

NEW TEST – Available Now

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HBV Prenatal Triple Panel

3018943, HBV TRI PN

Specimen Requirements:

Patient Preparation:

Collect: Serum separator tube (SST).

Specimen Preparation: Separate serum from cells ASAP or within 2 hours of collection.

Transfer 3.00 mL serum to an ARUP standard transport tube.

Effective Date: April 21, 2025

(Min: 1.750 mL)

Transport Temperature: Refrigerated.

Unacceptable Conditions: Heparinized plasma. Specimens containing particulate material

or obvious microbial contamination. Heat-inactivated or

severely hemolyzed.

Remarks:

Stability: After separation from cells: Ambient: 12 hours; Refrigerated: 7

days; Frozen: 30 days (avoid repeated freeze/thaw cycles).

Methodology: Quantitative Chemiluminescent Immunoassay (CLIA) /

Qualitative Chemiluminescent Immunoassay (CLIA)

Performed: Sun-Sat

Reported: 1-2 days

Note: The HBV core total assay tests for IgG and IgM antibodies, but

does not differentiate between them. HBsAb results greater than 1,000.00 IU/L are reported as greater than 1,000.00 IU/L. If results for HBsAg screen are reactive (=1.0), then HBsAg Confirmation Prepatal will be added. Additional charges apply

Confirmation, Prenatal will be added. Additional charges apply.

CPT Codes: 86706; 86704; 87340; if reflexed, add 87341

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Reference Interval:



Effective Date: April 21, 2025

Test Number	Components	Reference Interval	
	Hepatitis B Surface Antigen, Prenatal	Negative	
	Hepatitis B Surface Antigen, Prenatal		
	Hepatitis B Surface Antigen, Prenatal		
	Hepatitis B Core Antibodies, Total	Negative	
	Hepatitis B Surface Antibody		
		Less than 10.00 IU/L	Negative
		Greater than or equal to 10.00 IU/L	Positive

HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.