Hepatitis B Virus (HBV) Quantitative Real-time PCR  
Test Code: 1100

Clinical and Procedure  
Clinical Utility  
Hepatitis B quantitative DNA PCR can be used in conjunction with clinical presentation and other laboratory markers of disease status as an aid in managing individuals infected with HBV. Results from the assay can potentially be used to assess disease progression and to monitor the efficacy of antiviral therapy by measuring changes in HBV DNA levels during the course of therapy. Viral load tests should not be used to diagnose HBV infection.

Procedure  
Extraction of DNA from specimen; amplification and detection of hepatitis B genotypes A through H using real-time, quantitative PCR. An internal control is added to ensure the extraction was performed correctly and the PCR reaction was not inhibited. RealTime HBV is a product of Abbott Laboratories. It is FDA approved for in vitro diagnostic use.

Specificity  
Detects all 8 HBV genotypes. The primers and probes used in this assay are specific for HBV.

Turnaround Time  
Same day (within 8 - 12 hours from receipt of specimen), Tuesday through Saturday.

Specimen Information  
<table>
<thead>
<tr>
<th>Specimen Type</th>
<th>Order Code</th>
<th>CPT Code</th>
<th>NY Approved</th>
<th>Volume</th>
<th>Assay Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>plasma</td>
<td>1101</td>
<td>87517</td>
<td>Yes</td>
<td>2 mL (min. 0.7 mL)</td>
<td>10 IU/mL to 1.0 x 10^9 IU/mL. HBV DNA detected below 10 IU/mL will be reported as &quot;&lt;10 IU/mL&quot;. Reported in 2 formats: IU/mL and Log10 IU/mL.</td>
</tr>
<tr>
<td>serum</td>
<td>1110</td>
<td>87517</td>
<td>Yes</td>
<td>2 mL (min. 0.7 mL)</td>
<td>10 IU/mL to 1.0 x 10^9 IU/mL. HBV DNA detected below 10 IU/mL will be reported as &quot;&lt;10 IU/mL&quot;. Reported in 2 formats: IU/mL and Log10 IU/mL.</td>
</tr>
</tbody>
</table>

Special Instructions  
- Collect 4-5 mL whole blood in EDTA.  
- Centrifuge within 6 hours of draw and transfer 2 mL plasma to a sterile, screw top tube.  
- If shipped ambient, separated plasma fraction must arrive within 24 hours of draw.

Shipping
Ship Monday through Friday. Friday shipments must be labeled for Saturday delivery. All specimens must be labeled with patient's name and collection date. A Viracor Eurofins test requisition form must accompany each specimen. Multiple tests can be run on one specimen. Ship specimens FedEx Priority Overnight® to: Viracor Eurofins, 1001 NW Technology Dr, Lee's Summit, MO 64086

**Causes for Rejection**
Specimens beyond their acceptable length of time from collection as listed in the specimen handling or specimen types other than those listed.

**Disclaimer**
Specimens are approved for testing in New York only when indicated in the Specimen Information field above.

The CPT codes provided are based on Viracor Eurofins' interpretation of the American Medical Association's Current Procedural Terminology (CPT) codes and are provided for general informational purposes only. CPT coding is the sole responsibility of the billing party. Questions regarding coding should be addressed to your local Medicare carrier. Viracor Eurofins assumes no responsibility for billing errors due to reliance on the CPT codes illustrated in this material.