

Hepatitis B Viral Load (HBV-DNA) with Age and Sex Stratifications in Bangladeshi People

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Abstract

Background: Hepatitis B virus surface antigen (HBsAg) is used for the detection of Hepatitis B virus (HBV) infection and to predict disease progression.

Objectives: The main objective of this study was to observe the pattern of HBV viral load levels among people in terms of age and sex distribution in Bangladesh.

Method: Blood specimen was collected from 585 objects with HBsAg Positive and assayed for the quantity of hepatitis B virus using PCR based technique.

Results: It is found from the study that the mean viral load was 353,500,000 DNA copies/ml for 20-35 age group, 249,300,000 DNA copies/ml for 35-50 age group and 104,800,000 DNA copies/ml for 50-65 age group. The median HBV viral loads for male and female were 58,494 and 103,287 DNA copies/ml, respectively.

Conclusion: High viral load was observed in the 21-35 and 36-50 age group while females are at most risk due to HBV infection as their viral load is higher compared to male.

Keywords: Viral load; HBV infection; HBV DNA; Bangladesh

Introduction

Hepatitis B virus (HBV) infection is considered as a serious public health concern worldwide since more than 350 million people are chronic carriers of HBV [1]. Persistent HBV infection is a risk factor for the development of Hepatocellular Carcinoma (HCC) [2]. According to Liver Foundation of Bangladesh incidence of Hepatitis-B infection in Bangladesh is about 4%-7% of total population. About 3.5% of pregnant mothers in Bangladesh are the carriers of the hepatitis B virus. Among them those are HBeAg positive (about 90%) will transmit the virus to their offspring [3]. According to a study published in 1996, 36% of the Hepatocellular carcinoma is associated with hepatitis B infection in Bangladesh [4]. Influence of age and sex on the development of HBV carrier state has been studied and a consistent relation was found [5].

Hepatitis B Virus surface Antigen (HBsAg) is an important marker for the detection of HBV infection and also it can be used to predict disease progression [6]. In Bangladesh although there is setup for quantitative detection of HBV viral load but these tests are quite expensive and rare [7]. It should be also consider that co-infections and super-infections with other hepatitis viruses (e.g. HCV) may also occur [8]. Therefore, person's disease history, age, sex, vaccination status and previous tests results should be considered to guide

appropriate testing. The purpose of our study was to present the pattern of HBV viral load levels among people in terms of age and sex distribution in Bangladesh.

Materials and Methods

All samples were collected from six different metropolitan city of Bangladesh. Metropolitan cities have supply water system for drinking purpose and also sewage system parallel, most of the slum area have no proper sanitation system especially in the slum are of Dhaka city and hospitals do not have legitimate waste management system.

Specimen collection and processing

A bio-data was raised for each of the 585 subjects containing details of their age and sex. All subjects were informed about this study and only included if they permitted to use their data. Inclusion criteria for the subjects were age from 12 to 80 years old and HBsAg Positive. Only the first test carried out by a patient was included in this study. The specimen of choice for the diagnosis of HBV infection was blood. In brief, 5 ml of blood was collected from each individual into EDTA treated tubes and centrifuged for 10 minutes. The plasma was separated and stored at -20°C until analyzed. Patients had been previously confirmed as HBsAg positive and well aware about their inclusion in this study, prior to assessing the HBV viral load test.

HBV DNA viral load assay

Samples were assayed for the quantity of hepatitis B virus according to the SMART CYCLER HBV monitor test Version 2.0, a PCR based technique. The HBV viral load results were expressed in DNA copies/ml.

Statistical analyses

Graph Pad Prism 5 was used for statistical analysis and test of significance was done using Kruskal-Wallis statistical packages [9]. Differences of $p < 0.05$ were taken to be statistically significant at 95% confidence interval.

Values\Age Distribution (Years)	<20	20-35	36-50	51-65	>65
Mean	1,562,000	273,800,000	228,100,000	104,000,000	20,334
Median	21,546	28,749	57,646	54,823	22,565
Lower 95% CI of mean	-793,239	120,400,000	77,820,000	-5,926,000	-23,054
Upper 95% CI of mean	3,918,000	427,200,000	378,300,000	214,000,000	63,723

Table 1: The age distribution of the HBV viral load

The participant's age range was from 20 to 65 Years. We have equally classified the age range into three categories: 20-35, 36-50 and 51-65 years of old (Table 1). As the life expectancy of Bangladeshi people is 69.8 years [10]; therefore this age distribution is covered the entire population of Bangladesh except younger population and children. The mean viral load found 353,500,000 DNA copies/ml for 20-35 age group, while it was 249,300,000 DNA copies/ml for 35-50 age group and 104,800,000 DNA copies/ml for 50-65 age group. The median HBV viral loads for male and female were 58,494 and 103,287 DNA copies/ml, respectively ($P > 0.005$) (Table 2). Male to female ratio was 2:1.

Values/Sex Distribution	Male	Female
Mean	218,800,000	209,800,000
Median	36,627	36,548
Lower 95% CI of mean	113,300,000	57,840,000
Upper 95% CI of mean	324,300,000	361,700,000

Table 2: The sex distribution of the HBV viral load

Discussion

Infection with HBV is under control in developed countries, but it is still a serious public health problem in developing countries like Bangladesh. In this study we examined the viral load pattern of those assessing laboratory services at our site, being the only laboratory at present where HBV viral load is carried out commercially within the country. As such samples were received from a wide range of localities across the country. There was a wide age range observed by the patients, with the highest prevalence being between 30-39 years (Figure 1).

Results

In brief, HBV viral load is found minimum 256 and maximum 15,645,417,077 DNA copies/ml among the participants sample. The mean value of the viral load is observed using statistical analysis 263,000,000 DNA copies/ml with 95% Confidence Interval (CI) (lower mean: 157,100,000; upper mean: 368,900,000). Mean Value at 75% Percentile is calculated 6,169,000 DNA copies/ml.

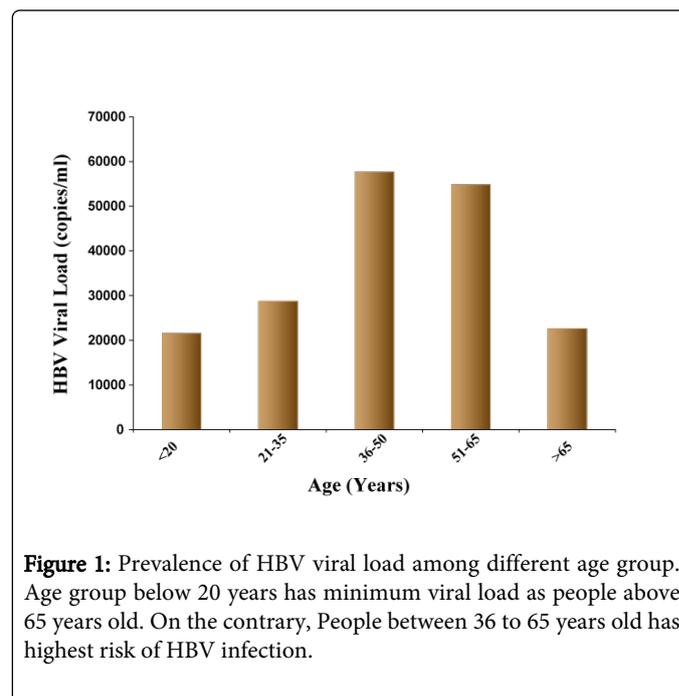


Figure 1: Prevalence of HBV viral load among different age group. Age group below 20 years has minimum viral load as people above 65 years old. On the contrary, People between 36 to 65 years old has highest risk of HBV infection.

Infection with HBV is a serious public health concern in Bangladesh like other developing countries. According to WHO [11], 2-5% of the general population in the Indian subcontinent is chronically infected with HBV. In this study viral load pattern was examined in the blood sample collected from various site of the country. In the wide age range of patients highest prevalence was observed between 36-50 years (29.47% of the population). Males have relatively higher prevalence than female. Country like Bangladesh males are the main work resource; they travel to the different part of the country for business or job purpose, drink unhygienic restaurant water or supply water from roadside small tea shop. Therefore males are

likely to be exposed to the contamination rather than females. The lower and upper limit of viral load found in the analysis for the Hepatitis B monitor assay is 128,300,000 and 301,900,000 DNA copies/ml respectively.

HBV viral load is important for clinical monitoring and treatment of individual with high HBV viral load as it is an independent predictor of liver cancer risk. Measurements of HBV viral load and genotype may help to define which male HBV carriers aged 30 years or older are at high risk for HCC [12]. Our study showed that HBV is more prevalent in male group and in the 36-50 age group. According to a Taiwanese study the risk of HCC is closely associated with HBV. They also showed that HBV DNA levels were persistently elevated in patients at highest risk of liver cancer [13]. Another study in the Gambia, West Africa showed that High-level HBV DNA (>10,000 copies/ml) was strongly associated with both HCC and cirrhosis (17- and 39-fold increased risk); even Lower level HBV viremia (200–10,000 copies/ml) confer a significant risk of HCC [14]. Though Antiviral treatment is the only way to reduce morbidity and mortality from chronic HBV infection but The implementation of mass immunization programs, which have been recommended by the World Health Organization since 1991, have dramatically decreased the incidence of HBV infection among infants, children, and adolescents in many countries [15].

However regular/routine screening of the HBV can alter that morbidity rate in Bangladesh and also the risk could be minimized if the National programme on immunization (NPI) scheme for HBV vaccination is implemented.

Conclusion

In conclusion, high viral load observed in the 21-35 and 36-50 age group where risk of HCC is involved. There were no study performed to compare the HBV viral load of metropolitan population with rural population. Also it would be nice if geographical distribution can be shown by further study. However, it is recommended that there should be health education for public and health care providers and also screening and vaccination of all special risk groups. National vaccination program and mass awareness about HBV infection can reduce the risk of viral infection and other associated disease.

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Conflict of Interest

There is no conflict of interest

References

1. Organization WH (2010) Introduction of hepatitis B vaccine into childhood immunization services. Department of Vaccines and Biologics, World Health Organization: Geneva 8-9.
2. Sherman M (2009) Risk of hepatocellular carcinoma in hepatitis B and prevention through treatment. *Cleve Clin J Med* 76 Suppl 3: S6-S9.
3. Bangladesh Lfo (2013) Incidence of Hepatitis-B infection in Bangladesh.
4. Khan M and Ahmad N (1996) Seroepidemiology of HBV and HCV in Bangladesh. *International Hepatology Communications* 5: 27-29.
5. Okwurawe A (2011) Experience with Hepatitis B viral load testing in Nigeria. *African Journal of Clinical and Experimental Microbiology* 12: 101-105.
6. Kao JH (2008) Diagnosis of hepatitis B virus infection through serological and virological markers. *Expert Rev Gastroenterol Hepatol* 2: 553-562.
7. Hasan KN, Rumi MA, Hasanat MA, Azam MG, Ahmed S, et al. (2002) chronic carriers of hepatitis B virus in Bangladesh: a comparative analysis of HBV-DNA, HBeAg/anti-HBe, and liver function tests. *Southeast Asian J Trop Med Public Health* 33: 110-117.
8. Sato S, Fujiyama S, Tanaka M, Yamasaki K, Kuramoto I, et al. (1994) Coinfection of hepatitis C virus in patients with chronic hepatitis B infection. *J Hepatol* 21: 159-166.
9. Kruskal-Wallis anova was performed using GraphPad Prism version 6.0 for Windows. GraphPad Software, San Diego California USA.
10. Population of Bangladesh.
11. Hepatitis B (2014) in Fact Sheet 204.
12. Yu MW, Yeh SH, Chen PJ, Liaw YF, Lin CL, et al. (2005) Hepatitis B virus genotype and DNA level and hepatocellular carcinoma: a prospective study in men. *J Natl Cancer Inst* 97: 265-272.
13. Yu MW, Chang HC, Chen PJ, Liu CJ, Liaw YF, et al. (2002) Increased risk for hepatitis B-related liver cirrhosis in relatives of patients with hepatocellular carcinoma in northern Taiwan. *Int J Epidemiol* 31: 1008-1015.
14. Mendy ME, Welzel T, Lesi OA, Hainaut P, Hall AJ, et al. (2010) Hepatitis B viral load and risk for liver cirrhosis and hepatocellular carcinoma in The Gambia, West Africa. *J Viral Hepat* 17: 115-122.
15. WHO (2000) Hepatitis B. World Health Organization Fact Sheet 204.