

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

Patient Name REPORT , HH		DOB	Report Printed 4/21/2020
Specimen Type Serum		Client Information Test RightFax 5 Upstairs Left	MiraVista ID M00000984
Specimen ID			
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	30.8	EU	POSITIVE	03/18/2020
anti-Histoplasma IgM EIA	23.9	EU	POSITIVE	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.

This document contains confidential privileged information. The recipient of the information is prohibited from disclosing the contents to another party without authorization. If you are not the intended recipient, you are hereby notified that disclosure of the contents is strictly prohibited. Please notify MiraVista Diagnostics immediately if you received this information in error.

Patient Name REPORT , HH	Specimen ID	Specimen Type Serum	Collection Date 01/01/2001
------------------------------------	--------------------	-------------------------------	--------------------------------------