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

Patient Age/Gender: 44 years Male
Printed: 23-Sep-19 08:07:20

Procedure	Result	Units	Ref Interval	Accession	Collected	Received	Reported/ Verified
Histoplasma Antigen, Serum Interp	**Detected ***		[Not Detected]	19-263-900176	20-Sep-19 17:21:00	20-Sep-19 17:21:00	20-Sep-19 17:24:45
Histoplasma Antigen, Serum	**Not Quantified *f**		[Not Detected]	19-263-900176	20-Sep-19 17:21:00	20-Sep-19 17:21:00	20-Sep-19 17:24:45

20-Sep-19 17:21:00 Histoplasma Antigen, Serum:
Histoplasma Antigen was detected, but at a level below 0.19 ng/mL. Antigen detected at a level below 0.19 ng/mL cannot be accurately quantified by this assay.

20-Sep-19 17:21:00 Histoplasma Antigen, Serum Interp:
INTERPRETIVE INFORMATION: Histoplasma Antigen Quantitative by EIA, Serum

The quantitative range of this assay is 0.19-60.0 ng/mL. Antigen concentrations less than 0.19 ng/mL or 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, serology and/or radiographic evidence, to aid in the diagnosis of histoplasmosis.

Cross-reactivity 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: www.aruplab.com/CS