Test Definition: FHSAG

MVista Histoplasma Ag, CSF

Overview

Method Name

EnzymeImmunoassay(EIA)

NY State Available

Yes

Specimen

Specimen Type

CSF

Specimen Required

Specimen Type: Spinal Fluid

Sources: CSF

Container/Tube: Sterile container

Specimen Volume: 2 mL

Collection Instructions:Â Collect 2 mL of spinal fluid (CSF). Ship refrigerated. 2 mL of spinal fluid. Send specimen in a plastic, screw-capped vial refrigerated.

Specimen Minimum Volume

0.8 mL

Reject Due To

Hemolysis	NA
Lipemia	NA
Icterus	NA
Other	Specimen that is too viscous to pipette. Tissue, biopsy, sputum, bronchial brush, tracheal aspirate, FNA, bone marrow aspirate, stool or samples in transport media, fixative or Isolator tubes

Specimen Stability Information

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

Clinical and Interpretive

Reference Values

Reference interval: None Detected

Results reported as ng/mL in 0.4 Ã¢â‚¬â€œ 19 ng/mL range

Results above the limit of detection by below 0.4 ng/mL are reported as 'Positive, Below the Limit of Quantification'.

Results above 19.0 ng/mL are reported as 'Positive, Above the Limit of Quantification'.

Cautions

Interfering Substances & Cross-Reactivities:

Sodium hydroxide and sputolysin. Cross-reactivity occurs between blastomycosis and histoplasmosis and in paracoccidioidomycosis, penicillosis, coccidioidomycosis, aspergillosis and porotrichosis.

Performance

PDF Report

No

Day(s) and Time(s) Test Performed

Monday through Friday

Analytic Time

2 days

Maximum Laboratory Time

4 - 8 days

Performing Laboratory Location

MiraVista Diagnostics

Fees and Codes

Fees

- Authorized users can sign in to Test Prices for detailed fee information.
- Clients without access to Test Prices can contact Customer Service 24 hours a day, seven days a week.
- Prospective clients should contact their Regional Manager. For assistance, contact Customer Service.

Test Classification

This test was developed and its performance characteristics determined by MiraVista Diagnostics. It has not been cleared or approved by the FDA; however, FDA clearance or approval is not currently required for clinical use. The results are intended to be used as the sole means for clinical diagnosis or patient management decisions.

CPT Code Information

87385

LOINC® Information

Test ID	Test Order Name	Order LOINC Value
FHSAG	MVista Histoplasma Ag, CSF	51754-0

Result ID	Test Result Name	Result LOINC Value
Z1722	Result:	51754-0
Z1034	Interpretation	Not Provided