Original Research Article SEROLOGICAL MARKERS OF HEPATITIS B VIRUS INFECTIVITY AMONG HEPATITIS B SURFACE ANTIGEN NEGATIVE BLOOD DONORS AT THE UNIVERSITY COLLEGE HOSPITAL, IBADAN.

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8 ABSTRACT

9 Background: Transmission of HBV infection has been documented from hepatitis B surface
10 antigen negative blood donations.

Objective: To determine the prevalence of serological markers of hepatitis B virus infectivity
 among hepatitis B surface antigen negative blood donors.

Materials and Methods: A cross-sectional study was carried out among 490 blood donors who were negative for HBsAg. anti-HBc and other viral markers such as anti-HBs, HBeAg and anti-HBe were tested using ELISA kits by DIAPRO Diagnostic Bioprobes Milano, Italy.

Results: The mean age of participants was 32.5 years (± 9.5), majority were males, 462 (94.3%). Eighty-three (16.9%) were positive for anti-HBc, out of which 35 (7.1%) had anti-HBc alone, 30 (6.1%) had both anti-HBc and anti-HBs while 18 (3.7%) were positive for anti-HBc, anti-HBs and anti-HBe. Antibody to HBsAg (anti-HBs) was detected in 54 (11%) donor samples, of which 6(1.2%) were positive for anti-HBs alone. The number of donors positive for anti-HBeAg was 18 (3.7%). However, no subject was positive for HBeAg.

Conclusion: This study has showed that some blood units containing other markers of HBV are being transfused to recipients even after screening for HBsAg is negative. These blood units are potentially infectious and can cause post-transfusion hepatitis in the recipients. There is need to consider introduction of testing for other markers of HBV infection in our blood banks.

26 Keywords: Blood donors, anti-HBc, HBsAg negative, HBV infectivity

27 INTRODUCTION

The detection of hepatitis B surface antigen (HBsAg) in blood is the mainstay in the diagnosis 28 and screening for HBV infection in most developing countries, including Nigeria.^(1,2) However, it 29 has been reported that transmission of HBV infection by blood transfusion still occurs in a 30 proportion of cases even if the transfused blood tested negative for HBsAg using highly sensitive 31 assays.^(3,4) Therefore, Hepatitis B virus (HBV) remains a major risk of transfusion-transmitted 32 infection. The other modes of HBV transmission are perinatally (mother to child), close 33 interpersonal contact with blood and other body, unsafe injection practices and sexual contact.⁽⁵⁾ 34 Nucleic acid testing (NAT) of all collected units of blood would give near zero risk of 35 transfusion-associated HBV⁽⁶⁾. However, NAT has not been adopted in most developing 36 37 countries, including Nigeria due to cost.

It is estimated that worldwide more than two billion people have been infected by HBV and 257 million have chronic infection. The HBV carrier rate variation is 1-20% worldwide. HBV infection accounts for 500,000 to 1.2 million deaths each year.⁽⁷⁾ Studies have shown that the prevalence of HBV infection is relatively higher in the tropics particularly African region, where it has been reported to be endemic for HBV infection, accounting for the high number of patients chronically infected with HBV. ⁽⁸⁾

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The production of antibodies against HBsAg confers protective immunity and can be detected in patients who have recovered from HBV infection or in those who have been vaccinated. Antibody to HBcAg is detected in almost every patient with previous exposure to HBV. The Immunoglobulin M (IgM) subtype is indicative of acute infection or reactivation, whereas the IgG subtype is indicative of chronic infection. Antibody to HBeAg is suggestive of a nonreplicative state and one in which the antigen has been cleared.⁽⁹⁾

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52 Several studies have reported the prevalence of HBsAg positivity among blood donors from 53 various regions of Nigeria^{10,11}. However data are scarce on serological markers of Hepatitis B 54 virus infectivity among hepatitis B surface antigen negative blood donors. This study determined 55 the prevalence of antibodies to hepatitis B core antigen (anti-HBc), anti-HBeAg, anti-HBs, and 56 HBeAg with the aim to determine the presence of previous HBV infection in Nigerian blood

donors that might have been missed by isolated assay HBsAg. 57 an of This would help in reducing the risk of transfusion of HBV-infected blood units with its 58 attendant complications like liver cirrhosis and hepatocellular carcinoma. 59

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62 Material and Method

This was a descriptive cross-sectional study. The study population consisted of 490 consecutive consenting HBsAg negative blood donors who were also negative for HIV, HCV, Syphilis seen at the blood bank of the University College Hospital, Ibadan over a 6-month periodh. Ethical approval was obtained from the Joint Ethical committee of the University of Ibadan and University College Hospital Ibadan before the commencement of the study.

Semi-structured, self-administered questionnaire was used to obtain subjects' sociodemographic details. Five (5mls) millimeters of venous blood was collected in a plain vacutainer tubes from the participants after obtaining a written informed consent. The blood was allowed to clot and sera separated by centrifugation at room temperature at 3000 gyration, and stored at -20° C in the deep freezer until analyzed.

All samples were screened for HBsAg, using Monolisa HBsAg ULTRA by BIORAD which is a
sandwich third generation enzyme linked immunosorbent assay (ELISA) according to the
manufacturer's instructions. All samples found to be negative for HBsAg were further tested for
anti-HBs, anti-HBc, HBeAg and anti-HBeAg using HBsAb ELISA Kit (DIA.PRO Milano Italy),
HBcAb ELISA kit (DIA.PRO Milano Italy), HBeAg & Ab ELISA Kit (DIA.PRO Milano Italy)
respectively.

Data collected were subjected to descriptive and inferential statistical analysis using the SPSS
version 20 (SPSS Inc, Illinois, USA). Quantitative variables were summarized using mean and
standard deviation while qualitative variables were summarized in frequencies and proportions.
Level of significance was set at 5%.

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86 **RESULTS**

A total number of 550 blood donors were screened for Hepatitis B surface antigen using ELISA,
out of which 60 (10.9%) were positive while 490 (89.1%) tested negative.

Of the 490 that tested negative for HBsAg, 462 (94.3%) were males and 28 (5.7%) were females giving a male to female ratio of 17:1. Their ages ranged from 18-60 with a mean of 32.5 ± 9.5

91 years. More than half of them were married, 290 of 490 (59.2%) while the remaining 200

- 92 (40.8%) were single. Forty-two (8.6%) had primary education while 237 (48.4%) and 210 (43%)
 93 had secondary and tertiary education respectively. Two hundred and one (41%) were employed
- while 71 (14.5%) and 218 (44.5%) were unemployed and students/ housewives respectively.

Eighty-nine of the 490 (18.2%) prospective donors considered fit for blood donation based on 95 Hepatitis B surface antigen negativity were found to be positive for at least one other serological 96 marker (anti-HBc, anti-HBs and anti-HBe) of Hepatitis B virus infection. The sero-prevalence of 97 anti-HBc was 16.9% (83 of 490), out of which 35 (7.1%) were positive for anti-HBc alone, 98 30(6.1%) were positive for both anti-HBc and anti-HBs and 18 (3.7%) were positive for anti-99 HBc, anti-HBs and anti-HBe. Anti-HBs was detected in a total of 54 (11%) donor samples, 100 however, only 6 (1.2%) were positive for anti-HBs alone. The prevalence of anti-HBe was 3.7% 101 (18 of 490). No subject was positive for HBeAg (Table 1 and 2). 102

Table 1: Prevalence of hepatitis B virus markers in Hepatitis B surface antigen negative blood donors.

HBV marker	No	Prevalence (%) (n =490)
Anti- HBc	83	16.9
Anti- HBs	54	11.0

Anti-		
HBe	18	3.7
HBoAg		
преяд		<u>^</u>
	0	0
Total		
	89 ^a	18.2

- a = presence of more than one marker is common
- 107 Anti-HBc= antibody to hepatitis B core antigen; anti-HBs = antibody to hepatitis B surface
- *antigen; anti-HBe = antibody to hepatitis B e antigen.*
- **Table 2: Serological characteristics of Hepatitis B surface antigen negative blood donors.**

Characteristics	Number (Percent %)
Anti-HBc only	35 (7.1)
Anti-HBc + Anti-HBs	30 (6.1)
Anti-HBc+ Anti-HBs + Anti-HBe	18 (3.7)
Anti-HBs only	6 (1.2)
Anti-HBe only	0
HBeAg	0

113 **DISCUSSION**

HBV transmission through blood transfusion is still a great source of concern despite screening 114 for hepatitis B surface antigen (HBsAg) in blood; which is the mainstay of diagnosis for HBV 115 infection in most blood banks in developing countries, including Nigeria. The first serological 116 marker of HBV infection is HBV DNA, followed by HBsAg and HBeAg. Thereafter, anti-HBc, 117 anti-HBe and anti-HBs appear. Antibody to hepatitis B core antigen is the first antibody to 118 appear following acute hepatitis B infection and persists at high level following resolution of 119 infection. ⁽¹²⁾ It is a marker of acute, chronic or resolved infection, although, the degree of 120 protection depends on anti-HBs levels. Anti-HBc remains detectable for life (13) and its 121 significance in screening of blood donors as a way of reducing the residual risk of post 122 transfusion hepatitis B infection has been investigated.⁽¹⁴⁾ 123

There have been concerns about risk of transfusion transmissible HBV infection from blood donors in whom anti-HBc is the only detectable hepatitis B virus marker with no evidence of HBsAg or anti-HBs, particularly in highly endemic regions. The prevalence of "anti-HBc only" in this study of 7.1% is similar to prevalence of 8% reported by Pourazar *et al* among Iranian blood donors ⁽¹⁵⁾. El-Zaatari *et al* ⁽¹⁶⁾ and Salawu *et al* ⁽¹⁷⁾ reported a lower prevalence of 3.7% and 4.4% in Lebanon and Ife respectively among blood donors. However, higher prevalence rates of 18.9% and 30.1% respectively were reported by Asim et al. ⁽¹⁸⁾and Panigrahi *et al* ⁽¹⁹⁾.

The variations in the seroprevalence of anti-HBc in blood donors may be due to differences in 131 the prevalence of HBV infection in these regions. It may also be due to difference in the 132 specificity, sensitivity and positive predictive value of the test method. Likewise, the difference 133 in the socio-cultural practices such as tattooing, scarifications, may explain the variations 134 observed. Countries with intervention measures and health policies such as access to health care, 135 136 immunization practices as found in developed countries are bound to have lower prevalence rate reported. Co-infection of HBV with Human immunodeficiency virus and Hepatitis C virus as 137 suggested by some authors could down-regulate the synthesis of HBsAg.^(20,,21) However, none of 138 the samples in this study tested positive for either HCV or HIV. 139

The importance of anti-HBc in screening for occult HBV infection has been argued extensively.
Studies have demonstrated that some HBsAg-negative individuals but anti-HBc positive

142 continue HBV replication.^(22, 23) The infectivity of blood donations positive for anti-HBc only 143 was reported by Allain *et al* ⁽²⁴⁾ as 4% in immune competent recipients. However, Mosley *et al* 144 ⁽²⁵⁾ reported 17% infectivity of anti-HBc only blood products, although the immune status of the 145 recipients was not indicated. In order to determine the rate of HBV transmission via anti-HBc-146 positive and HBsAg-negative blood donations in this environment, a retrospective studies on 147 regular blood donors and their respective recipients will be necessary.

In this study, both anti-HBc and anti-HBs were found in 30 of 89 (33.7%) individuals 148 constituting 6.1% of the total number (30/490) tested for HBV markers. These subjects were 149 150 considered to be previously infected and to have become immune to HBV infections. It has been documented that blood components positive for anti-HBc and anti-HBs do not appear to transmit 151 HBV and there is clearly an inverse correlation between anti-HBs level and infectivity.⁽²⁶⁾ 152 However, on the contrary, the presence of anti-HBs is not a sign of total HBV eradication as 153 being suggested by Thedja *et*al⁽²⁷⁾. Reactivation of HBV infection despite high levels anti-HBs 154 levels has been revealed by Gartner et al (28) and further reported by Levicnik-Stezinar et al (29). 155 Manzini et al (30) observed that some blood donors with high titres of anti-HBs, over 100IU/L 156 still had detectable HBV DNA. In a more recent study, Ashim et al (31) reported HBV DNA 157 positive cases were detected in donors with low titres of anti-HBc positive and anti-HBs positive 158 159 antibodies.

160 No participant was found positive for hepatitis B e antigen in this study. This is similar to 161 findings by Japhet *etal* ⁽³²⁾ in Ife, Nigeria, but in contrast with finding by Salawu *etal* ⁽¹⁷⁾ who 162 reported a prevalence of 0.22% (1 of 459) in Ile-Ife.

In similar studies done in Africa, El-Ghitany *et al* ⁽³³⁾ reported 0.4% among Egyptian blood donors, while Ashim *etal* $(2010)^{(31)}$, in India found none of the subjects positive for HBeAg. The presence of HBeAg is associated with relatively high infectivity and severity of HBV infection.

This study reveals under-diagnosis of HBV infection with the use of only HBsAg as its surrogate marker and suggests that anti-HBc antibody should be tested routinely in addition to surface antigen in our blood banks.

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170 CONCLUSION

The result in this study highlights the high prevalence of Hepatitis B core antibody in Hepatitis B surface antigen negative blood donors in Ibadan, Southwestern Nigeria. There is need to further screening of our blood donors for other serological markers of HBV even if we cannot embark on Nucleic acid testing due to cost.

175 **RECOMMENDATION**

- Anti-HBc screening of blood donations should be advocated as part of the National
 policy on screening in blood banks with the view of curtailing transmission of HBV
 through this route.
- There is need for a large multi-centre study to determine prevalence of occult hepatitis
 B infection among blood donors in Nigeria and its implications for blood transfusion.
- Retrospective studies should be carried out on regular blood donors and their
 respective recipients to determine the rate of HBV transmission via anti-HBc-positive
 and HBsAg-negative blood donations.
- 184 LIMITATIONS OF THE STUDY
- The sample size may not be fully representative of the entire donor population of the
 blood donors of the hospital.
- 187 2. Hepatitis B virus DNA was not done due to limited resources.

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