MiraVista DIAGNOSTICS
Rapid Fungal Testing. Accurate Results.

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

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.