

Laboratory Director: Deborah E. Blue, MD, MT(ASCP)

Patient Name REPORT , H		DOB	Report Printed 4/21/2020
Specimen Type Serum		Client Information Test RightFax 5 Upstairs Left	MiraVista ID M00001033
Specimen ID			
Collection Date 01/01/2001	Received Date 03/18/2020	Fax 317-455-2189	

LAB SERVICES REPORT

Test	Result	Unit	Interpretation	Report Date
310 MVista® Histoplasma Ag Quantitative EIA				
Histoplasma Antigen EIA	None detected	ng/mL	NEGATIVE	03/18/2020

Test Parameters:

Reference Interval: None Detected

Reportable Range: 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 Quantification'

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

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.

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