

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

Patient Name REPORT , HH		DOB 01/01/2001	Report Printed 4/21/2020
Specimen Type Serum		Client Information Test RightFax 5 Upstairs Left	MiraVista ID M00000981
Specimen ID 123456			
Collection Date 01/01/2001	Received Date 03/17/2020	Fax 317-455-2189	

LAB SERVICES REPORT

Test	Result	Unit	Interpretation	Report Date
326 MVista® Histoplasma Antibody IgG IgM EIA				
anti-Histoplasma IgG EIA	Negative	EU	NEGATIVE	03/18/2020
anti-Histoplasma IgM EIA	Negative	EU	NEGATIVE	03/18/2020

Test Parameters:

Reference Interval: Negative
 Reportable Range: Results below 8.0 are Negative
 Results between 8.0 and 9.9 are Intermediate
 Results 10.0 and above are Positive

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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