MiraVista DIAGNOSTICS
Rapid Fungal Testing. Accurate Results.

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Patient Name		DOB	Report Printed
REPORT , H			4/21/2020
Specimen Type		**Client Information**	**MiraVista ID**
Serum		Test RightFax 5 Upstairs Left	M00001034
Specimen ID			
Collection Date	**Received Date**	**Fax** 317-455-2189	
01/01/2001	03/18/2020		

LAB SERVICES REPORT

Test	Result	Unit	Interpretation	Report Date
310 MVista® Histoplasma Ag Quantitative EIA				
Histoplasma Antigen EIA	13.88	ng/mL	**POSITIVE**	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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