Test Definition: FHIST

MVista Histoplasma Ag, S

Overview

Method Name

EnzymeImmunoassay(EIA)

NY State Available

Yes

Specimen

Specimen Type

Serum

Specimen Required

Specimen Type: Serum

Container/Tube: Red or SST

Specimen Volume: 2 mL

Collection Instructions: Draw blood in a plain red-top tube(s). (Serum gel tube is acceptable.) Spin down and send 2 mL serum refrigerate in a plastic vial

Specimen Minimum Volume

1.2 mL

Reject Due To

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

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Serum	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 19.0 ng/mL range

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

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

Cautions

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

Performance

PDF Report

No

Day(s) and Time(s) Test Performed

Monday through Friday

Analytic Time

1 day

Maximum Laboratory Time

3 - 6 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 not intended to be used as the sole means for clinical diagnosis or patient management decisions.

CPT Code Information

87385

LOINC® Information

Test Definition: FHIST

MVista Histoplasma Ag, S

Test ID	Test Order Name	Order LOINC Value
FHIST	MVista Histoplasma Ag, S	51753-2

Result ID	Test Result Name	Result LOINC Value
Z1711	Result:	51753-2
Z1035	Interpretation	Not Provided