MAYO CLINIC LABORATORIES

Test Definition: FBMO

MVista Blastomyces Quantitative Antigen, Fluid

Reporting Title: MVista Blastomyces Ag, Fluid
Performing Location: MiraVista Diagnostics

Specimen Requirements:

Submit only one of the following:
Specimen Type: CSF or Bronchoalveolar Fluid
Container/Tube: Sterile leak-proof container
Specimen Volume: 2 mL
Collection Instructions:
CSF: Collect 2 mL of spinal fluid (CSF) in sterile leak-proof container. Send refrigerated in a plastic screw cap vial.
Bronchoalveolar Lavage: Collect 2 mL in sterile leak-proof container. Send refrigerated in a plastic screw cap vial.
NOTE:
1. Specimen type is required.
2. Separate order required for each specimen.
3. Sputolysin, sodium hydroxide, and potassium hydroxide treatment degrade the analyte detected in the assay.

Specimen Minimum Volume:

CSF: 0.8 mL; BAL: 0.5 mL

Specimen Type	Temperature	Time	Special Container
Varies	Refrigerated (preferred)	14 days	
	Ambient	14 days	
	Frozen		

Ask at Order Entry (AOE) Questions:

Test ID	Question ID	Description	Type	Reportable
FBMO	Z5523	Specimen Type	Plain Text	Yes

Result Codes:

Result ID	Reporting Name	Type	Unit	LOINC®
Z5523	Specimen Type	Alphanumeric		31208-2
Z5524	Result:	Alphanumeric	ng/mL	Not Provided
Z5525	Interpretation	Alphanumeric		Not Provided

LOINC and CPT codes are provided by the performing laboratory.

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Supplemental Report:
No

CPT Code Information:

87449

Reference Values:

Reference Interval: None Detected
Reportable Range: 0.31 ng/mL - 20.00 ng/mL
Results above 20.00 ng/mL are reported as 'Positive, Above the Limit of Quantification'