

Original Research Article

SEROPREVALENCE OF HEPATITIS B VIRUS AMONG PREGNANT WOMEN ATTENDING ANTE-NATAL CLINIC AT GENERAL HOSPITAL ARGUNGU, KEBBI STATE NIGERIA

ABSTRACT:

Hepatitis B virus infection is caused by Hepatitis B virus, and the virus can be transmitted from infected mother to her new born child during pregnancy. This research work was aimed at determining the prevalence of Hepatitis B virus among pregnant women attending ante-natal clinic (ANC) in General Hospital Argungu. 300 serum samples were assayed using Hepatitis B surface antigen (HBsAg) Rapid Test Strip manufactured by Lab ACON Hangzhou Biotest Biotech Co., Ltd. 38 (12.7%) of the participants have antibodies to HBV. From the research the highest prevalence of HBsAg was found among the age group 20 – 29 years and lowest among the age group of 30 – 39 years. 2nd trimester (4 – 6 months) had the highest prevalence rate of 11.3%, followed by 3rd trimester (7 – 9 months) with 1.3%, while the 1st trimester (1 – 3 months) had zero prevalence (0%). Those that shared sharp objects had the prevalence of 4.6%. Those that had blood transfusion had prevalence of 1.0% while those that are unvaccinated had the highest prevalence of 12.3%. The family type or status i.e. monogamy or polygamy, from the three hundred subjects screened, two hundred and thirty nine family type of the subjects were monogamous and sixty one were polygamous which represent 11.2% and 18.0% respectively. It was observed that fifty had primary education, two hundred and thirty six had secondary education, twenty had tertiary education and ninety four had informal education. Those that had secondary education had the highest prevalence of 7.0% while those with primary education have the least prevalence with 1.0%. This study shows that there are tendencies of vertical transmission from these infected mothers to their new born babies. It is therefore recommended that more studies with advance technology such as PCR should be encouraged and more research should be conducted on a large population in other states of the country so as to establish the endemicity of HBV.

Keywords: sero prevalence, HBV, Pregnant Women, General Hospital.

INTRODUCTION

31
32

33 Hepatitis B virus (HBV) is a DNA virus belonging to the family Hepadna-viridae with Hepatitis
34 B Surface Antigen (HBsAg) being a complex antigen found on its surface [1, 2]. The
35 recognition of hepatitis B virus was first made by Blumberg. When testing the serum of an
36 Australian Aborigine, which he described as Australian antigen and is later termed hepatitis B
37 surface antigen [3]. Hepatitis B virus has been recognized as one of the public challenges
38 worldwide with approximately two billion people infected, an estimated 1 – 2 million annual
39 deaths due to infection and about 400 million persons being chronic carriers [4].

40 In human, hepatitis B virus is among the most important cause of acute inflammation and
41 necrosis of the liver and it is an etiological agent of hepato-cellular carcinoma. HBV attack the
42 liver and cause lifelong infection, cirrhosis of the liver, liver cancer, liver failure and death [5].
43 Hepatitis B virus is transmitted parenterally and most common by transfusion of HBV infected
44 blood or blood products, intravenous drug abuse, from mother to child, needle stick injury, ear
45 piercing, tattooing and other tribal ceremonies, barbers razors etc. infection may also be spread
46 by fomites, sharing of toothbrush, abrasion and sexual contact (hetero or homosexual) with
47 infected persons [6]. Neonates born of chronically infected mothers are 20% at risk of the
48 infection progressing to chronic phase [7].

49 Since detection of HBsAg in serum is an indicative of either acute or chronic phase of HBV
50 infection, this investigation was carried out to detect the prevalence of HBsAg in the Sera of
51 Pregnant women attending General Hospital Argungu with a view of establishing the
52 seroprevalence of HBV infection among pregnant women attending the hospital.

MATERIALS AND METHOD

53

Study Area

54 The research was designed in order to study the prevalence rate of Hepatitis B surface antigen
55 infection among pregnant women in Argungu metropolis. General Hospital Argungu was used
56 for the purpose of this study.
57

Ethical Clearance:

58 Ethical clearance for the study were obtained from ethical committee of General Hospital
59 Argungu. Informed consent was obtained from the Patients.
60

Study Population

61 The study population comprised of three hundred pregnant women attending ANC in Argungu
62 metropolis, kebbi state.
63

64 **Sample Collection**

65 Five milliliter of blood samples was collected by vein puncture from the anti-cutibal foci after
66 swabbing with 70% alcohol from each subject aseptically. The blood was allowed to clot; the
67 serum was centrifuged at 2500 rpm for 20 minutes. It was then separated into sterile sample
68 bottle and labeled with their antenatal number, and was sued for HBV assay.

69 **Laboratory Methods**

70 The ACON rapid test kit was used to test the samples for HBV antibodies. This is a rapid
71 chromatographic immunoassay for the qualitative detection of antibodies to HBV in serum or
72 plasma. The specificity and sensitivity of ACON kits is 98.2% - 100% and 97.2% - 99.8%
73 respectively [8]

74 **Principle of the Test**

75 The ACON HBsAg Rapid Test Strip (serum/plasma) is a qualitative, solid phase, two-site
76 sandwich immunoassay for the detection of HBsAg in whole blood, serum or plasma. The
77 membrane is pre-coated with anti-HBsAg antibodies on the test line region of the strip. During
78 testing, the whole blood, serum or plasma specimen reacts with anti-HBsAg antibodies
79 conjugated particles. The mixture migrates upward on the membrane chromatographically by
80 capillary action to react with anti-HBsAg antibodies on the membrane and generate a coloured
81 line. The presence of this coloured line in the test region indicates a positive result, while its
82 absence indicates a negative result. To serve as a procedural control, a colored line will always
83 appear in the control line region indicating that proper volume of specimen has been added and
84 membrane wicking has occurred.

85 **Procedure**

86 The test strip and the test samples were allowed to equilibrate to room temperature prior to
87 testing. The test strip was removed from the sealed foil pouch. The tape from the test card was
88 peeled off, and the test strip was stocked in the middle of the test card with arrows pointing down
89 on the test card. By holding the dropper vertically, 3 drops of serum (approximately 75µl) was
90 transferred onto the “specimen pad” of the test strip, and the timer was started. The result was
91 read after 15 minutes.

92 **Interpretation of Test Results**

- 93
- 94 • POSITIVE: Two distinct coloured lines appear. One line should be in the control region (C) and another line should be in the test region (T).
 - 95 • NEGATIVE: One coloured line appears in the control region no apparent coloured line appears in the test region (T).
 - 96 • INVALID: Control line fails to appear. Insufficient specimen volume or incorrect procedural techniques are the most likely reasons for control line failure. Review the procedure and repeat the test with a new test strip. If the problem persists, discontinue using the test kid immediately.
- 97
98
99
100

101

102

RESULTS

103 A total of three hundred (300) serum samples were collected from pregnant women attending
104 ante-natal clinic, General Hospital Argungu. Out of the three hundred serum samples screened
105 for HBsAg, thirty eight 38 (12.7%) women were positive for Hepatitis B surface antigen and 262
106 (87.44%) were negative for Hepatitis B surface antigen. (Table).

107 Table 2 shows the age distribution of HBsAg. The age group below 20 years has the highest
108 prevalence of 15.0% followed by 20 - 29 years age group with 12.4% while the 30 – 39 years
109 age group has the least prevalence of 10.0%, followed by 40 – 49 years age group with zero
110 prevalence.

111 Table 3. Shows the Prevalence of HBsAg in relation to trimester of subjects. 2nd trimester (4 – 6
112 months) had the highest prevalence rate of 13.9%, followed by 3rd trimester (7 – 9 months) with
113 8.2%, while the 1st trimester (1 – 3 months) had zero prevalence (0%).

114 Table 4. Show the prevalence of HBsAg in relation to risk factors. Those that shared sharp
115 objects had the prevalence of 17.5%. Those that had blood transfusion had prevalence of 18.7%
116 while those that are unvaccinated had prevalence of 15.8%. The family type or status i.e.
117 monogamy or polygamy, from the three hundred subjects screened, two hundred and thirty nine
118 family type of the subjects were monogamous and sixty one were polygamous which represent
119 11.2% and 18.0% respectively.

120 Table 5. Shows prevalence of HBsAg in relation to educational status of subjects. From the table,
121 it was observed that fifty had primary education, two hundred and thirty six had secondary
122 education, twenty had tertiary education and ninety four had informal education. Those that had
123 tertiary education had the highest prevalence of 30.0% while those with primary education have
124 the least prevalence with 6.0%.

125

126 **TABLE 1: Overall Result of HBsAg Prevalence**

127

Total number	No. of positive (%)	No. of negative (%)
300	38 (12.66)	262 (87.4)

128 HBsAg = Hepatitis B surface antigen, No. = number, % = percent.

129 **TABLE 2: Age Distribution of HBsAg Among the Patients**

130

Age (years)	No. screened	No. positive	(%)
Below 20	80	12	15.0
20 – 29	177	22	12.5
30 – 39	40	4	10.0
40 – 49	3	0	0.0
Total	300	38	

131 HBsAg = Hepatitis B surface antigen, No. = number, % = percent.

132 **TABLE 3: Distribution of HBsAg Based on Trimester**

133

Trimester	No. screened	No. positive	(%)
1 st (1-3 months)	8	0	0
2 nd (4-6 months)	243	34	11.3
3 rd (7-9 months)	49	4	1.3
Total	300	38	

134 HBsAg = Hepatitis B surface antigen, No. = number, % = percent, 1st = First, 2nd = Second, 3rd =
135 Third.

136

137 **TABLE 4: Distribution of HBsAg with Respect to Risk Factors**

138

Age (years)	No. screened	No. positive	(%)
1. Sharing with sharp object			
Yes	80	14	17.5
No	220	24	10.91
Total	300	38	
2. Blood transfusion			
Yes	16	3	18.75

No	284	35	12.32
Total	300	38	

3. Vaccination

Yes	67	1	1.49
No	233	37	15.88
Total	300	38	

4. Family status

Monogamy	239	27	11.29
Polygamy	61	11	18.03
Total	300	38	

139 HBsAg = Hepatitis B surface antigen, No. = number, % = percent.

140 **TABLE 5:** Distribution of HBsAg Based on Educational Status

141

Education	No. screened	No. positive	(%)
Primary	50	3	6.0
Secondary	136	21	15.4
Tertiary	20	6	30.0
Informal	94	8	8.5
Total	300	38	

142 HBsAg = Hepatitis B surface antigen, No. = number, % = percent.

143

DISCUSSION

144 The prevalence rates of HBV vary according to the endemicity of the infection in a given area.
 145 Kong *et al.*, [9] reported prevalence rate of 10.0% among pregnant women in Hong Kong, Lin *et*
 146 *al.*, [10] reported 12.0% prevalence rate from Taiwan, while 17.3% was reported for Burkina
 147 Faso [11]. In Nigeria, 11.6% prevalence rate has reported from Maiduguri, 4.3% from Port
 148 Harcourt, 5.7% from Ilorin, in Lagos, prevalence was reported to be 4.4% and 8.3% from Zaria
 149 [12, 13, 14, 15 and 16]. Very high prevalence rate are mostly reported from the developing
 150 nations in Asia and Africa.

151 Hepatitis B is one of the diseases of mankind and is a serious global health problem, caused by
 152 the hepatitis B virus. It has been established that HBV infection can be transmitted from mother
 153 to child during birth. High prevalence of HBV among pregnant women increases chances of

154 HBV in children. From the result obtained in this study, out of 300 samples screened for HBsAg,
155 38 samples were found positive to hepatitis B virus infection (12.7%). This is in agreement with
156 earlier reports of 13.8%, 10.0%, 11.6% and 12.0% from Lagos, Hong Kong, Maidurugi and
157 Taiwan respectively [15, 9, 12, and 10].

158 Within Nigeria, results from this study is higher than the 4.3%, 5.7% and 8.3% reported from
159 Port Harcourt, Ilorin and Zaria respectively [1, 2 and 11]. The decrease in prevalence rates among
160 some Nigerians could be due to anti HBsAg vaccination policy of the government. Detection of
161 HBsAg among the study population has confirmed statement that detection of HBsAg in serum
162 is indicative of active acute or chronic hepatitis B virus infection [17].

163 On the basis of age group, the highest prevalence rate (15.0%) was found among those below 20
164 years, followed by 20 - 29 years with 12.4% while 40 – 49 years had 0.0% prevalence. This age
165 of infection correlate well with the age of greatest sexual activity especially among women of
166 childbearing age, supporting the role of sexual intercourse in the transmission of hepatitis B virus
167 infection. In this study, women of their second trimester of pregnancy had the highest prevalence
168 of 13.9%, contrary to observations of Lilavati *et al*, [8] that the third trimester in pregnant women
169 had the highest prevalence rate.

170 Considering various risk factors, pregnant women with history of blood transfusion have the
171 highest prevalence of 18.7%, indicating the significance of screening blood for HBV infection,
172 followed by sharing sharp object with 17.5%, which might be one of the most pre-disposing of
173 HBV infection among these pregnant women. From the study, it was observed that highest
174 number of HBV infectious was found among polygamy type of family (18.0%), while there are
175 few positive cases of HBV infection among monogamy family type, this shows that family type
176 (monogamy or polygamy) does not have much significant in the prevalence of HBV infection in
177 Argungu metropolis, this is because the spread of most STD's does not depend on family type
178 but depend on so much on the faithfulness of partners which are involved. Those who belong to
179 the polygamy family who are infected may be due to sharing of husband who is unfaithful or
180 who becomes infected by an unfaithful co-wife.

181

182
183

CONCLUSION

184 The conclusion from this study is that it is evident that HBV infection is present or occurred
185 among these pregnant women hence there is still need to educate them about the danger
186 associated with this virus infection, its possible routes of transmission and possibilities of vertical
187 transmission to their new born babies from infected mothers.

188
189

RECOMMENDATIONS

190 Based on the result obtained in this study the following are recommended.

- 191 1. Every pregnant woman for ante-natal visit should be screened for HBsAg and
192 government should subsidize HBsAg screening not only for pregnant women but also for
193 those preparing for pregnancy so that adequate precaution should be taken.
- 194 2. There should be campaign to create awareness on the modes of transmission, the risk
195 factors as well as how to control the spread of HBV should be intensified and increase
196 where there is no trust for one another.
- 197 3. Blood for transfusion, blood should be properly screened with latest and modern
198 equipment and reagents that can detect minute antibody or antigen in the blood.
- 199 4. Sharing of personal items such as tooth brush, razor blades should be discouraged among
200 the populace.
- 201 5. Health personnel in close contact with infected individuals should be given HBV vaccine
202 and possible precautions to avoid hospital infection
- 203 6. Infected individual should be treated to reduce spread of the virus in the community.
- 204 7. Routine vaccination of previously unvaccinated children and vaccination of adults at
205 increased risk for infection.
- 206 8. prevention of perinatal HBV infection through routine screening of all pregnant women
207 for HBV infection and by providing immunoprophylaxis to infants born to infected
208 women or to women of unknown infection status.

209

210

211

212

213

214

REFERENCES

215

- 216 1. Brooks, G. F., Carroll, K. C., Butel, J. S., Morse, S. A. Medical Microbiology, 24th
217 Edition; International Edition, McGraw Hill Publishers, New York, USA; 2007.
- 218 2. Hallinger, F. B. and Dienstag, J. L. Hepatitis B and Hepatitis D virus in: Murray, P. R.,
219 Baron, E. J., Tenover, F.C. and Tenover, R. H. (eds): Manual of clinical
220 microbiology, 7th Edition, American Society for Microbiology Asna Press, Washington
221 DC: USA; 1990. P. 1025 – 1042
- 222 3. Rajesh, B., Rattan.L. I. Essentials of medical microbiology, 4th edition. Jaypee Brothers
223 Medical Publisher (P) LTD; 2008. P. 391 – 396.
- 224 4. WHO. The modes of HIV transmission. Fact sheet. 2000.

- 225 5. Koneman, B. W., Allen, S. D., Winn, Jr U. N. C. Diagnostic of infectioncsued by
226 viruses, Chlamydia, rickettsia and related organism and diagnostic microbiology. J.P.
227 Lippingcolt 6th Edition; 1992. P. 1000 – 1006.
- 228 6. Ugwuja, E., Ugwu, N. Seroprevalence of hepatitis B surface antigen and liver function
229 tests among adolescents in Abakaliki, South Eastern Nigeria. *The Internet Journal of*
230 *Tropical Medicine*. 2009; **6** (1): 220 – 229.
- 231 7. Lin, K. W., Kirchner, J. T. *Hepatitis B Journal of American Academy of family*
232 *physicians*. 2004; **69** (1): 75 – 86.
- 233 8. Blumberg, B. S. The discovery of Australian antigen and its relation to viral hepatitis.
234 *Vitro*. 1971; 7:22
- 235 9. Kong MS, Liang DC, Shau WY, Chen DS. Universal hepatitis B vaccination in Taiwan
236 and the incidence of hepatocellular carcinoma in children. *Taiwan Childhood Hepatoma*.
237 *N Engl J Med*. 1997; 336(26):1855-1859.
- 238 10. Lin, H. H., Kao, J. H., Chang, T. C., Hsu, H.Y., Chen, D.S. Secular trend of age specific
239 prevalence of hepatitis B surface and antigenemia in pregnant women in Taiwan. *Journal*
240 *of Medical Virology*. 2003; **69** (1): 75 – 86.
- 241 11. Collenberg, E., Ouedraogo, T., Ganame, J., Ackernscher, H. Kynas-wolf, G., Becher, H.,
242 Kouyate, B. Krauslich, H.C., Sangave, L., Tiet, D.M. Sero-prevalence of six different
243 viruses among pregnant women and blood donors in rura and urban Burkina Faso: A
244 comparative analysis. *Journal of Medical Virology*. 2006; **78** (5): 638 – 192.
- 245 12. Harry, T. O., Bajani, M. D., Moses, A. E. Hepatitis B virus infection among blood donors
246 and pregnant women in Maiduguri, Nigeria. *East African Medical Journal*. 1994; **70**: 596
247 – 597.
- 248 13. Akani, C. I., Ojule, A. C., Oporum, H. C., Ejilemele, A. A. Seroprevalence of HBsAg in
249 pregnant women in part of Port Harcourt, Nigeria. *Post Graduate Medical Journal*. 2005;
250 **12** (4): 266 – 270.
- 251 14. Agbede, O. O., Iseniya, J. O., Kolewale, M. O., Ojuowa, A. Risk factors and
252 Seroprevalence of Hepatitis B antigenemia in mothers and their preschool children in
253 Ilorin, Nigeria. *Therapy*. 2007; **4** (1): 67 – 72.
- 254 15. Nasidi, A. T. O., Vyazor, S. O., Numumbe, G. M. R., Azzan, B. B., Ancinlev, V. A.
255 Prevalence of Hepatitis B infection marker in two different geographical areas of
256 Nigeria.Proceedings of the first international conference. Lagos, Nigeria; 1983.
- 257 16. Luka, S. A., Ibrahim, M. B., Iliya, S. N. Seroprevalence of hepatitis B surface antigen
258 among pregnant women attending Ahmadu Bello University Teaching Hospital Zaria.
259 *Nigerian Journal of Parasitology*.2008; **29** (1): 38 – 41.
- 260 17. Lilavati, G., Chandra, M.P., Umakanta, N. Incidence of HBsAgcarriers state in pregnancy
261 in eastern Orissa. *Journal of Obbstetric and Gynaecology India*. 2004; **54** (2): 136 – 138.
- 262
263