

# Foresight®

## HBsAb Quantitative EIA Test Kit Package Insert

REF	I231-1271	English
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An enzyme immunoassay (EIA) for the quantitative detection of Hepatitis B Surface Antibody (HBsAb) including IgG, IgM and IgA antibodies in human serum or plasma.

For professional *in vitro* diagnostic use only.

### INTENDED USE

The HBsAb Quantitative EIA Test Kit is a one step enzyme immunoassay for the quantitative detection of Hepatitis B Surface Antibody (HBsAb) including IgG, IgM and IgA antibodies in human serum or plasma. It is intended as a screening tool for previous exposure to the Hepatitis B virus (HBV), follow-up of infected individuals, monitoring after vaccination or susceptibility to infection and determination of need for vaccination.

### SUMMARY

Hepatitis B virus is a spherical enveloped, partially double-stranded DNA virus of the Hepadnaviridae family.<sup>1</sup> The Hepatitis B infection of the liver is transmitted through sexual contact, blood borne exposure, transmission from mother to child during delivery, sharing of objects that pierce the skin, child-to-child and household contact.<sup>2,3,4,5</sup> HBV infection has been linked to a variety of mild to chronic liver diseases, including cirrhosis, and hepatocellular carcinoma.<sup>4</sup> In some cases, the virus may persist for a lifetime. Annually, 1 million people die from chronic active hepatitis, cirrhosis or primary liver cancer. Hepatitis B affects millions of people worldwide and is considered a global public health problem.<sup>4,5</sup>

HBsAg is one of the earliest markers that appear in the blood following infection with the Hepatitis B virus. The body's immune response to infection includes the development of specific antibodies to HBsAg. These antibodies appear a few weeks after HBsAg is cleared from the blood and the virus can no longer be passed on to others. The appearance of HBsAb is associated with recovery and is regarded as the marker for immunity. These antibodies also appear as a result of successful immunization.<sup>3,5</sup> Therefore, the detection and monitoring of antibodies to HBsAg has become an important tool in the screening and monitoring of infected individuals as well as an indicator of those successfully vaccinated against HBV.

The HBsAb Quantitative EIA Test Kit is an immunoassay for the quantitative detection of the presence of Hepatitis B Surface Antibody (HBsAb) including IgG, IgM and IgA antibodies in serum or plasma specimen.

### PRINCIPLE

The HBsAb Quantitative EIA Test Kit is a solid phase quantitative enzyme immunoassay based on the sandwich principle for the detection of HBsAb including IgG, IgM and IgA antibodies in human serum or plasma. The microwell plate is coated with recombinant HBsAg. During testing, the specimens are added to the antigen coated microwell plate and then incubated. If the specimen contains HBsAb, it will bind to the antigen coated on the microwell plate and simultaneously bind to the conjugate to form immobilized antigen-HBsAb-conjugate complexes. If the specimen does not contain HBsAb, the complexes will not be formed. After initial incubation, the microwell plate is washed to remove unbound materials. Substrate A and substrate B are added and then incubated to produce a blue color, indicating the amount of HBsAb present in the specimen. Sulfuric acid solution is added to the microwell plate to stop the reaction which produces a color change from blue to yellow. The color intensity, which corresponds to the amount of HBsAb present in the specimen, is measured with a microplate reader at 450/630-700 nm or 450 nm.

### PRECAUTIONS

- For professional *in vitro* diagnostic use only. Do not use after expiration date.
- Do not mix reagents from other kits with different lot numbers.
- Avoid cross contamination between reagents to ensure valid test results.
- Follow the wash procedure to ensure optimum assay performance.
- Use Plate Sealer to cover microwell plate during incubation to minimize evaporation.
- Use a new pipet tip for each specimen assayed.
- Ensure that the bottom of the plate is clean and dry and that no bubbles are present on the surface of the liquid before reading the plate. Do not allow wells to dry out during the assay procedure.
- Do not touch the bottom of the wells with pipette tips. Do not touch the bottom of the microwell plate with fingertips.
- Do not allow sodium hypochlorite fumes from chlorine bleach or other sources to contact the microwell plate during the assay as the color reaction may be inhibited.
- All equipment should be used with care, calibrated regularly and maintained following the equipment manufacturer's instructions.

### HEALTH AND SAFETY INFORMATION

- Some components of this kit contain human blood derivatives. No known test method can offer complete assurance that products derived from human blood will not transmit infectious agents. Therefore, all blood derivatives should be considered potentially infectious. It is recommended that these reagents and human specimens be handled using established good laboratory working practices.
- Wear disposable gloves and other protective clothing such as laboratory coats and eye protection

while handling kit reagents and specimens. Wash hands thoroughly when finished.

- ProClin™ 300 is included as a preservative in the Conjugate, Concentrated Wash Buffer, Substrate and Calibrators. Avoid any contact with skin or eyes.
- Do not eat, drink or smoke in the area where the specimens or kits are handled. Do not pipette by mouth.
- Avoid any contact of the Substrate A, Substrate B, and Stop Solution with skin or mucosa. The Stop Solution contains 0.5M sulfuric acid which is a strong acid. If spills occur, wipe immediately with large amounts of water. If the acid contacts the skin or eyes, flush with large amounts of water and seek medical attention.
- Non-disposable apparatus should be sterilized after use. The preferred method is to autoclave for one hour at 121°C. Disposables should be autoclaved or incinerated. Do not autoclave materials containing sodium hypochlorite.
- Handle and dispose all specimens and materials used to perform the test as if they contained infectious agents. Observe established precautions against microbiological hazards throughout all the procedures and follow the standard procedures for proper disposal of specimens.
- Observe Good Laboratory Practices when handling chemicals and potentially infectious material. Discard all contaminated material, specimens and reagents of human origin after proper decontamination and by following local, state and federal regulations.
- Neutralized acids and other liquids should be decontaminated by adding sufficient volume of sodium hypochlorite to obtain a final concentration of at least 1.0%. A 30 minute exposure to a 1.0% sodium hypochlorite may be necessary to ensure effective decontamination.

### STORAGE AND STABILITY

- Unopened test kits should be stored at 2-8°C upon receipt. All unopened reagents are stable through the expiration date printed on the box if stored between 2-8°C. Once opened, all reagents are stable for up to 3 months after the first opening date if stored between 2-8°C. Return reagents to 2-8°C immediately after use.
- Allow the sealed pouch to reach room temperature before opening the pouch and remove the required number of strips to prevent condensation of the microwell plate. The remaining unused strips should be stored in the original resealable pouch with desiccant supplied at 2-8°C and can be used within 3 months of the opening date. Return the remaining unused strips and supplied desiccant to the original resealable pouch, firmly press the seal closure to seal the pouch completely and immediately store at 2-8°C.
- Concentrated Wash Buffer may be stored at room temperature to avoid crystallization. If crystals are present, warm up the solution at 37°C. Working Wash Buffer is stable for 2 weeks at room temperature.
- Do not expose reagents especially the Substrate to strong light or hypochlorite fumes during storage or incubation steps.
- Do not store Stop Solution in a shallow dish or return it to the original bottle after use.

### SPECIMEN COLLECTION AND PREPARATION

- The HBsAb Quantitative EIA Test Kit can be performed using only human serum or plasma collected from venipuncture whole blood.
- EDTA, sodium heparin, and ACD collection tubes may be used to collect venipuncture whole blood and plasma specimens. The preservative sodium azide inactivates horseradish peroxidase and may lead to erroneous results.
- Separate serum or plasma from blood as soon as possible to avoid hemolysis. Grossly hemolytic, lipidic or turbid samples should not be used. Specimen with extensive particulate should be clarified by centrifugation prior to use. Do not use specimens with fibrin particles or contaminated with microbial growth.
- Serum and plasma specimens may be stored at 2-8°C for up to 7 days prior to assaying. For long term storage, specimens should be kept frozen below -20°C.
- Bring specimens to room temperature prior to testing. Frozen specimens must be completely thawed and mixed well prior to testing. Specimens should not be frozen and thawed repeatedly.
- If specimens are to be shipped, they should be packed in compliance with local regulations covering the transportation of etiologic agents.

### REAGENTS AND COMPONENTS

#### Materials Provided

No.	Reagent	Component Description	Quantity	
			96 wells/kit	480 wells/kit
	HBsAb Microwell Plate	Microwell plate coated with recombinant HBsAg	1 plate (96 wells/plate)	5 plates (96 wells/plate)
1	HBsAb Conjugate	Purified HBsAg bound to peroxidase; Preservative: 0.1% ProClin™ 300	1 x 8 mL	5 x 8 mL
2	Concentrated Wash Buffer (25x)	Tris-HCl buffer containing 0.1% Tween 20; Preservative: 0.1% ProClin™ 300	1 x 40 mL	5 x 40 mL
3	Substrate A	Citrate-phosphate buffer containing hydrogen peroxide; Preservative: 0.1% ProClin™ 300	1 x 8 mL	5 x 8 mL

4	Substrate B	Buffer containing tetramethylbenzidine (TMB); Preservative: 0.1% ProClin™ 300	1 x 8 mL	5 x 8 mL
5	Stop Solution	0.5M Sulfuric acid	1 x 8 mL	5 x 8 mL
6	HBsAb Calibrator 1	Diluted human serum non-reactive for HBsAb, HCV, HIV-1, and HIV-2; Preservative: 0.1% ProClin™ 300	1 x 12 mL	5 x 12 mL
7	HBsAb Calibrator 2	Diluted human serum containing 10 mIU/ml HBsAb and negative for HCV, HIV-1, and HIV-2; Preservative: 0.1% ProClin™ 300	1 x 1 mL	5 x 1 mL
8	HBsAb Calibrator 3	Diluted human serum containing 50 mIU/ml HBsAb and negative for HCV, HIV-1, and HIV-2; Preservative: 0.1% ProClin™ 300	1 x 1 mL	5 x 1 mL
9	HBsAb Calibrator 4	Diluted human serum containing 100 mIU/ml HBsAb and negative for HCV, HIV-1, and HIV-2; Preservative: 0.1% ProClin™ 300	1 x 1 mL	5 x 1 mL
10	HBsAb Calibrator 5	Diluted human serum containing 200 mIU/ml HBsAb and negative for HCV, HIV-1, and HIV-2; Preservative: 0.1% ProClin™ 300	1 x 1 mL	5 x 1 mL
	Plate Sealers		2	10
	Package Insert		1	1

#### Materials Required But Not Provided

- Freshly distilled or deionized water
- Sodium hypochlorite solution for decontamination
- Absorbent paper or paper towel
- Water bath or incubator capable of maintaining 37°C ± 2°C.
- Calibrated automatic or manual microwell plate washer capable of aspirating and dispensing 350 µL/well
- Disposable gloves
- Automated processor (optional)
- Calibrated micropipettes with disposable tips capable of dispensing 50 µL
- Graduated cylinders for wash buffer dilution
- Vortex mixer for specimen mixing (optional)
- Disposable reagent reservoirs
- Calibrated microplate reader capable of reading at 450 nm with a 630-700 nm reference filter, or reading at 450 nm without a reference filter
- Timer

### DIRECTIONS FOR USE

Allow reagents and specimens to reach room temperature (15-30°C) prior to testing. The procedure must be strictly followed. Assay must proceed to completion within time limits. Arrange the controls so that well A1 is the Blank well. From well A1, arrange the controls in a horizontal or vertical configuration. The procedure below assigns specific wells arranged in a vertical configuration. Configuration may depend upon software.

Step	Detailed Procedure	Simplified Procedure
	<ul style="list-style-type: none"> <li>Prepare Working Wash Buffer by diluting the Concentrated Wash Buffer 1:25. Pour the contents of the bottle in a graduated cylinder and fill it with freshly distilled or deionized water to 1000 mL for 96 wells/plate testing. The Working Wash Buffer is stable for 2 weeks at 15-30°C.</li> <li><b>Note:</b> If crystals are present in the Concentrated Wash Buffer, warm it up at 37°C until all crystals dissolve.</li> <li>Remove unused strips from the microwell plate, and store in the original resealable pouch at 2-8°C.</li> </ul>	<ul style="list-style-type: none"> <li>Prepare Working Wash Buffer by diluting the Concentrated Wash Buffer 1:25</li> <li>Remove and store unused strips at 2-8°C</li> </ul>
0	<ul style="list-style-type: none"> <li>Leave A1 as Blank well.</li> </ul>	<ul style="list-style-type: none"> <li>Leave A1 as Blank well</li> </ul>
1	<ul style="list-style-type: none"> <li>Add 50 µL of Calibrator 1 in wells B1 and C1.</li> <li>Add 50 µL of Calibrator 2 in wells D1 and E1.</li> <li>Add 50 µL of Calibrator 3 in wells F1 and G1.</li> <li>Add 50 µL of Calibrator 4 in wells H1 and A2.</li> <li>Add 50 µL of Calibrator 5 in wells B2 and C2.</li> </ul> <p>The colors of Calibrator 1-5 gradually change from yellow to blue.</p>	<ul style="list-style-type: none"> <li>B1 and C1: Add 50 µL Calibrator 1</li> <li>D1 and E1: Add 50 µL Calibrator 2</li> <li>F1 and G1: Add 50 µL Calibrator 3</li> <li>H1 and A2: Add 50 µL Calibrator 4</li> <li>B2 and C2: Add 50 µL Calibrator 5</li> </ul>
2	<ul style="list-style-type: none"> <li>Add 50 µL of specimen to assigned wells starting at D2.</li> </ul>	<ul style="list-style-type: none"> <li>Starting D2: Add 50 µL specimen</li> </ul>
3	<ul style="list-style-type: none"> <li>Add 50 µL of Conjugate to each well except for the Blank well. (Red Reagent).</li> </ul>	<ul style="list-style-type: none"> <li>Add 50 µL of Conjugate to each well except for the Blank well</li> </ul>
4	<ul style="list-style-type: none"> <li>Mix gently by swirling the microwell plate on a flat bench for 30 seconds.</li> <li>Cover the microwell plate with the Plate Sealer and incubate in a water bath or an incubator at 37°C ± 2°C for 30 minutes ± 2 minutes.</li> </ul>	<ul style="list-style-type: none"> <li>Mix gently</li> <li>Cover the microwell plate with the Plate Sealer and incubate at 37°C for 30 min</li> </ul>

5	<ul style="list-style-type: none"> <li>Remove the Plate Sealer.</li> <li>Wash each well 5 times with 350 µL of Working Wash Buffer per well, then remove the liquid.</li> <li>Turn the microwell plate upside down on absorbent tissue for a few seconds. Ensure that all wells have been completely washed and dried.</li> </ul> <p>Note: Improper washing may cause false positive results.</p>	<ul style="list-style-type: none"> <li>Remove the Plate Sealer</li> <li>Wash each well 5 times with 350 µL of Working Wash Buffer</li> <li>Turn the microwell plate upside down on absorbent tissue</li> </ul>
6	<ul style="list-style-type: none"> <li>Add 50 µL of Substrate A to each well. (Clear Reagent)</li> <li>Add 50 µL of Substrate B to each well. (Clear Reagent)</li> </ul> <p>Then a blue color should develop in wells containing Positive specimens.</p>	<ul style="list-style-type: none"> <li>Add 50 µL of Substrate A to each well</li> <li>Add 50 µL of Substrate B to each well</li> </ul>
7	<ul style="list-style-type: none"> <li>Mix gently then cover microwell plate with Plate Sealer and incubate in a water bath or incubator at 37°C ± 2°C for 15 minutes ± 1 minute.</li> </ul>	<ul style="list-style-type: none"> <li>Mix then cover microwell plate with Plate Sealer and incubate at 37°C for 15 min</li> </ul>
8	<ul style="list-style-type: none"> <li>Remove the Plate Sealer.</li> <li>Add 50 µL of Stop Solution to each well. (Clear Reagent)</li> </ul> <p>Then a yellow color should develop in wells containing Positive specimens.</p>	<ul style="list-style-type: none"> <li>Remove the Plate Sealer</li> <li>Add 50 µL of Stop Solution to each well</li> </ul>
9	<ul style="list-style-type: none"> <li>Read at 450/630-700 nm within 30 minutes.</li> </ul> <p>Note: Microwell plate can also be read at 450 nm, but it is strongly recommended to read it at 450/630-700 nm for better results.</p>	<ul style="list-style-type: none"> <li>Read at 450/630-700 nm within 30 min</li> </ul>

#### AUTOMATED PROCESSING

Automatic EIA microplate processors may be used to perform the assay after validating the results to ensure they are equivalent to those obtained using the manual method for the same specimens. Incubation times may vary depending on the processors used but do not program less incubation times than the procedure listed above. When automatic EIA microplate processors are used, periodic validation is recommended to ensure proper results.

#### VALIDATION REQUIREMENTS AND QUALITY CONTROL

1. Calculate the Mean Absorbance of Calibrators 1-5 by referring to the table below.

##### Example of Calibrator 2 Calculation

Item	Absorbance
Calibrator 2: Well D1	0.165
Calibrator 2: Well E1	0.161
Total Absorbance of Calibrator 2	0.165 + 0.161 = 0.326
Mean Absorbance of Calibrator 2	0.326/2 = 0.163
Blank Absorbance: Well A1	0.005
Mean Absorbance of Calibrator 2 – Blank Absorbance	0.163 - 0.005 = 0.158

2. Check the validation requirements below to determine if the test results are valid.

Item	Validation Requirements
Blank Well	Blank Absorbance should be < 0.050 if read at 450/630-700 nm Note: It should be < 0.100 if read at 450 nm
Calibrator 1	Mean Absorbance after subtraction of Blank Absorbance should be < 0.105
Calibrator 2	Mean Absorbance after subtraction of Blank Absorbance should be > 0.105
Calibrator 4	Mean Absorbance after subtraction of Blank Absorbance should be > 0.500

**NOTE:** The test results are considered invalid if the above validation requirements are not met. Repeat the test or contact your local distributor.

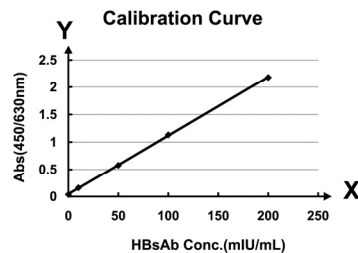
#### INTERPRETATION OF RESULTS

Draw the calibration curve and obtain quantitative specimen results.

1. Subtract the Blank Absorbance from the Mean Absorbance of each Calibrator, then plot them on the Y-axis against their concentration in mIU/mL on the X-axis on a linear graph paper and draw the calibration curve. Draw the best fitted line through data points to obtain a standard curve. Refer to an example of the calibration curve at right.

**NOTE:** Do not use the calibration curve at right to make any calculation. A calibration curve must be performed for each run.

2. Obtain quantitative specimen results from their absorbance after subtraction of Blank Absorbance



by using the calibration curve.

**NOTE:** Specimens that have absorbance above Calibrator 5 should be pre-diluted using Calibrator 1 and retested. The concentration must be multiplied by the dilution factor. Automated reading and calculation may be performed using linear regression function on suitable computer programs.

Results	Concentration
Negative	< 9 mIU/mL
Positive	> 11mIU/mL
Equivocal*	9 - 11mIU/mL

\***NOTE:** For Equivocal results, the specimen should be retested. Specimens that are repeatedly Equivocal after retest should be confirmed using an alternate method. If the results remain Equivocal, collect a new specimen in two weeks. If the new specimen is Positive, the specimen is presumed to be Positive.

#### LIMITATIONS

- The HBsAb Quantitative EIA Test Kit is used for the detection of HBsAb in human serum or plasma. Diagnosis of an infectious disease should not be established based on a single test result. Further testing, including confirmatory testing, should be performed before a specimen is considered positive. A non-reactive test result does not exclude the possibility of exposure. Specimens containing precipitate may give inconsistent test results.
- As with all diagnostic tests, all results must be interpreted together with other clinical information available to the physician.
- As with other sensitive immunoassays, there is the possibility that non-repeatable reactive results may occur due to inadequate washing. The results may be affected due to procedural or instrument error.
- Erroneous result may be due to fibrin particles and microbial contamination.

#### PERFORMANCE CHARACTERISTICS

##### Clinical Sensitivity and Specificity

The HBsAb Quantitative EIA Test Kit has correctly identified specimens of a seroconversion panel and has been compared with a leading commercial HBsAb Quantitative EIA test using clinical specimens. The results show that the clinical sensitivity of the HBsAb Quantitative EIA Test Kit is 99.3%, and the clinical specificity is 99.2%.

##### HBsAb Quantitative EIA vs. Other EIA

Method	Other EIA		Total Results
	Positive	Negative	
HBsAb Quantitative EIA	Positive	1139	1154
	Negative	8	1839
<b>Total Results</b>	1147	1854	3001

Clinical Sensitivity: 99.3% (98.6-99.7%)\*

Clinical Specificity: 99.2% (98.7-99.6%)\*

Overall Agreement: 99.2% (98.9-99.5%)\*

\*95% Confidence Interval

##### Reproducibility

**Intra-Assay:** Within-run precision has been determined by using 20 replicates of one specimen.

**Inter-Assay:** Between-run precision has been determined by 3 independent assays on the same sample. Test 20 replicates of the same with three different lots of the HBsAb Quantitative EIA Test Kit.

Specimen	Intra-Assay			Inter-Assay		
	Mean Absorbance	Standard Deviation	Coefficient of Variation (%)	Mean Absorbance	Standard Deviation	Coefficient of Variation (%)
1	1.081	0.063	5.828	1.074	0.071	6.611

#### BIBLIOGRAPHY

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- Centers for Disease Control. Viral Hepatitis B Fact Sheet.
- Richman, D., R. Whitley, F. Hayden. *Clinical Virology*. New York: Churchill Livingstone Inc., 1997.
- World Health Organization. World Health Organization Hepatitis B Fact Sheet. N°204. Revised October 2000.
- World Health Organization. Hepatitis B. 2002.

#### Index of Symbols

	Consult instructions for use		Tests per kit		Manufacturer
	For <i>in vitro</i> diagnostic use only		Use by		Catalog #
	Store between 2-8°C		Lot Number		Substrate B
	HBsAb		Substrate A		Calibrator 1
	Wash Buffer (25x)		Conjugate		Calibrator 3
	Calibrator 1		Calibrator 2		Stop Solution
	Calibrator 4		Calibrator 5		Package Insert
	Microwell Plate		Plate Sealer		



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