Title: MIC34200-CSF Culture

Issuing Authority: Director of Health Services

Next Review Date:

Type: Laboratory Services Program SOP

Policy Number: Date Approved:

PROGRAM Standard Operating Procedure – Laboratory Services		
Title: MIC34200 – CSF Culture	Policy Number:	
Program Name: Laboratory Services		
Applicable Domain: Lab, DI and Pharmacy Services		
Additional Domain(s):		
Effective Date:	Next Review Date:	
Issuing Authority:	Date Approved:	
Director of Health Services		
Accreditation Canada Applicable Standard:		

GUIDING PRINCIPLE:

Bacterial meningitis is the result of infection of the meninges (lining around the brain). Specimens include central nervous system shunt fluid, external ventricular drainage fluid and cerebro-spinal fluid (CSF). The examination of CSF from patients suspected of having meningitis is always considered a STAT procedure.

PURPOSE/RATIONALE:

To determine the presence or absence of bacterial pathogens in CSF specimens.

SCOPE/APPLICABILITY:

This procedure applies to Medical Laboratory Technologists processing specimens for CSF culture.

SAMPLE INFORMATION:

Special	Refer to Policy 17-02-V1: Specimens Containing Suspected		
Precautions	Risk Group 3 Pathogens		
Туре	CSF collected into clean, sterile, leak-proof tube		
	Central nervous system shunt fluid		
Source	Fluid from Ommaya reservoirs		
Source	External ventricular drainage fluid		
	CSF from lumbar puncture		
	Generally, 1 to 3 mL of CSF is required for the bacterial		
Volume	culture		
Volume	If viral, fungal, or mycobacterial testing is required, then		
	at least 3 to 4 mL should be sent for referral		
Stability	Transport to the laboratory immediately		
Storage	If a delay in processing is anticipated, hold specimens at		
Requirements	room temperature, do NOT refrigerate		

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	physician to prioritize requests
Cuit auia fau	2. Leaking specimens should be processed, but alert the
Criteria for	physician of the possibility of contamination
rejection	3. Improperly collected, labeled, transported, or handled
	specimens should be processed. SCM40110-Waiver of
	Responsibility form needs to be filled out by the
	responsible nurse

REAGENTS and/or MEDIA:

- Blood agar (BA), Chocolate agar (CHO), MacConkey agar (MAC) and Thioglycollate broth (THIO)
- Identification reagents: catalase, oxidase, Staph latex test, Strep latex test, etc.

SUPPLIES:

- Sterile red top vacutainer tube
- Disposable inoculation needles
- Alcohol pads
- Ringed cytology microscope slides
- Wooden sticks

EQUIPMENT

- Biosafety cabinet
- 35° ambient air and 35° CO₂ incubators
- Vitek 2 and supplies

SPECIAL SAFETY PRECAUTIONS:

Containment Level 2 facilities, equipment, and operational practices for work involving infectious or potential 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

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PROCEDURE INSTRUCTIONS:

Step		Action		
Proce	Processing specimens for CSF culture			
1	>1 mL received	 Centrifuge specimen at 3500 rpm for 10 minutes Transfer supernatant to labeled red top tube 		
	<1 mL received	Do not centrifuge		
2	 Place 1 to 2 drops Streak for isolated Prepare smear by prepare smear by prepare	et, using a sterile pipette: of sediment or mixed CSF onto BA, CHO, and MAC growth using a disposable inoculation needle olacing 1 to 2 drops of CSF onto a clean ringed oe slide and allow the drop(s) to form one large drop re also planted to THIO broth and held for 14 days		
3		in the CO ₂ incubator lection tube, supernatant tube, MAC and THIO (if O ₂ incubator		
4	•	d perform gram stain. Gram stain must be read Refer to MIC20115-Gram Stain Procedure.		
5	hours of 08:00 to 20: <1 hour. Outside the	mmediately. During the regular Microbiology lab 00, turnaround time for these gram stains is regular Microbiology lab hours, Microbiology called in if ordering physician determines the stain ately.		
6		sitive CSF gram stain results to ordering location		

Common bacterial causes of acute meningitis by age group			
Neonate	 E.coli S.agalactiae	L.monocytogenes	
Infants/Children	S.agalactiaeH.influenzae	S.pneumoniaeN.meningitides*+	
Adolescents and Young Adults	S.pneumoniaeN.meningitidis*+		
Older Adults	S.pneumoniaeN.meningitides*+H.influenzae	S.agalactiaeL.monocytogenes	
Ventriculoperitoneal Shunt Infections	Coagulase-negative StaphylococcusS.aureusC.albicans	Corynebacterium spp.Propionibacterium acnes	

^{*} Risk group 3 organism. If suspected, refer to 17-02-V1: Specimens Containing Suspected Risk Group 3 Pathogens

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⁺ All work-up should be performed in the BSC

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INTERPRETATION OF RESULTS:

Step	Action
1	Ensure growth on culture media correlates with gram stain results. If discordant results are found between the gram stain and growth: Re-examine smear and culture plates Check for anaerobic growth Re-incubate media to resolve Consider re-smearing or re-planting specimen
2	 Observe BA and CHO plates at 24 hours, 48 hours, and 72 hours Observe MAC plate at 24 hours and 48 hours Observe THIO on day 2, 5, 10 and 14 if applicable
3	If growth is observed, perform biochemical testing to report preliminary ID of the isolate.
4	Provide genus and species identification as soon as possible. If a preliminary identification cannot be made after 24 hours, release a preliminary culture report using the gram stain morphology.

REPORTING RESULTS:

REPORTING RESULTS.			
IF	REPORT		
No growth after 1 day	PRELIM: • Report: "No Growth after 1 Day. Further report to follow"		
No growth after 3 days	FINAL: • Report: "No growth aerobically after 3 days"		
Shunt fluid: No growth after 3 days	INTERIM:Report: "No growth aerobically after 3 days"		
Shunt fluid: No growth after 14 days	FINAL: • Report: "No growth anaerobically after 14 days"		
Growth of organism	 Report organism identification List quantitation as "Isolated" Report susceptibility results as per ASTM Freeze isolate(s) and log into stored isolates log 		
H. influenzae or N.meningitidis isolated	 Must be sent immediately to Alberta Precision Laboratories for typing Refer to MIC36600-Microbiology Organism Referral Freeze isolate(s) and log into stored isolates log 		
S.pyogenes, S.agalactiae, S.pneumoniae, H. influenzae or N.meningitidis isolated	 Any S.pyogenes, S.agalactiae, S.pneumoniae, H.influenzae or N.meningitidis isolated from blood culture specimens must be sent to NML for International Circumpolar Surveillance (ICS) program Refer to MIC36600-Microbiology Organism Referral Freeze isolate(s) and log into stored isolates log 		

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NOTE:

 Refer to Reportable Diseases – Public Health Act as of September 2009 for reporting to 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 isolates to APL
- Refer to MIC36400-Referral of Category B Specimens to DL for sending isolates to DynaLIFE
- Refer to MIC36500-Referral of Category B Specimens to NML for sending isolates to NML
- Refer to MIC36600-Microbiology Organism Referral

LIMITATIONS:

- 1. A positive culture generally indicates infection with the organism.
- 2. Lack of pus cells in CSF does not rule out infection, especially in Listeriosis.
- 3. The most common cause of community acquired bacterial meningitis is Streptococcus pneumoniae.
- 4. Direct bacterial antigen testing is not recommended.
- 5. Since THIO is mainly a broth for anaerobes and does not support the growth of the most common pathogens in CSF, it is not recommended for routine CSF culture but should be used when Shunt Infection is suspected.

CROSS-REFERENCES:

- 17-02-V1: Specimens Containing Suspected Risk Group 3 Pathogens
- LQM70620-Laboratory Critical Results List-Microbiology
- MIC20115-Gram Stain Procedure
- MIC35000-Reportable Diseases Notification
- MIC36100-Nosocomial Infection Notification Job Aid
- MIC36200-Referral of Category A Specimens to APL
- MIC36300-Referral of Category B Specimens to APL
- MIC36400-Referral of Category B Specimens to DynaLIFE
- MIC36500-Referral of Category B Specimens to NML
- Refer to MIC36600-Microbiology Organism Referral
- SCM40110-Waiver of Responsibility

REFERENCES:

- 1. Clinical Microbiology Procedures Handbook, 4th edition, ASM Press, 2016
- 2. 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, 11th edition, ASM Press, Washington, D.C.
- 3. Policy B-0160: Specimens Containing Suspected Risk Group 3 Pathogens for Primary Specimen Handling Flow Chart

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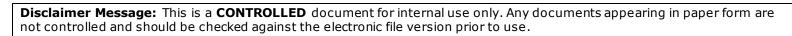
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APPROVAL			
Date		 	

REVISION HISTORY:

REVISION	DATE	Description of Change	REQUESTED BY
1.0	11 Jan 17	Initial Release	L. Steven
2.0	04 Dec 18	Updated to include new Vitek 2 instrument	L. Steven
3.0	30 Jan 21	Procedure reviewed and added to NTHSSA policy template	L. Steven
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