MAYO CLINIC LABORATORIES

Test Definition: FHST

MVista Histoplasma Ag Quantitative EIA

Reporting Title: MVista Histoplasma Antigen
Performing Location: MiraVista Diagnostics

Specimen Requirements:

Submit only 1 of the following specimens:

Bronchial Washing
Collect 2 mL of Bronchial Washing in leak proofed container.
Ship refrigerate.

Required:

1. Label specimen appropriately (Bronchial Washing)

Body Fluid
Collect 2 mL of Body Fluid in leak proofed container.
Ship refrigerate.

Required:

1. Label specimen appropriately (Type of Body Fluid)

Note: MiraVista will test most body fluids with the following disclaimer: The reference range and other method performance specifications have not been established for this test in this type of Body Fluid. The test results should be integrated into the clinical context for interpretation.

Note: Minimum volume does not allow for repeats.

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

Ask at Order Entry (AOE) Questions:

Test ID	Question ID	Description	Type	Reportable
FHST	Z1746	Specimen Type	Plain Text	Yes

Result Codes:

Result ID	Reporting Name	Type	Unit	LOINC®
Z1746	Specimen Type	Alphanumeric		31208-2
Z1747	Result	Alphanumeric	ng/mL	57766-8
Z1748	Interpretation	Alphanumeric		59464-8

LOINC® and CPT codes are provided by the performing laboratory.

Supplemental Report:
No

CPT Code Information:
87385

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”.