MAYO CLINIC LABORATORIES

Test Definition: FHIST

MVista Histoplasma Ag, S

Overview

Method Name
Enzyme Immunoassay (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

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 Minimum Volume
1.2 mL

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container

Serum	Refrigerated (preferred)	14 days	
	Ambient	14 days	
	Frozen		

Clinical & Interpretive

Reference Values

Reference interval: None Detected

Reportable Range: Positive Results reported in ng/mL from 0.20 ng/mL to 20.00 ng/mL.

Positive Results above 20.00 ng/mL are reported as “Above the Limit of Quantification”.

Cautions

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

Performance

PDF Report

No

Performing Laboratory Location

MiraVista Diagnostics

Fees & Codes

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

Result ID	Reporting Name	LOINC®
Z1711	Result:	51753-2
Z1035	Interpretation	Not Provided