

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

<b>Patient Name</b> REPORT , B		<b>DOB</b>	<b>Report Printed</b> 4/21/2020
<b>Specimen Type</b> Serum		<b>Client Information</b> Test RightFax 5 Upstairs Left	<b>MiraVista ID</b> M00001040
<b>Specimen ID</b>			
<b>Collection Date</b> 01/01/2001	<b>Received Date</b> 03/18/2020	<b>Fax</b> 317-455-2189	

**LAB SERVICES REPORT**

Test	Result	Unit	Interpretation	Report Date
<b>316 MVista® Blastomyces Ag Quantitative EIA</b>				
Blastomyces Antigen EIA	1.40	ng/mL	<b>POSITIVE</b>	03/18/2020

**Test Parameters:**

Reference Interval: None Detected

Reportable Range: Results reported as ng/mL in 0.2 - 14.7 ng/mL range

Results above the limit of detection but below 0.2 ng/mL are reported as 'Positive, Below the Limit of Quantification'

Results above 14.7 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.

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.

<b>Patient Name</b> REPORT , B	<b>Specimen ID</b>	<b>Specimen Type</b> Serum	<b>Collection Date</b> 01/01/2001
-----------------------------------	--------------------	-------------------------------	--------------------------------------