Sensitivity comparison of ELISA and Rapid Screening Techniques for the Detection of HBsAg among Chronic Liver Disease (CLD) Patients in a Tertiary Care Hospital, South Bihar, India

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ABSTRACT

BACKGROUND
Hepatitis B is a major health problem in India. Based on the prevalence of hepatitis B carrier in India is the intermediate endemic level of hepatitis B surface antigen (HBsAg). This study aims to determine the frequency of HBV by screening of chronic liver disease (CLD) patients. This study also aims to compare the sensitivity between two diagnostic tests; one step rapid test strip device and Enzyme Linked Immunosorbent Assay (ELISA).

MATERIALS AND METHODS
A cross-sectional study was carried out in adult patients with liver disease attending the Hepatology OPD, Tertiary Care Hospital in Nalanda (Bihar), India. Age, gender and clinical history of the patients were recorded. Blood specimen was screened for Hepatitis B surface Antigen (HBsAg) using one step rapid test strip device and Enzyme Linked Immunosorbent Assay (ELISA).

RESULTS
Five hundred patients were enrolled in the study. The mean age of infected patients in the study group was 37.7 ± 1.32 years (range 19 to 76 years). Of the 500 samples tested, 18.4% were positive for HBsAg by HEPA-SCAN HBsAg ELISA Test and 13.4% were positive by Hepacard one step rapid test. Considering the results of HEPA-SCAN HBsAg ELISA Test, the sensitivity, specificity, PPV and NPV of ELISA were 97.87%, 99.75%, 98.92% and 99.51% respectively. The sensitivity, specificity, PPV and NPV of Hepacard one step rapid test were 97.10%, 99.76 %, 98.52% and 99.54% respectively.

CONCLUSION
This study shows that frequency of HBV is high in India and the incidence is greater in males than the females. We also noted that in comparing both methods (ELISA method and the rapid test strip) for assessing the presence of HBsAg, ELISA test method was found to be more sensitive than the rapid test strip device. Therefore, we recommend strongly, that ELISA method be used to confirm test results obtained from the one step rapid test, when screening for chronic liver disease (CLD).

KEYWORDS
Chronic Liver Disease, ELISA Methods, Hepatitis B, Rapid Test Strip.

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BACKGROUND
Hepatitis B Virus, which causes serious liver damage is one of the WHO’s target for global eradication by 2020.[1] It is a major public health problem worldwide and is more prevalent in the developing countries.[2,3] More than 2 billion people are infected with HBV worldwide while some 280 million are chronic carriers, harbouring the virus in their liver.[4]

Hepatitis B is a major health problem in India. Based on the prevalence of hepatitis B carrier state in the general population, countries are classified as having high (8% or more), intermediate (2-7%), or low (less than 2%) HBV endemicity. India is at the intermediate endemic level of hepatitis B, with hepatitis B surface antigen (HBsAg) prevalence between 2% and 10% among the populations studied. The prevalence does not vary significantly by region in the country. The number of HBsAg carriers in India has been estimated to be over 40 million (4 crore). Hepatitis B is one of the major diseases of mankind and is a serious global public health problem. Of the 2 billion people who have been infected with the hepatitis B virus (HBV) in the world, more than 360 million have chronic (lifelong) infections. These chronically infected people are at high risk of death from cirrhosis of the liver and liver cancer, diseases that kill about one million each year. In India, of the 25 million infants born every year, over one million run the lifetime risk of developing chronic HBV infection. Estimates indicate that annually over 100,000 Indians die due to illnesses related to HBV infection.[5] Hepatitis B Virus (HBV) infection is a global public health problem. It is estimated that approximately 360 million people are infected worldwide with the virus.[6]

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hepatitis is a systemic disease primarily involving the liver. Most of the cases of acute viral hepatitis are caused by Hepatitis A (HAV), Hepatitis B (HBV) or Hepatitis C Viruses (HCV). HBV has a double standard DNA encoding for P, X, core and surface proteins. The complex antigen found on the surface of HBV is called Hepatitis B surface antigen (HBsAg). Antibodies against HBV proteins are other immunological markers of infection, of which Anti-Hepatitis B core antigen, Hepatitis B envelope antigen and Hepatitis B envelope antibody are also identified shortly after HBsAg, and are important markers of past or present HBV infection. In a typical Hepatitis B infection, Hepatitis B surface antigen (HBsAg) will be detected within 2 to 5 weeks before symptoms or jaundice develop.[7]

Presently, both serological and molecular screening tests are employed for the diagnosis and patient monitoring of HBV infection. Nucleic acid test (NAT) is preferable in terms of its simplicity, rapidness and sensitivity but in some cases of occult HBV especially, NAT may miss some positive samples. NAT has the ability to decrease window period. However, it has been shown that NAT is too costly for testing individual samples. Among all HBsAg assays, ELISA techniques are the most frequently used because of their effectiveness. In many developed countries, HBsAg screening is usually done with ELISA techniques.[8] Blood transfusion services are a vital part of modern health care system, with every unit of blood there is 1% chance of transfusion associated problem including transfusion transmissible diseases. Transfusing infected blood to unsuspected patients in need is a crime. It is mandatory to test each and every unit of donor blood for antibodies to HIV-1 and HIV-2, syphilis, HbsAg and HC Virus. ELISA is the recommended and preferred screening technique for blood banking.[10] Many blood banks still do not have this facility and rather prefer rapid screening kits because they are easy to perform, cheap and are user friendly kits, and do not require sophisticated equipment and elaborate training. A preliminary study was therefore conducted to evaluate the efficacy of these rapid testing kits for screening of blood donors.[11] HBsAg rapid test strip is a rapid screening test for the qualitative detection of HBsAg in whole blood, serum or plasma specimen. The test utilises a combination of monoclonal and polyclonal antibodies to selectively detect elevated levels of HBsAg in whole blood, serum or plasma.[12] Enzyme-linked Immunosorbent Assay (ELISA) is an enzymatic immunoassay technique of the “sandwich” type for the detection of Hepatitis B virus in human serum or plasma. The test uses monoclonal antibodies selected for their ability to bind themselves to the various sub-types of HBsAg now recognised by the World Health Organization (WHO) and the most part of variant HBV strains. [13, 14]

The present study was hence carried out to find out the prevalence of HBsAg in liver disease patients and to compare the routinely Central Pathology Section used HBsAg detection kit using Hepacard one step rapid test with the ELISA for in-vitro qualitative detection of HBsAg test kit in human serum (HEPA-SCAN HBsAg ELISA Test was used).

MATERIALS AND METHODS

Study Design

A cross-sectional study was carried out in adult patients with liver disease attending the Hepatology OPD, Tertiary Care Hospital in Nalanda, (Bihar), India after obtaining institutional ethics committee approval over a period of Two years from January 2015 to January 2017. All the consecutive patients during the study period who were above 18 years of age and gave written informed consent were included in the study. Age, gender and clinical history of the patients were recorded in the case record form after obtaining written informed consent.

Patient’s Inclusion Criteria

This study included a total of five hundred (500) patients enrolled in the study. The mean age of infected patients in the study group was 37.7 ± 1.32 years (range 19 to 75 years), i.e. a cohort study of selected chronic liver disease (CLD) patients visiting the Hepatology OPD, Vardhaman Institute of Medical Sciences, Pawapuri, Nalanda, Bihar and Associated Hospital of Bihar between January 2015 and January 2017. The inclusion criteria for the selection of the CLD patients used for this study were presence of jaundice, ascites, hepatomegaly and oedema.

This prospective study was conducted in the Department of Pathology, Vardhaman Institute of Medical Sciences, Pawapuri and Associated Hospital of Bihar, India from January 2015 to January 2017. The 500 serum/plasma samples were screened using Hepacard one step rapid test for the detection of hepatitis B surface antigen (HBsAg) (Manufactured by Diagnostic Enterprises, Plot no.26, Indl. Estate, Sector -1, Parwanoo-173220, HP). It is a rapid immunochromatographic assay designed for qualitative determination of HBsAg in human serum. It is for in-vitro diagnostic use with sensitivity of 97.10% and specificity of 99.76%.

Detection of HBsAg using ELISA Method

The 500 subjects were also screened using ELISA method for hepatitis B. The ELISA test is a solid-phase microtitre plate coated with monodonal antibodies to human IgM which is based on sandwich principle. ELISA for in-vitro qualitative detection of HBsAg test kit in human serum (HEPA-SCAN HBsAg ELISA Test) was used. It is for in-vitro qualitative detection use with sensitivity of 97.87% and specificity of 99.75%.

Statistical Analysis

The X² (Chi-square) test and analysis using the statistical software (SPSS version 18) was performed for quantitative variables to check for relationship in detecting HBV infection. Percentages were calculated directly for HBV infection. P = 0.05 was used as the accepted significance level.

RESULTS

Five hundred patients were enrolled in the study. The prevalence of HBsAg was found to be 17.8%. The mean age of infected patients in the study group was 37.7 ± 1.32 years (range 19 to 76 years). Three hundred and Forty six (69.22%) were male and One hundred and Fifty four (30.78%) were female. The positivity amongst the male population was 18.20% which was higher than the female population (15.58%), but the difference was statistically significant (p = 0.05) (Table-1).

Most of the CLD patients were in the age group of 19 - 30 years of age 28.26% followed by 35.86% in 31 - 45 years, 30.43% in 46 - 60 years and 5.43% in 61 - 75 (Table 1).
Seroprevalence of HBV among 500 CLD patients in present study was 18.4%. HBV infection was seen mostly in the age group of 31 - 45 yrs. (35.86%) (Chart Table No. 2).

Of the 500 samples tested, 18.4% were positive for HBsAg by HEPA-SCAN HBsAg ELISA Test [Fig-1] and 13.4% were positive by Hepacard one step rapid test [Fig/Table-2] (Chart Table-1). Considering the results of HEPA-SCAN HBsAg ELISA Test, the sensitivity, specificity, PPV and NPV of ELISA were 97.87%, 99.75%, 98.92% and 99.51% respectively. The sensitivity, specificity, PPV and NPV of Hepacard one step rapid test were 97.10%, 99.76 %, 98.52% and 99.54% respectively. (Table-4)

Education level of 34.74% of the study participants were illiterate, 32.60% had primary education, 10.86% had upto secondary education and 27.17% were upto high school. Only 5.43% were graduates. HBV infection was maximum in illiterates 32/92 (34.74%) and patients having education upto primary level 30/92 (32.60%) (Table 3).

<table>
<thead>
<tr>
<th>Age</th>
<th>Number</th>
<th>Positive (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>19-30</td>
<td>76</td>
<td>26 (28.26)</td>
</tr>
<tr>
<td>31-45</td>
<td>216</td>
<td>33 (35.86)</td>
</tr>
<tr>
<td>46-60</td>
<td>178</td>
<td>28 (30.43)</td>
</tr>
<tr>
<td>61-75</td>
<td>30</td>
<td>5 (16.67)</td>
</tr>
<tr>
<td>Male</td>
<td>346</td>
<td>66 (19.07)</td>
</tr>
<tr>
<td>Female</td>
<td>154</td>
<td>26 (16.68)</td>
</tr>
</tbody>
</table>

Table 1. Age and Gender Wise Distribution of Patients

Education Status of HBsAg Patients

<table>
<thead>
<tr>
<th>Educational Status</th>
<th>CLD NO</th>
<th>CLD %</th>
<th>HBV +ve NO</th>
<th>HBV +ve %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Illiterate</td>
<td>125</td>
<td>25</td>
<td>32</td>
<td>34.78</td>
</tr>
<tr>
<td>Primary school</td>
<td>175</td>
<td>35</td>
<td>30</td>
<td>32.60</td>
</tr>
<tr>
<td>Secondary school</td>
<td>100</td>
<td>20</td>
<td>10</td>
<td>10.86</td>
</tr>
<tr>
<td>High school</td>
<td>60</td>
<td>12</td>
<td>15</td>
<td>16.30</td>
</tr>
<tr>
<td>Graduate</td>
<td>40</td>
<td>8</td>
<td>5</td>
<td>5.43</td>
</tr>
<tr>
<td>Total</td>
<td>500</td>
<td>100</td>
<td>92</td>
<td>100</td>
</tr>
</tbody>
</table>

Table 3. Educational Status of HBsAg Patients

| Sensitivity | 97.87% | 99.76% |
| Specificity | 99.75% | 99.76% |
| Positive Predictive Value (PPV) | 98.92% | 98.52% |
| Negative Predictive Value (NPV) | 99.51% | 99.54% |

Table 4. Sensitivity Comparison of ELISA and Rapid Screening Techniques for the Detection of HBsAg among Chronic Liver Disease (CLD) patients when ELISA was taken as Gold Standard
DISCUSSION
It is estimated by the World Health Organisation that there are about 350 million chronic carriers of hepatitis B spreading in every continent in Asia, America, Europe and Africa. In this study, two methods were used (rapid strip test device and ELISA test) to check for sensitivity in screening for hepatitis B surface antigen among chronic liver disease patients. A percentage of the subjects (97.87%) tested positive using ELISA method while 97.10% tested positive using the rapid test strip.

Viral hepatitis is the most common cause of chronic liver disease throughout the world. In India, HBV is reported to be responsible for 70% of chronic hepatitis cases and 80% of cirrhosis of liver cases. The mean age of the infected subjects in the present study was 43 years which is in concordance with a study published by Arora and Mann. Nayak et al in their study have suggested that 30% of chronic carriers get infected vertically and remaining get infection horizontally from those who got it vertically. The chances of horizontal infection occurring through close contact with carriers, use of unsafe injections, and an association with a number of sociocultural practices increases with advancing age. Hepatitis B vaccination became available in 1981 and has been included in the Universal Immunization Program in India as late as 2007-2008. This has contributed significantly in reducing the prevalence thereafter. However, in the present study, the prevalence of Hepatitis B infection gradually decreased with age from 34.9% in 19-30 years age group to 13.4% in patients older than 45 years of age. This possibly could be due to the other causes of chronic liver disease in older age group. In our study, 17.8% of patients with liver disease had hepatitis B infection which is similar to the study by Kumar et al., (17.34%). Other studies in India have reported HBsAg detection rate varying between 12.2-51% in liver disease patients. The difference in prevalence rate in a given region is dependent on the degree of endemicity in that region. The higher prevalence in the present study as compared to that in the general population is expected since the patients enrolled were liver disease patients attending the Hepatology OPD with majority having symptoms like jaundice, anorexia, nausea, vomiting, and haematemesis.

A high frequency rate has been seen in ELISA method when compared to rapid strip test device. So, there is a great and urgent need for people in this locality and beyond to screen their blood for HBV and it should be screened using the ELISA technique which has been found to be more sensitive and accurate as compared to the rapid test strip device method.

CONCLUSION
This study shows that frequency of HBV is high in India and the incidence is greater in males than the females. We also noted that in comparing both methods (ELISA method and the rapid test strip) for assessing the presence of HBsAg, ELISA test method was found to be more sensitive than the rapid test strip device. Therefore, we recommend strongly, that ELISA method be used to confirm test results obtained from the one step rapid test, when screening for chronic liver disease (CLD).

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