Lab Dept:	Serology
Test Name:	HEPATITIS Be ANTIGEN (HBeAg)
General Information	
Lab Order Codes:	HBEA
Synonyms:	Hepatitis HBe Ag; HBeAg; Hepatitis Be Viral Antigen
CPT Codes:	87350 – Hepatitis Be antigen
Test Includes:	Hepatitis Be Antigen reported as positive, negative or equivocal.
Logistics	
Test Indications:	Determining the presence or absence of detectable hepatitis B virus e antigen in monitoring infection status of individuals with chronic hepatitis B
	Determining infectivity of hepatitis B virus (HBV) carriers
	Monitoring serologic response of chronically HBV-infected patients receiving antiviral therapy
Lab Testing Sections:	Serology - Sendouts
Referred to:	Mayo Clinic Laboratories (MML Test: EAG)
Phone Numbers:	MIN Lab: 612-813-6280
	STP Lab: 651-220-6550
Test Availability:	Daily, 24 hours
Turnaround Time:	1 - 3 days; test performed Monday - Saturday
Special Instructions:	N/A
Specimen	
Specimen Type:	Blood
Container:	SST (Gold or Marble) tube
Draw Volume:	2.2 mL (Minimum: 1.8 mL) blood

Processed Volume:	0.7 mL (Minimum: 0.6 mL) serum
Collection:	Routine venipuncture
Special Processing:	Lab Staff: Centrifuge specimen, remove serum aliquot into a screw- capped round bottom plastic vial within 2 hours of collection. Store and ship at frozen temperatures. Forward promptly.
	Specimen stable frozen (preferred) for 90 days, refrigerated for 6 days, ambient for 72 hours.
Patient Preparation:	For 24 hours before specimen collection, patient should not take multivitamins or dietary supplements (e.g., hair, skin, and nail supplements) containing biotin (vitamin B7).
Sample Rejection:	Specimens other than serum; warm specimens; gross lipemia; grossly icteric; gross hemolysis; mislabeled or unlabeled specimens
Interpretive	
Reference Range:	Negative
Critical Values:	N/A
Limitations:	Disappearance of hepatitis B virus e antigen or appearance of HBe antibody in serum does not completely rule-out chronic hepatitis B carrier state or infectivity.
	Specimens should not be taken from patients receiving therapy with high biotin doses (ie, >5 mg/day) until at least 8 hours following the last biotin administration.
	Performance characteristics of this assay have not been established in patients younger than 2 years or in populations of immunocompromised or immunosuppressed patients. This assay is not licensed by US Food and Drug Administration for testing cord blood specimens or screening donors of blood, plasma, human cell, or tissue products.
	Performance characteristics have not been established for the following specimen characteristics: -Grossly icteric (total bilirubin level >40 mg/dL) -Grossly lipemic (intralipid level >2200 mg/dL) -Grossly hemolyzed (hemoglobin level >2200 mg/dL) -Specimen containing particulate matter
Methodology:	Electrochemiluminescence Immunoassay (ECLIA)
References:	Mayo Clinic Laboratories April 2024

Updates:

4/6/2004: Test moved from Memorial Blood Center of Minneapolis to Mayo Medical Laboratories. 1/16/2017: Update to SST.

4/23/2024: Updated optimal and minimum specimen volumes, changed methodology, updated limitations, added specimen stability, added patient preparation.