

**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 – 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, penicilliosis, 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 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