ARUP LABORATORIES | aruplab.com
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Tracy I. George, MD, Chief Medical Officer

Patient Report

Patient Age/Sex: 7 years Female

Specimen Collected: 07-Mar-22 14:46

Histoplasma Galactomannan Ag Quant, Urn **|Received: 07-Mar-22 14:49** **Report/Verified: 07-Mar-22 15:16**

Procedure	Result	Units	Reference Interval
Histoplasma Galactomannan Ag Quant,Urn	24.0	ng/mL	
Histoplasma Galactomannan Ag Interp,Urn	**Detected** * f1 i1		Not Detected

Result Footnote

f1: Histoplasma Galactomannan Ag Interp, Urn

Cross-reactivity with other endemic mycoses (Blastomyces and Coccidioides) may occur. Positive test results should be correlated with other clinical findings and relevant exposure history.

Test Information

i1: Histoplasma Galactomannan Ag Interp, Urn

INTERPRETIVE DATA: Histoplasma Galactomannan Antigen
Quantitative by EIA, Urine
Less than 0.4 ng/ml = Not Detected
0.4-0.7 ng/mL = Detected (below the limit of quantification)
0.8-24.0 ng/mL = Detected
Greater than 24.0 ng/mL = Detected (above the limit of quantification)

The quantitative range of this assay is 0.8-24.0 ng/mL. Antigen concentrations between 0.4-.07 or >24.0 ng/mL fall outside the linear range of the assay and cannot be accurately quantified.

This EIA test should be used in conjunction with other diagnostic procedures, including microbiological culture, histological examination of biopsy samples, and/or radiographic evidence, to aid in the diagnosis of histoplasmosis.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

Unless otherwise indicated, testing performed at:
ARUP Laboratories
500 Chipeta Way, Salt Lake City, UT 84108
Laboratory Director: Tracy I. George, MD

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