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3 **SEROPREVALENCE OF HEPATITIS B SURFACE ANTIGEN IN**  
4 **PREGNANT WOMEN OF GENERAL HOSPITAL AGBOR, DELTA**  
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8 **Abstract:**  
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10 Reportedly transmitted through unprotected sexual intercourse with infected person(s),  
11 experts have estimated new cases of hepatitis B virus (HBV) infections to be over 70,000 per  
12 year in the United States. With little or no records of such in Nigeria, this study investigated  
13 the prevalence of hepatitis B surface antigen at different trimesters of pregnancy in women  
14 who visit general hospital Agbor, Delta state, Nigeria; for antenatal care. A total of one  
15 hundred (100) pregnant and fifty (50) non-pregnant (control) women were ethically recruited  
16 for the exercise. They were then sub-grouped by age and duration of pregnancy (trimester);  
17 and an Acon serological strip was used to obtain blood samples from each subject. Obtained  
18 blood was then assayed for the presence of hepatitis B virus (in serum) and compared with  
19 those of control (non-pregnant) group. Following careful comparison of differences in mean  
20 (using the Analysis of variance), study found a 5% prevalence rate (of hepatitis B) in  
21 pregnant than non-pregnant (2% prevalence) women. Study also observed a statistically  
22 significant increase in hepatitis B surface antigen for non-pregnant women of age bracket 20-  
23 24 years (2.6% prevalence) to pregnant women of between 20-24 years. HBV infection  
24 therefore has high prevalence rate in pregnant than non-pregnant women as they are often  
25 more exposed to unprotected sexual intercourse. We recommend regular and continuous  
26 HBV screening in pregnancy to help circumvent HBV infection related ailments and  
27 complications. The same is also suggested for non-pregnant women for purpose of proper  
28 vaccination.

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29 **Keywords:** Hepatitis B, Pregnancy, Sero-prevalence, Antenatal  
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31 **INTRODUCTION**

32 Hepatitis, inflammation of the liver caused by viruses, bacterial infections, or continuous  
33 exposure to alcohol, drugs, or toxic chemicals, such as those found in aerosol sprays and  
34 paint thinners [1&2]. Hepatitis can also result from an autoimmune disorder, in which the  
35 body mistakenly sends disease-fighting cells to attack its own healthy tissue, in this case the  
36 liver. Irrespective of its cause, hepatitis reduces the liver's functional abilities, including its  
37 filtering prowess for harmful infectious agents from the blood, as well as its capacity to store  
38 blood sugar and converting it to usable energy forms that are necessary for life [3-5].

39 Depending on cause and overall health infected individual, the symptoms of hepatitis  
40 vary significantly, with selected cases showing few noticeable symptoms [6]. If present  
41 however, symptoms may include general weakness and fatigue, loss of appetite, nausea,

42 fever, and abdominal pains/tenderness. Another symptom is jaundice, which apparently  
43 occurs as a yellowing of the skin and eyes due to the liver's failure to break down excess  
44 yellow-colored bile pigments in the blood. In acute hepatitis, symptoms often subside without  
45 treatment within a few weeks or months. About 5 percent of cases develop into an incurable  
46 form of the disease called chronic hepatitis, which may last for years [7&8]. Chronic hepatitis  
47 causes slowly progressive liver damage that may lead to cirrhosis, a condition in which  
48 healthy liver tissue is replaced with dead, nonfunctional scar tissue. In some cases, cancer of  
49 the liver develops [6].

50 Although it has many different causes, hepatitis most often results from infection by  
51 one of several hepatitis viruses. All hepatitis viruses are contagious, but each is differently  
52 passed from one person to another [9]. Over the las decade, the WHO issued its first  
53 guideline for the treatment of chronic hepatitis B in Nigeria, positing that globally, some over  
54 240 million people have chronic hepatitis B infection with increasing risk of dying, and  
55 highest rate more to be found in Africa and Asia[7]. In Nigeria, even though no specific  
56 treatment option is fully documented, effective medicines however exists that can prevent  
57 this condition in people. However, due to lack of clear evidence-based guide and poor living  
58 standard, most sufferers who need these medicines cannot afford them [7].

59 Recently, A WHO report has it that HBV is a common infectious disease that  
60 accounts for a major cause of global health problems. The report further posits that an  
61 estimated 2 billion earthlings are sero-positive of past or present HBV infection, with 350  
62 million of such cases reportedly chronically infected, putting sufferers at risk of HBV-related  
63 liver diseases [10]. In another study, HBV was reported to be 50 to 100 times more infectious  
64 than the AIDS due to HIV, and 10 times more infectious than hepatitis C virus (HCV).  
65 Tentatively, it is an important cause of liver diseases; with associated co-morbidities and liver  
66 failure, cirrhosis and liver cancer [11].

67 Epidemiologically, the prevalence of HBV varies from as low as 2% in developed  
68 countries to about 8% in developing countries. Here, even though sex, age and socio-  
69 economic status are reported risk factors that exacerbate it [12], however, available studies  
70 have suggest its degree of endemicity to often correlate with predominant mode of  
71 transmission. HBV disease has an enormous impact on health and national economy of many  
72 countries, and its severity is highly variable and quite unpredictable. The minimum infectious  
73 dose is so low that such practices like sharing of tooth brush or a razor blade can elicit its  
74 quick transmittion [13 & 14]. Hepatitis B virus also shares similar routes of transmission with  
75 HIV [6], Currently having four recognized modes of transmission (Viral Hepatitis Prevention

76 Board, 1996); from mother to child at birth (prenatal), by contact with infected person  
77 (horizontal), by sexual contact and by exposure to blood or other infected fluids [7].

78 Irrespective of age group, HBV reportedly affects people across diverse spheres and  
79 ethnicities, predominantly more in young adults than the elderly [1]. Currently, Nigeria ranks  
80 high in the list of highly endemic HBV infected countries; with about 75% of its population  
81 reportedly likely to have been exposed to HBV at one time or the other in their life [2, 4, and  
82 9]. To this point, current study was crafted to investigate the prevalence of HBV surface  
83 antigen in pregnant women in Nigeria, using general hospital, Agbor as a case study

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### 85 **Aim of Study**

86 Study aimed at examining the sero-prevalence of hepatitis B surface antigen in pregnant  
87 women of general hospital agbor, Delta State. Specifically, study investigated the prevalence  
88 of HBV by age and trimester of pregnancy

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## 90 **Materials and Methods**

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### 92 **Study Design**

93 The study was analytical, and was designed to determine (by age and trimester), the  
94 incidences of Hepatitis B in pregnant women; specifically those that regularly attend  
95 antenatal screening at the general hospital in Agbor, Delta State.

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### 97 **Study Population**

98 The population for this study comprised of pregnant women who attended the antenatal care  
99 unit of the general hospital, Agbor, Delta State. A total of one hundred (100) pregnant women  
100 and fifty (50) non-pregnant women (control) were randomly sought from the population.

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### 102 **Sample Collection**

103 With the aid of a Pasteur pipette (specimen dropper), blood samples were collected from  
104 subjects (in 21 days, spanning a total of three weeks), and serology for hepatitis B was  
105 conducted on obtained blood samples to ascertain the presence of HBV in blood. Analysis of  
106 obtained sample for HBV positivity was done with the Acon Serological strips (ELISA).

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### 108 **Procedure**

109 About three (3) drops of blood from each collected sample were vertically dropped into a  
110 sample pad of test strip, which was placed on the test cards. Thereafter, a drop of buffer was  
111 dispensed on the strip containing sampled plasma or serum. The mixture was then left to react  
112 with the HBsAg antibodies and conjugated particles for a period of 15 minutes; following  
113 which result was read.

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### 115 **Principle**

116 The Hepatitis B surface antigen rapid strip test is a quantitative solid phase of a two site  
117 sandwich immunoassay procedure, used to detect the presence of hepatitis B serum antigen  
118 (HBsAg) in the whole blood serum and/or plasma. Here, the whole blood serum or plasma  
119 specimen reacts with anti HBsAg antibodies, and the conjugate migrate upwards to the  
120 membrane by capillary action. This leads to their reaction with the membrane to generate a  
121 colour line. However, presence of two colour lines will indicate a positive result, while a  
122 single colour line will indicate a negative result.

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### 124 **Ethical Considerations**

125 Ethical clearance was obtained from the general hospital, Agbor, before actual sample  
126 collection, consent forms were administered to seek participants' permission. Only subjects  
127 whose consent we got were actually investigated.

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### 129 **Results**

130 Table 1 (below) shows the prevalence rate of Hepatitis B surface antigen (HBsAg) amongst  
131 pregnant and non-pregnant women. From the table, total number of sampled pregnant women  
132 was 100 (test group). Of these, 5% were seen to be positive for HBsAg, with 2% of total non-  
133 pregnant women (100) showing positive for HBsAg. Apparently, 95% of sampled pregnant  
134 women were negative for HBsAg, with non-pregnant cases having 98% HBsAg negative.

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136 **TABLE 1: Prevalence of Hepatitis B surface antigen between Pregnant and Non-**  
137 **Pregnant Women**

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No Screened	No Positive	No Negative	Mean	p-value
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Pregnant Women	100	5(5%)	95(95%)	1.24	Insignificant
Non Pregnant Women (Control)	100	2(2%)	98(98%)	1.24	Insignificant

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Table II below shows the percentage prevalence by age of HBsAg in pregnant and Non-Pregnant women. Therefore, pregnant women between ages 20-24, 25-29 and 30-34 had a relatively high prevalence rate of 10%, 4.7% and 3.7% respectively. All were statistically significant (at  $p < .05$ ) upon comparison with Non-pregnant women.

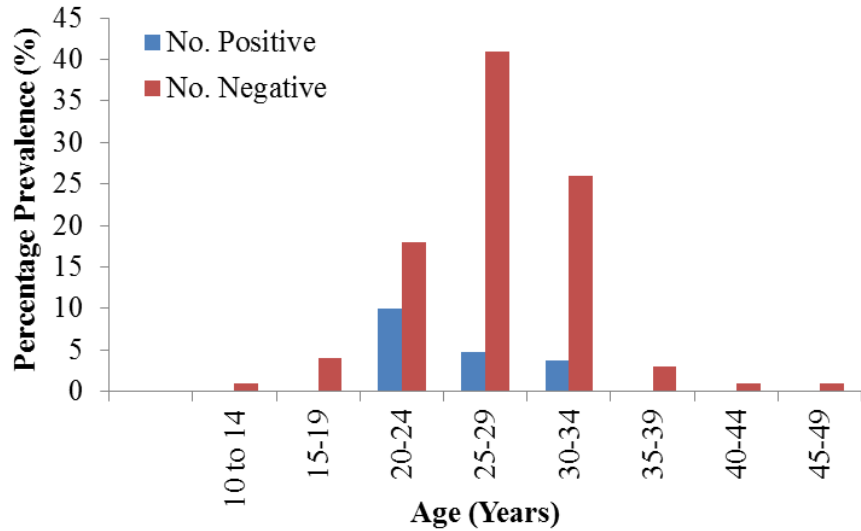
**TABLE II: Sero-Prevalence of Hepatitis B surface antigen by Age in Pregnant and Non-Pregnant Women**

Age (Years)	Pregnant Women	No. Positive	No. Negative	Non-Pregnant Women	No. Positive	No. Negative
10-14	1	0	1 (1%)	4	0	4 (4%)
15-19	4	0	4 (4%)	43	1 (2.3%)	42 (42%)
20-24	20	2 (10%)	18 (18%)	38	1 (2.6%)	37 (37%)
25-29	43	2 (4.7%)	41 (41%)	10	0	10 (10%)
30-34	27	1 (3.7%)	26 (26%)	2	0	2 (2%)
35-39	3	0	3 (3%)	1	0	1 (1%)
40-44	1	0	1 (1%)	2	0	2 (2%)
45-49	1	0	1 (1%)	0	0	0
Total	100	5	94	100	2	98

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The figure I (below) compares the percentage prevalence of Hepatitis B Virus (by age) for pregnant women. Here, significant number of sampled pregnant women between ages 20 through 34 years tested positive for HBV,

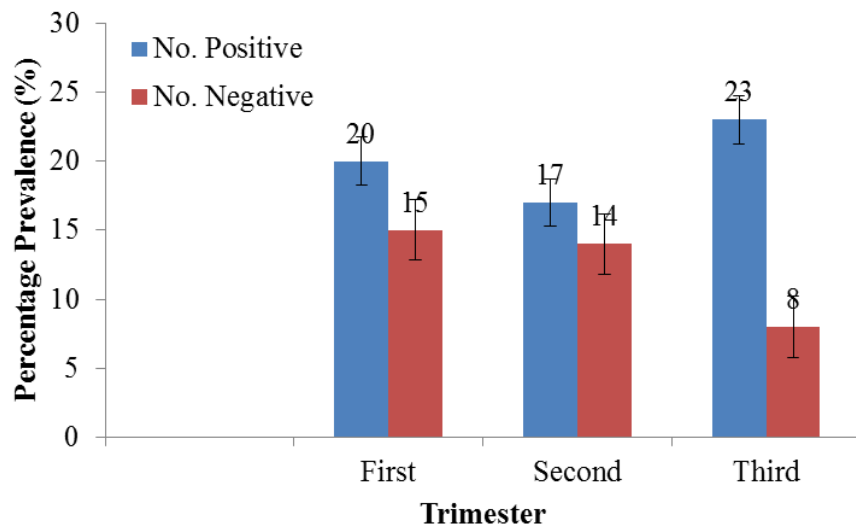
**Figure I: Comparing Percentage Hepatitis B Positives to Negatives (by age) in Pregnant Women**



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The figure II (below) represents changes in HBV with various trimesters of pregnancy in participants. Visible here is prevalence rates of (number of positive) 23%, 20% and 17% for Third, First and Second Trimesters respectively. This implies that sero-prevalence of HBV is trimester independent.

**Figure II: Comparing Percentage Prevalence of Hepatitis B by Trimester in Pregnant Women**



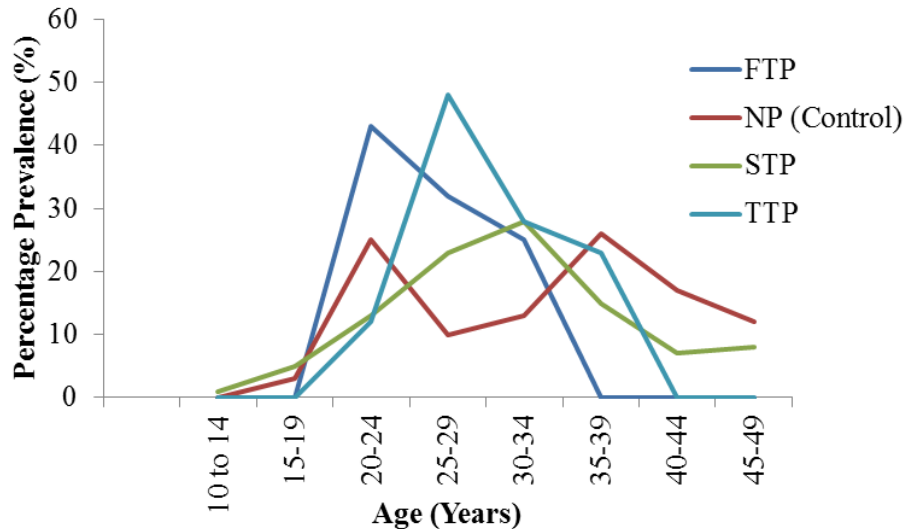
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Figure III (below) shows a comparison between sero-prevalence of HBV (by age) with various trimesters of pregnancy (for pregnant women) and non-pregnancy states. As seen, higher prevalence rate occur in women of age brackets 15-19, 20-24, 25-29, 30-34, and 35-39; with those between 25-29 posing highest upon comparison for all sampled women. Here,

173 FTP = First Trimester Pregnant women, NP=Non-Pregnant women, STP=Second Trimester  
174 Pregnant women, and TTP=Third Trimester Pregnant women

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176 **Figure III: Comparing Percentage Sero-Prevalence of Hepatitis B by Age and**  
177 **Trimester of Pregnancy**



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## 181 Discussion

182 According to the centre for disease control and prevention (CDC) 2005, the  
183 prevalence of hepatitis B infection varies in different parts of the world. Investigation on the  
184 prevalence rate of Hepatitis B surface antigen among pregnant women (test) in General  
185 Hospital, Agbor, Delta State. Investigation was also carried out on non-pregnant women  
186 (control). Result showed that the prevalence rate in pregnant women was higher than in non-  
187 pregnant women, being 5% in pregnant women and 2% in non-pregnant women in the  
188 general study population.

189 Studies have shown various percentage of Hepatitis B in pregnant women across  
190 different countries. The CDC goals of 2005 included an objective that by the year 2000, 90%  
191 of pregnant women would be screened in health centres before delivery. For current study,  
192 General Hospital, Delta state showed a 5% prevalence rate of HBsAg in pregnant as against  
193 2% rate for non-pregnant women. This prevalence is higher than that of in general population  
194 as reported by Zahedan of Iran [15] who showed less than 3% prevalence in Barbers. In  
195 another study among 103 barbers, Zahedan showed 8.7% prevalence for HBsAg as well.

196 Similarly, table II of current study shows Sero-Prevalence of Hepatitis B surface  
197 antigen by Age in Pregnant and Non-Pregnant Women. From the table, higher percentage of  
198 HBsAg positive subjects was seen in pregnant women of age 20-24 years with higher

199 percentage (41%) of same age bracket proven to be HBsAg negative. The exact reason for  
200 subjects within this age (20-24 years) having to be positive may not be farfetched. It is most  
201 likely traceable to the fact that it is the age with high level of productivity. Expectedly,  
202 unprotected sexual intercourse is probable reason; more so that the said women were married.  
203 This finding aligns with that of Berker et al (2009) [1] who recommended HBsAg test in  
204 pregnancy situations.

205 Hepatitis B virus is highly contagious. Usually, the disease is passed on during the birth  
206 process or during a vaginal delivery or a C-section [16]. When babies become infected with  
207 Hepatitis B, they have a 90% chance of developing a lifelong, chronic infection. As many as  
208 1 in 4 people with chronic Hepatitis B develop serious health problems. Hepatitis B can cause  
209 liver damage, liver disease, and liver cancer [15, 16]. In part with acute hepatitis B, vertical  
210 transmission occurs in up to 10% of neonates when acute infection occurs in third trimester  
211 [1]. Although the mother usually becomes jaundiced during the acute stage, 50% of cases  
212 have no symptoms, which is one of the reasons all pregnant women should routinely test for  
213 HBsAg at the first prenatal visit.

214 Figure III compares percentage sero-prevalence of Hepatitis B by age and trimester of  
215 pregnancy. Here, it is obvious that majority of pregnant women that tested seropositive for  
216 Hepatitis B were in their first and third trimesters, being of age brackets 20-24 and 25-29  
217 years respectively. Though the possible reason for this is inexplicable, one cannot but think  
218 that the prevalence rate was independent of trimester. For figure II also, third trimester of  
219 pregnancy appears to have highest prevalence for HBsAg than first (higher) and third  
220 trimesters respectively, with each recording a prevalence rates of 23%, 20% and 17%  
221 respectively for third, second and first trimesters. A statistically significant difference ( $p <$   
222  $.05$ ) was returned from differences in mean between trimesters with analysis of variance. This  
223 also implies no relationship (correlation) between prevalence of HBsAg and trimester of  
224 pregnancy.

225 In prenatal screening exercises, HBsAg tests are highly recommended (Eulerciary et  
226 al., 2003). If testing has not been during pregnancy, it should be done at the time of delivery.  
227 If a pregnant woman has a positive test, her infant should be vaccinated against hepatitis B,  
228 and made to receive hepatitis B immunoglobulin. This will help reduce the risk that the infant  
229 will become infected to a range from zero to 3% [16]. There is therefore no perfect report on  
230 the pregnancies of HBsAg positivity in other regions of the world, though in most countries,  
231 pregnant women also have higher prevalence rate of Hepatitis B than the non-pregnant  
232 women. This implies that pregnant women are most exposed to Hepatitis B infections due to



233 high exposure to sex. Also, due to the Hepatitis B infection in pregnant women, infections in  
234 neonates arise, causing hepatitis B in children. Also, due to documentation of high prevalence  
235 rate in pregnant than non-pregnant women, it becomes appropriate for pregnant women to  
236 take precautionary measures before, during and after pregnancy to ensure they are screened  
237 for HBsAg and get vaccinated against the virus. Vaccination is also suggested for non-  
238 pregnant women.

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## 240 **Conclusion**

241 Hepatitis is an inflammation of the liver. Some **types** can be caused by various serious  
242 diseases, and may be sexually transmitted. In view of this, current study investigated the  
243 serum prevalence of Hepatitis B surface Antigen by age (and trimester) in pregnant women  
244 from general hospital, Agbor, Delta State. Upon investigation, study observed a 5%  
245 prevalence rate (of hepatitis B) in pregnant than non-pregnant (2% prevalence) women. Study  
246 also observed a statistically significant increase in hepatitis B surface antigen for non-  
247 pregnant women of age bracket 20-24 years (2.6% prevalence) to pregnant women of  
248 between 20-24 years.

## 249 **Recommendations**

250 It is suggested that women regularly screen for presence of hepatitis B in their blood  
251 before, during, and after pregnancy for proper prognosis and vaccination against the virus.  
252 Awareness programmes should also be encouraged, especially for rural communities to keep  
253 the public apprised of the cause, transmission, and symptoms of infection from this virus. **We**  
254 **also recommend for further research in this area; approaching it from a more sophisticated**  
255 **way with higher sample size.**

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