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**Specimen Collected:** 8/4/2025 15:00 MDT

Histoplasma Galactomannan Ag Quant, Urn		Received: 8/6/2025 06:43 MDT		Report/Verified: 8/6/2025 12:42 MDT	
Procedure		Result		Units	
Histoplasma Galactomannan Ag Quant,Urn		2.164		Reference Interval	
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 0.4-0.7 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.

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\*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

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**Unless otherwise indicated, testing performed at:****ARUP Laboratories**

500 Chipeta Way, Salt Lake City, UT 84108

Laboratory Director: Jonathan R. Genzen, MD, PhD

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