

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

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

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.

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