MiraVista DIAGNOSTICS
Rapid Fungal Testing. Accurate Results.

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Patient Name		DOB	Report Printed
Example Patient		08/26/1961	11/30/2020
Specimen Type		**Client Information**	**MiraVista ID**
Urine		Example Laboratory	
Specimen ID			
20X-319MV003			
Collection Date	**Received Date**	**Fax**	
11/14/2020	11/16/2020		

LAB SERVICES REPORT

Test	Result	Unit	Interpretation	Report Date
310 MVista® Histoplasma Ag Quantitative EIA				
Histoplasma Antigen by EIA	None detected	ng/mL	NEGATIVE	11/16/2020

Test Parameters:

Reference Interval: None Detected
Reportable Range: Positive Results reported in ng/mL from 0.20 ng/mL to 20.00 ng/mL
Positive results above 20.00 ng/mL are reported as "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.