and report full identification
	Perform and report susceptibility testing if ANY of the following are
	true:
	$\circ$ 3 to 4+WBC in the gram stain
4	<ul> <li>Clinical diagnosis is infection</li> </ul>
-	<ul> <li>Patient is immunocompromised</li> </ul>
	<ul> <li>Multiple cultures are positive for the same organism</li> </ul>
	<ul> <li>If organism is an anaerobe:</li> </ul>
	<ul> <li>Perform and report full identification</li> </ul>
	Perform and refer to APL for susceptibility testing if ANY of the following and thread
	following are true:
	<ul> <li>Organism is a probable pathogen</li> </ul>
	<ul> <li>Organism is predominant in direct smear</li> </ul>
	• Multiple or previous cultures are positive for the same organism
	Multiple morphologies growing on plates:
	If organism is a probable pathogen:
	Perform and report full identification
	Perform and report susceptibility testing as per ASTM
	<u>If organism is a potential pathogen</u> :
	Perform minimal identification and list if ANY of the following are
	true:
	<ul> <li>Moderate to numerous epithelial cells in the gram stain</li> </ul>
	$\circ$ No WBC in the gram stain
	$\circ$ No clinical history that indicates infection was provided
	$\circ \geq$ 3 organisms growing, excluding probable pathogens
	<b>NOTE:</b> Mixed enteric Gram-negative rods should be reported as mixture
	of coliform organisms, not reported individually
5	<b>NOTE:</b> Mixed anaerobes should be reported as mixture of anaerobic
	organisms, not reported individually
	If none of the above are true:
	<ul> <li>Perform and report full identification</li> </ul>
	<ul> <li>Perform and report susceptibility testing as per ASTM if ANY of</li> </ul>
	the following are true:
	♦3 to 4+WBC in the gram stain
	Clinical diagnosis is infection
	Patient is immunocompromised
	Multiple cultures are positive for the same organism
	If organism is commensal flora:
	<ul> <li>Perform minimal identification and report as commensal flora</li> </ul>
	<b>NOTE:</b> Mixed commensal flora should be reported as commensal flora,
	not reported individually and not reported as mixed
	Perform a flood oxidase test on both chocolate and TM agar on day 3.
6	Sub any oxidase positive organisms to chocolate agar immediately.
	sub any onlase positive organisms to chocolate agai infinediately.

## REPORTING INSTRUCTIONS FOR LABIA AND PENIS CULTURE:

IF	REPORT
No growth after 3 days	<ul> <li>Report: "No Growth after 3 days"</li> </ul>
Growth of probable pathogen, NOT Neisseria gonorrhoeae	<ul> <li>Report organism full identification</li> <li>List quantitation</li> <li>Report susceptibility results as per ASTM</li> </ul>
Neisseria gonorrhoeae isolated NOTE: N.gonorrhoeae is a critical result if isolated on a child ≤13	<ul> <li>Add organism: "Neisseria gonorrhoeae"</li> <li>List quantification as: "Isolated"</li> <li>Report susceptibility results as per ASTM</li> <li>Add isolate comment &amp;REF6</li> <li>In order entry, copy report to OCPHO (HPU1)</li> <li>Refer isolate to APL for susceptibility testing</li> </ul>
Growth of	<ul> <li>Freeze isolate and log into stored isolates log</li> <li>Report organism full identification</li> </ul>
potential pathogen or commensal flora where full identification is required	<ul> <li>List quantitation</li> <li>If indicated by procedure, perform and report susceptibility testing as per ASTM</li> </ul>
Growth of potential pathogens where minimal identification and listing is required	<ul> <li>Report the minimal identification         <ul> <li>(i.e., Gram Negative Bacilli - Lactose</li> <li>Fermenter)</li> </ul> </li> <li>List quantitation</li> </ul>
Growth of commensal flora where minimal identification and listing is required	<ul> <li>Report: "Commensal flora"</li> <li>List quantitation</li> </ul>
Mix of enteric Gram-negative bacilli	<ul> <li>Report: "Mixture of coliform organisms"</li> <li>List quantitation</li> </ul>

#### NOTE:

If clinical history states query STI, sexual assault or requests *Neisseria gonorrhoeae* screen and *Neisseria gonorrhoeae* is NOT isolated:

#### > Report "No Neisseria gonorrhoeae isolated"

- Add culture comment {GENP
- > If growth of yeast is present, add culture comment **{GCY**

#### NOTE:

- Refer to Reportable Diseases Act-Public Health Act as of September 2009 for reporting to OCPHO (HPU1)
- Refer to LQM70620-Laboratory Critical Results List-Microbiology for results that need to be phoned to ordering location
- Refer to MIC36100-Nosocomial Infection Notification Job Aid to determine if organism needs to be copied to Infection Prevention and Control
- Refer to MIC36200-Referral of Category A Specimens to APL for sending category A isolates to APL
- Refer to MIC36300-Referral of Category B Specimens to APL for sending category B isolates to APL

# LIMITATIONS:

- 1. A negative genital specimen culture does not eliminate the possibility of a genital tract infection. Organisms such as viruses, Mycoplasmas and Chlamydia are not detected by routine culture.
- 2. Inadequate specimen collection, improper specimen handling and low organism levels in the specimen may yield a false negative result.
- 3. The presence of yeast may inhibit the growth of *Neisseria gonorrhoeae*. Although Thayer Martin agar contains Nystatin to inhibit the growth of yeast, inhibition of *Neisseria gonorrhoeae* should be considered on CHO if culture is positive for yeast species.
- 4. In prepubescent females, diptheroids and coagulase-negative staphylococci are predominant.
- 5. In the adult female, lactobacilli are predominant.
- 6. In postmenopausal women, fewer lactobacilli are present and a greater number of Enterobacteriaceae are predominant.

# **CROSS-REFERENCES:**

- MIC20115-Gram Stain Procedure
- MIC36100-Nosocomial Infection Notification Job Aid
- MIC36200-Referral of Category A Specimens to APL
- MIC36300-Referral of Category B Specimens to APL
- MIC40100-Suspect High Risk Organism Workup
- LQM70620-Laboratory Critical Results List-Microbiology

## **REFERENCES:**

- 1. Leber, A. (2016). *Clinical microbiology procedures handbook.* (4<sup>th</sup>ed.) Washington, D.C.: ASM Press
- Jorgensen J.H., Pfaller M.A., Carroll K.C., Funke G., Landry M.L., Richter S.S., Warnock D.W. (2015). *Manual of Clinical Microbiology*, 11<sup>th</sup> edition. Washington, D.C: ASM Press

## **APPROVAL:**

Date

#### **REVISION HISTORY:**

REVISION	DATE	Description of Change	REQUESTED BY
1.0	27 Nov 17	Initial Release	L. Steven
2.0	22 Feb 21	Procedure reviewed and added to NTHSSA policy template	L. Steven
3.0	27 Feb 23	Procedure reviewed	L. Steven

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