ARUP Laboratories
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*****Example Report*****

Patient Age/Gender: 19 years Male
Printed: 19-Jun-19 13:07:00

Procedure	Result		Units	Ref Interval	Accession	Collected	Received	Reported/ Verified
Histoplasma Antigen, Serum Interp	**Detected**	*		[Not Detected]	19-170-900119	19-Jun-19 12:21:00	19-Jun-19 12:21:00	19-Jun-19 12:24:40
Histoplasma Antigen, Serum	3.50		ng/mL		19-170-900119	19-Jun-19 12:21:00	19-Jun-19 12:21:00	19-Jun-19 12:24:40

19-Jun-19 12:21:00 Histoplasma Antigen, Serum Interp:
INTERPRETIVE INFORMATION: Histoplasma Antigen Quantitative by EIA, Serum

Less than 0.19 ng/mL = Not Detected
0.19-60.0 ng/mL = Detected
Greater than 60.0 ng/mL = Detected (above the limit of quantification).

The quantitative range of this assay is 0.19-60.0 ng/mL. Antigen concentrations greater than 60.0 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.

Crossreactivity with Blastomyces dermatiditis, Coccidioides immitis and possibly Talaromyces marneffei have been observed with this EIA. Other clinically and geographically relevant endemic mycoses should be considered in the case of a positive test result.

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement B: aruplab.com/CS