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| Section: Genital Tract Culture Manual | Subject Title: Vaginal Culture | |
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VAGINAL CULTURE

I. Introduction

Vaginal infections are occasionally caused by *Staphylococcus aureus* and beta-hemolytic streptococci (not *S. anginosus (milleri)* group), and in children, *Salmonella* and *Shigella*. Vaginal culture can be used for diagnosis. *Neisseria gonorrhoeae* (GC) and *Chlamydia trachomatis* (CT) will also cause vaginal infections but vaginal swabs are not the optimal specimen to detect these agents.

Toxic-shock syndrome may be associated with vaginitis or vaginal colonization due to *S. aureus* and beta-hemolytic streptococci (not *S. anginosus (milleri)* group). Vaginal culture may be helpful; positive cultures should be tested to determine if they are toxin-producing strains.

II. Specimen Collection and Transport

See Pre-Analytical – Specimen Collection QPCMI2001 [Vaginal Swab for Culture](#)

III. Reagents and Media

See [Analytical Process - Bacteriology Reagents/Materials/Media List QPCMI10001](#)

IV. Procedure

A. Processing of Specimens:

See Specimen Processing Procedure QPCMI06003 [Vaginal Swab for Culture](#)

- a) Direct Examination: not required
- b) Culture:

| Media | Incubation |
|--|-----------------------------------|
| Colistin Nalidixic Acid Agar (CNA) | CO ₂ , 35°C x 48 hours |
| Todd Hewitt Broth for Group B Strep (TH) | O ₂ , 35°C x 24 hours |
| Martin-Lewis Agar (ML) (if requested) | CO ₂ , 35°C x 72 hours |
| MacConkey (MAC) (for <12 years old) | CO ₂ , 35°C x 48 hours |

PROCEDURE MANUAL

TORONTO MEDICAL LABORATORIES / MOUNT SINAI HOSPITAL MICROBIOLOGY DEPARTMENT

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B. Interpretation of culture:

- a) Examine the CNA plate after 24 hours incubation for colonies suspicious of *S. aureus*, beta-hemolytic streptococci (not *S. anginosus (milleri)* group) (Refer to [Bacteria Workup Manual](#) for identification). Send *S. aureus* isolates to PHL for toxin testing and freeze all toxin-producing strain.
- b) If the original CNA plate has no suspicious colonies re-incubate and examine the next day.
- c) After 24 hours incubation, if the original CNA plate is negative for beta-hemolytic streptococci (not *S. anginosus (milleri)* group), subculture a drop of GBS broth onto CNA and incubate in O₂ at 35°C x 24 hours.
- d) Examine ML plate at 24 and 72 hours. For GC work-up, refer to [Bacteria Workup Manual](#).
- e) Examine MAC at 24 and 48 hours. Work-up oxidase-negative non-lactose-fermenters as per [Bacteria Workup Manual](#).

C. Susceptibility testing

Refer to Susceptibility Testing Manual.

V. Reporting Results

Culture:

Negative Report:

If toxic shock syndrome requested:

“No *Staphylococcus aureus* or beta-hemolytic streptococci isolated.”

If ML is set up:

“No *Neisseria gonorrhoeae* isolated”.

If vaginal swab is received for GC culture on adults, report with comment: “The recommended specimen for *Neisseria gonorrhoeae* culture is an endocervical swab.”

If MAC is set up:

Report “No Salmonella or Shigella isolated.”

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Positive Report:

If toxic shock syndrome requested:
Report all significant isolates with appropriate susceptibilities (do not quantitate).

If ML is set up:
“*Neisseria gonorrhoeae* isolated” (do not quantitate)

If MAC is set up:
Report all significant isolates with appropriate susceptibilities (do not quantitate).

Telephone all positive GC cultures to floor/ordering Physician. Refer to [Isolate Notification and Freezing Table QPCMI15003](#)

For all positive GC cultures, send a Communicable Disease Report to the Medical Officer of Health. Refer to [Communicable Disease Results Reporting Process QPCMI16000](#) and [Reportable Diseases to the Medical Officer of Health QPCMI16001](#).

VI. References

Schreckenberger, Paul. Clinical Microbiology Newsletter, 1992 p. 126.

Spiegel, C., Amsel, R., Holmes, K. Journal of Clinical Microbiology, July, 1983 p. 170-177.

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