



HBsAg Rapid Card Test Kit

A Rapid One Step Visual Test for the qualitative detection of Hepatitis B Surface antigen in human serum or plasma

For Professional Use

CE₁₀₂₃

IVD

 READ THIS PACK INSERT CAREFULLY BEFORE PERFORMING THE TEST

CATALOGUE No. : HBC-LF

Intended Use : HEPA-SCAN[®] HBsAg card is an immunochromatographic based assay for the qualitative detection of Hepatitis B surface antigen in human serum or plasma.

INTRODUCTION :

The most common agent of acute hepatitis is a virus. Of all the viruses in hepatitis, Hepatitis B virus is responsible for the most serious form of the diseases. HEPA-SCAN[®] is a rapid, qualitative, one-step immunoassay based on the immunochromatographic principle. This method employs unique combination of monoclonal dye conjugate (colloidal gold) and polyclonal solid phase antibodies to selectively identify Hepatitis B surface antigen with a high degree of sensitivity.

Test Principles :

The HBsAg test device consists of a sample window containing a Sample pad where the serum/plasma is to be added. The sample pad is held in contact with the porous membrane material. The membrane has three zones of antibody. The first is mobilized by the sample while the other two are immobilised. The mobile zone consists of colored colloidal gold particles sensitized with monoclonal antibody. The second zone consists of polyclonal antibody immobilized on the membrane (Test Line). The third zone (Control line) Consists of an anti-mouse immunoglobulin antibody which is also immobilised. If the serum contains HBsAg it will form a complex with anti-HBsAg colloidal gold conjugate and allows this to be trapped by the test line, causing the formation of a red line. The unbound colloidal gold particles continue to move along the strip by capillary action until they come in contact with the control line and are trapped, giving a red line demonstrating the validity of tests.

Storage & Stability :

Store between 2°C-30°C. Do not freeze. The kit is stable until the expiry date mentioned on the Pouch, when stored under the above condition.

Pack Size :

Available in packs of 20's, 50's & 100's

Contents of the kit :

One testing device, One 2ml dropper and silica gel as a dehydrant.

Material required but not provided :

- Sterilised Vial
- Autoclaved tips
- Precision Pipette
- Disposable gloves
- Sodium hypochlorite solution (Free available Chlorine 50-500mg/L).

Warnings & Precautions :

In order to obtain reproducible results, the following rules must be observed,

- Read this pack insert carefully.
- DO NOT FREEZE THE KITS. If refrigerated the kits should be brought to room temperature before testing. Assay should be conducted between 15-30°C.
- Do not use reagents after their expiry date printed on the label.
- Do not use the kits beyond their expiry date.
- Use only serum/plasma specimen. Further more do not use umbilical cord blood, because it prevents colloidal gold from migrating & can interfere with results.
- Carefully observe the prescribed number of drops to be added, 2-3 DROPS ONLY.
- Use the test device soon after it is removed from the pouch.
- Do not use the test device, if the pouch seal is broken.
- Specimen should not be repeatedly frozen & thawed
- Specimen with extremely high concentrations of red blood cells, fibrin should be re-centrifuged before using.
- If both test & control lines fairly weak, dilute the sample 1:10 using PBS and retest with a fresh Hepa-Scan Card. The reason for both line appearing weak is due to the phenomenon called Hook's effect whereby the HBsAg titer in the sample is abnormally high.
- Avoid any contamination among samples; for this purpose, disposable tips should be used for each sample.
- In order to avoid personal and environmental contamination
 - Use disposable gloves while handling potentially infectious material and while performing the assay.
 - Do not pipette reagents by mouth.

- n). Do not smoke, eat drink or apply cosmetics during the assay.
 o). Avoid splashing or forming aerosols.
 p). Discard all materials and specimens as if capable of transmitting infection. The preferred method of disposal is autoclaving for a minimum of one hour at 121°C. Liquid wastes not containing acid may be mixed with sodium hypochlorite in volumes such that the final mixture contains 50-500 mg/L available chlorine. Allow 30 minutes for decontamination to be completed.
 q). Liquid waste containing acid must be neutralized with a proportional amount of base pair prior to the addition of sodium hypochlorite.
 r). Spills should be wiped up thoroughly using either an iodophor disinfectant or sodium hypochlorite solution. Materials used to wipe up spills should be added to biohazardous waste matter for proper disposal.
 s). Optimal results will be obtained by strict adherence to the test protocol. Accurate and precise pipetting, as well as following the exact time and temperature requirements is essential.
 t). Once the assay has been started, all steps should be performed without interruption.

u). Storage and Stability :

Store between 2-30°C. Do not freeze. The kit is stable until the expiry date mentioned on the pouch, when stored under the above condition.

v). For Invitro Diagnostic Use only.

w). For single use only.

x). Avoid using hemolytic, lipaemic, ecteric or bacterially contaminated specimens. Otherwise they may give erroneous results.

Specimen :

Fresh Serum / Plasma can be used for testing.

Specimen Collection and Preparation :

a). Collect blood in a clean, dry, sterilized vial and allow it to clot. Separate the serum by centrifugation at 5000 r.p.m. for 15 minutes at room temperature.

b). If serum is not to be assayed immediately it should be stored at 2-8°C or if storing more than 3 days, then freeze the specimen at -20°C or below.

Assay Procedure :

a). Bring the kit and samples to room temperature.

b). Remove the device from the pouch just prior to testing.

c). Place the device on a flat surface.

d). Add 2-3 Drops of serum/plasma into the sample window.

e). Read the positive in results in 10 minutes & Negative results in 20 minutes. **DO NOT READ ANY RESULT BEYOND TWENTY MINUTES.**

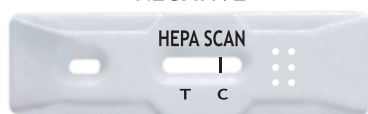
f). Any line appearing after 20 minutes would be of no diagnostic value.

Interpretation of Test Result :

Negative

The presence of only one Band at "C" within the result window indicates a Negative Result.

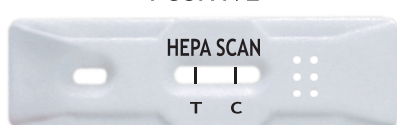
NEGATIVE



Positive

The presence of two Bands at "C" & "T" within the Result window indicates the Positive Result.

POSITIVE



Invalid

If no lines appear or only test line appears within the result window, after performing the test, the result is considered invalid. The direction may not have been followed correctly or the silicagel might have turned white. Repeat the test with a new device.

INVALID (Retest)



Sensitivity : The HEPA-SCAN® Test Device kit is able to detect upto 0.5ng/ml of HBsAg in a sample. 100% Sensitivity.

Specificity : Specificity of HEPA-SCAN® HBsAg Card test device is 99.5%.

Limitation of the test :

a). HEPA-SCAN® HBsAg Card Test detects HBsAg in human serum or plasma and is only a screening test. All reactive samples should be confirmed by supplemental assays like ELISA or RIA. Therefore, for a definitive diagnosis, the patient's clinical history, symptomology as well as serological data should be considered. The results should be reported only after complying with above procedure.

b). The assay is only validated for serum and plasma from individual bleeds and not for poo's of serum or other body fluids.

c). A non-reactive result does not exclude the possibility of exposure to or infection with HBsAg.

d). It should be noted that repeated false reactive results may occur due to non-specific binding of the sample to the membrane.

e). The presence of anti-HBsAg does not imply a Hepatitis B infection but may be indicative of recent and / or past infection by HBsAg.

f). Patients with auto-immune liver diseases may show falsely reactive results.

g). The kit works best when used with fresh samples and when all the kit components are at room temperature (25±5°C). Samples which have been frozen and thawed several times contain particles which can block the membrane, hence resulting in improper flow of reagents and high background colour which may make the interpretation of result difficult.

h). Optimum test performance depends on strict adherence to the test procedure as described in this manual. Any deviation from test procedure may lead to erratic result.

i). Adding more than 2-3 drops of serum/plasma will result in erratic result.

References :

1). NCCLS Document M29-T2 (1991). Protection of laboratory workers from instrument biohazards. Vol. 11, No. 15.

2). Blumberg. B.S., Alter, H.J. and Visnich, S.A. "New" Antigen in Leukemia sera, J.A.M.A. 191:541-546, 1965.

EN 980:2008 (E) MEDICAL DEVICES SYMBOL					
	Temperature Limitation		Date of Manufacture		In vitro Diagnostic Device
	Batch Code		Company name & address		Refer Operating Instructions
	Use by		Company Name		Authorised Representative in European Community
	Do Not Reuse		Sufficient for		KEEP AWAY FROM SUNLIGHT
	KEEP DRY		NON-STERILE		NEGATIVE CONTROL
	POSITIVE CONTROL				



M/S PDB Consulting
 Bruno Rosen, Flehenburg
 48, 42489 Walfrath, Germany
 P : + 49 - 2058 - 8989881
 Fax : + 49 - 2058 - 8989882

Mfg. in India by :

BHAT BIO-TECH INDIA (P) LTD.

No. 11-A, 4th Cross, Veerasandra Indl. Area,
 Electronic City, Bangalore - 560 100
 Phone : 080-4351 4000 (30 Lines)

Fax : 080-4351 4001
 www.bhatbiotech.com

Rev-7, 2010-09-28

