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# Bacterial Identification and Susceptibility – Anaerobic Information Sheet

# **Overview**

#### **MDL Test Name**

Bacterial Identification and Susceptibility - Anaerobic

#### **MDL Test Code**

BAC IDSUSN

#### **Ask at Order Questions**

Source

# **Specimen Source**

Pure culture isolate on suitable agar (solid) medium; slant or plate in an anaerobic environment.

# **Specimen Requirements**

#### Container/Tube

Suitable agar (solid) medium for isolate submitted. Submit a sealed plate in an anaerobic transport bag to maintain viability.

#### Specimen Volume (minimum)

Pure culture with viable growth must be submitted.

# Sample Stability Time

Stability varies with organism and transport system; usually up to 48 hours at ambient temperatures but fastidious organisms should be sent promptly.

#### **Transport/Storage Conditions**

Ambient  $(20 - 25^{\circ}C)$ 

#### **Patient Preparation / Collection Instructions**

N/A

# **Performance**

# Days Performed

Daily; Monday – Sunday

# Report Available (TAT) - (Once received at MDL)

3 – 5 days

## **Specimen Retention Time**

7 days

# **Method Description**

 Identification methods may include any of the following: conventional biochemical testing, matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry, and commercial identification panels.
 Genus and species are reported on anaerobic bacterial isolates whenever possible.

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- Susceptibility testing will be minimal inhibitory concentration (MIC) (gradient strip diffusion).
- NOTE: Susceptibility testing is performed depending on source and identification. If the source and identification do not warrant susceptibility testing per CLSI guidelines, MDL will notify the client prior to continuing with testing.

#### Reference Values

- Identification of organism.
- Susceptibility results are reported as minimal inhibitory concentration (MIC) in µg/mL. Breakpoints are used to categorize an organism as susceptible, susceptible-dose dependent, intermediate, resistant, or nonsusceptible according to either U.S. Food and Drug Administration (FDA) and/or Clinical Laboratory Standards Institute (CLSI).

#### **Cautions**

- Testing cannot be performed on mixed cultures, the submitted isolate must be pure.
- Testing ability depends on the viability of the organism submitted.
- Verify labeled appropriately and securely sealed for transport to avoid breaking or leakage. Recommend securing the lid with parafilm or similar material. Place in an individually sealed bag and then in an anaerobic transport bag.
- Invitro susceptibility does not guarantee a clinical response. Therefore, the
  decision to treat with a particular agent should not be based solely on the
  antimicrobial susceptibility testing result.