HOTLINE: Effective May 16, 2022

2009418 ***Histoplasma*** **Galactomannan Antigen Quantitative by EIA, Urine** **HISTOGM U**

Specimen Required: Collect: Random urine.
Specimen Preparation: Transfer 2 mL urine to an ARUP Standard Transport Tube.
Storage/Transport Temperature: Refrigerated.
Unacceptable Conditions: Specimens other than urine. Urine in boric acid. Serum; refer to test *Histoplasma* Antigen by EIA, Serum (ARUP test code 0092522).
Stability (collection to initiation of testing): Ambient: 24 hours; Refrigerated: 2 weeks; Frozen: 2 weeks

Interpretive Data:
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 US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.