The molecular epidemiology and Biochemical profiling of HBV infection in the North-eastern part of India

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Abstract:

In India also HBV is considered a major public health problem. Among the general population of India the prevalence of HBV was recorded as a range from 2%-8% and found that India is in the intermediate HBV endemic zone. The population in the North east India is one of the most indigenous tribes with different ethnicity and has a higher incidence of drug abusers as well as HIV infection. The present case control prospective study has been undertaken to explore the viral and host genetic risk factors for the hepatitis B infection in Assam using different diagnostic tools. The present study was designed to express on many important aspects of molecular epidemiology of HBV which are very much important for identifying the population, at the risk of acquiring HBV and the developing of severity of liver disease and also propose about the transmission through different modes. From the overall samples collected 5.12% were found as the HBV positive cases and were further studied for the above objective. The male population (74%) are more prone to HBV related liver disease than women (26%). The biochemical parameters and risk factors were studied to find the severity. Also to study the seroprevalence of HBV infection among the healthy blood donors and Healthcare workers among the North eastern population, samples were first done with HBsAg ELISA and then confirmed with PCR. Out of the 751 HBV positive samples (5.12%) it was found that 4.5% were positive for blood donor and 0.6% were positive for health care workers. The blood donors with mean age 28.12±4.30 with Male: Female ratio of 31:3 and found as significantly associated with the disease (p < 0.05).

Introduction

The hepatitis B virus (HBV) is also considered as a serious public health problem globally and it is also the major cause of chronic hepatitis, cirrhosis, and hepatocellular carcinoma (HCC). It was calculable that close to two billion individuals has medical science proof of past or gift HBV infection. There are more than 350 million people who are infected with this chronic carriers of the HBV (1). Also due to the high HBV virus related morbidity and mortality, the worldwide illness burden of haemoprotein is concerning as very important (2). The prevalence rate of the chronic HBV infection varies greatly in several a part of the world. This prevalence of chronic HBV infection which is worldwide could be categorized as very high, intermediate and low endimicity (3). The incident rate of the HBV infection is different throughout the regions of the world and which is also depend on the different factors(4). The Hepatitis B infection is extremely endemic in the developing regions with giant population like South East Asia, China, geographic region and therefore the Amazon Basin, wherever a minimum of 8 May 1945 of the population are HBV chronic carrier. Most infections occur during the infancy itself which may also transmit from their mother during pregnancy, which is very much useful to study. Since most of the infections in the kids are without any symptoms, there is a very little proof of the acute illness associated with HBV, however the rates of chronic disease and carcinoma in adults are very high (5).

Hepatitis B is moderately endemic partly of Japanese and Southern Europe, the centre East, Japan, and a part of South America. There are 10–60% of the population have been infected with this HBV and 2-7% are considered as chronic carriers. Acute illness associated with HBV is common in these areas as a result of several infections occur in adolescents and adults; but, the high rates of chronic infection are maintained largely by infections occurring in infants and children(6). The infection rate of HBV is generally low in most of the developed areas, like North America, Northern and Western Europe and Australia, which is considered as low endemic areas. HBV infections are rate as around 5–7% of the population, and only 0.5–2% of the population are considered as chronic carriers in these regions (7). In these areas, generally most of the HBV infections occurs in the adolescents and young adults in comparatively well-defined risky teams, as well as injection user, homosexual males, health care staff, patients UN agency need regular blood transfusion or haemodialysis.

Transmission of Hepatitis:

HBV is unfolding through contact with infected body fluids and also the solely natural host is human. The fluid connective tissue or blood is generally considered as the most important mode of transmission for various diseases, but other body fluids have also been implicated, including semen and saliva (8). Currently, 3 modes of HBV transmission are recognized: perinatal, sexual and parenteral/percutaneous transmission. There is no reliable evidence that airborne infections occur and faces are not a source of infection. HBV isn't transmitted by contaminated food or water, insects or other vectors. Transmission of HBV from carrier mothers to their babies will occur throughout the perinatal amount, and appears to be the most important factor in determining the prevalence of the infection in high endemicity areas, notably in China and geographic region. Sexual transmission of hepatitis B may be a major supply of infection altogether areas of the globe, especially in the low endemic areas, such as North America. Hepatitis B is taken into account to be a sexually transmitted sickness (STD). The risk of chronicity is low (less than 5%) for transmission through sexual contact, intravenous drug use, acupuncture, and transfusion(9). Individuals in danger for these transmission modes sometimes acquire HBV infection throughout adolescence or adulthood while not immune tolerance. Instead, the sickness progresses on to the immune clearance section and is of short length, that most likely accounts for prime spontaneous recovery.

Material and methods:

The North-eastern region, where the study was conducted, is a less developed region of India in terms of economic, social, and health indices. Insufficient health services and lack of public awareness of health-related issues have increased the prevalence rate of diseases, particularly communicable diseases. This population based case-control study was carried out by Bioengineering and technology Department, Gauhati University, Department of Gastroenterology, Gauhati Medical College, Guwahati, Assam; Department of Medicine, Regional Institute of Medical Sciences Regional Medical College,Imphal; Deptartment of Medicine, NEIGRIHMS, Shillong, Meghalaya; General Hospital, Naharlagun, Medical Adviser to the Hon'ble Chief Minister of Sikkim, STNM Hospital, Gangtok, Sikkim; Dept of Medicine, Agartala Govt. Medical College, Agartala, Tripura; Directorate of Medical Education, General Hospital, Naharlagun, Arunachal Pradesh. Arunachal Pradesh.

The study was carried out from Nov 2012 to May 2015. A total of 7130(48.40%) liver disease cases were collected included HBV,HCV,HAV,HEV, Aloholic liver disease, Nonalcoholic liver disease and Blood donors (48.05%) and health care worker (03.53%).

This cases were based on clinical and laboratory parameters of both Male ans Felmale with same number of control samples. Cases were pathologically and/or cytologically confirmed subjects of both sexes (male and female) and for HCC cases with serum alpha-HBV fetoprotein level greater than 400 ng/ml combined with at least one positive image on angiography, sonography, liver scan, and/or computed tomography scan were included. After obtaining written informed consent, trained social investigators collected the data from the eligible subjects by face-to-face interview using a pre-coded, closed ended questionnaire. The sociodemographic characteristics included participants' age, sex, educational status, occupational status and location of residence etc. The questionnaire included ranges of information including sociodemographic characteristics of participants, history of alcohol drinking habits, type of alcoholic beverages, dietary habits. Alcohol drinking habits were categorized into three categories, viz., non-drinker, former drinker, and current drinker. Subjects who were reported that they were regularly drink alcoholic beverage during index year were defined as current drinker, those who reported that they had stopped drinking the year before the index year or in the former were defined as former drinker, and those who reported that they never drank before were defined as non-drinker or abstainers. Diets and the dietary history of patients and controls were recorded based on 6 months of recall. Food frequency questions that contain details of dietary practices were included in the questionnaire. The frequency of consumption was classified as never eaten, occasionally, once a week, 2–4 times a week, and more than 4 times a week.

Clinical data analysis by Virological methods:

All those patients who were clinically diagnosed as Acute Viral Hepatitis (AVH), Fulminant Hepatic Failure (FHF), chronic active hepatitis (CAH), Cirrhosis, Hepatocellular Carcinoma (HCC) with HBsAg positive will be included in the study. Voluntary blood donors and the Health care workers (HCW) will be included in this study. Professional blood donors, high risk group like IV drug abusers were excluded from the study. The HBV suspected patient's blood sample were collected in both EDTA and Clot activator vials. Serum/Plasma was

separated by centrifugation method. Hepatitis B surface Antigen, Anti Hepatitis B core IgM, Anti HCV, Hepatitis A core IgM, Hepatitis E core IgM were done using specific kit. HBsAg positive cases were subjected to HBV DNA extraction and further Polymerase Chain Reaction with specific Primers. All the liver disease patients were examined by qualified Gastroenterologists to determine the different symptoms of liver disease which includes nausea, vomiting, loss of appetite, jaundice, dark urine and pale stool. Based on HBsAg positive and PCR positive status were categorized as chronic carriers and were sub-categorized as chronic hepatitis B or cirrhosis after co-relating with the clinical details; else they were categorized as acute HBV cases. The HBeAg status was further screened by ELISA using commercially available kits.

Statistical analysis:

Descriptive statistics was performed to describe different characteristics of the HBV mediated liver disease. Univariate and multiple logistic regression was performed to assess the factors associated with HBV. The multiple logistic regression analysis was performed to calculate adjusted odds ratio (Adj. OR) and corresponding 95% CI for determining independent associations between the factors and HBV status after adjusting for potential factors. Only variables that were found to be significantly associated with HBV status in univariate analysis at 5% level of significance were included in multiple regression model. This study was approved by institutional ethical committee of Gauuhati University and Indian Council of Medical Research, New Delhi.

Results:

The burden of the viral hepatitis B infection in North-eastern states of India is not well characterised till date. To fulfil the objectives of the study a surveillance was conducted across different states of North-eastern part of India. To study the epidemic-prone diseases, information on the outbreaks of all forms of viral hepatitis, including A, B, C and E were collected. The data were collected from the Department of Gastroenterology, Gauhati Medical College, Guwahati, Assam; Department of Medicine, Regional Institute of Medical Sciences Regional Medical College, Imphal; Deptartment of Medicine, NEIGRIHMS, Shillong, Meghalaya; STNM Hospital, Gangtok, Sikkim; Dept of Medicine, Agartala Govt. Medical College, Agartala, Tripura; Directorate of Medical Education, General Hospital., Naharlagun, Arunachal Pradesh. A total number of 14,729 samples with different types of liver diseases were screened for hepatitis B virus infection from all the centres, these screened data include 7130 (48.40%) different types of liver disease cases (HAV,HCV,HBV,HEV),7077 (48.05%) prospective blood donors and 522 (3.55%) health care workers and are depicted in table 1 and figure1. To study the objectives of polymorphism and expression a total number of 294 control samples were included. From different parts of North-eastern states different liver disease cases were screened and further they were studied for HBV DNA positivity.

Table1: Different categories of sample screened

STUDY CATEGORY	NUMBER OF SCREENED	SAMPLE
LIVER DISEASE	7130(48.40%)	
BLOOD DONOR	7077(48.05%)	
HEALTHCARE WORKER	522(3.53%)	

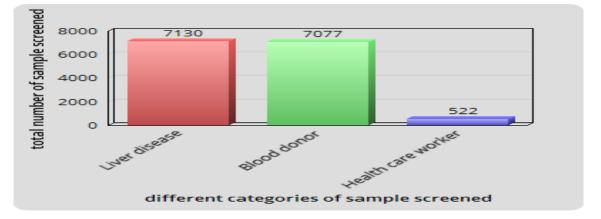


Figure 1: Showing different categories of samples screened from all the States.

Amplification of HBV viral samples:

All the screened samples were further amplified by nested PCR primers and found around 5.12% were HBV DNA positive and amplification of PCR result has been shown in figure2. In figure 3 graphically the comparison was done among the overall liver disease cases and HBV DNA positive cases. All the screened samples were further amplified by nested PCR primers and found around 5.12% were HBV DNA positive (Figure3; Table2).



Figure 2: HBV DNA detection by Nested PCR Method; Lane 1: 100 bp ladder; Lane 2,4,5,6,7: HBV Positive samples; Lane3: HBV DNA Negative sample.

STUDY GROUPS	Overall Cases (n=14,729)	HBV DNA positive
		(n=751) (5.12%)
AVH	781(5.3%)	221 (29.4%)
FHF	25(0.16%)	03 (0.3%)
САН	3682(24.74%)	308 (41%)
CIRRHOSIS	2543(17.26%)	142 (19%)
HCC	162(1%)	162(1%)
BLOOD DONOR	7077(48%)	34 (4.5%)
HEALTH CARE WORKER	522(3.54%)	05 (0.6%)

Table2: Distribution of HBV DNA positivity from the overall study samples.

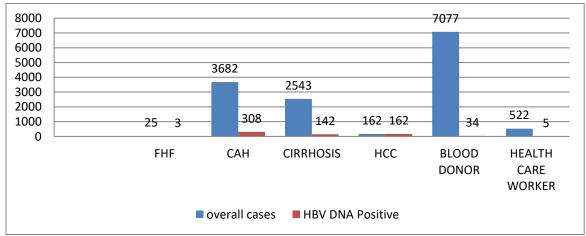


Figure 3: Comparision of overall liver disease cases and HBV positive cases

All the HBsAg positive cases were studied with demographical, biochemical, clinical and serological profile and compared with HBV DNA positive cases expressed in tabl3 and figure4. The overall distribution of different HBV cases has been shown state wise. The biochemical parameters like ALT, AST, Total Bilirubin, Direct Bilirubin, Haemoglobin, Prothrombin time, serum creatining, viral load, Na+, K+ were recorded. The p value was calculated by using unpaired t test. The study groups mean \pm S.D values were compared with the control samples and were represented in table 4. All the biochemical parameters were found as statistically significant and all are highly associated with the disease (p<0.05).

Table3 : Overall collected samples and the HBV positive samples in all the States.

States of NE	AVH	FHF	CAH	CIRRHOSIS	HCC	BLOOD	HEALTH
						DONOR	CARE
	(n=221)	(n=3)	(n=308)	(n=142)	(n=38)	(n=34)	WORKER
							(n=5)
ASSAM	71(32.1%)	0	79(25.6%)	52(37%)	12(31.6%)	8(23.5%)	0

MEGHALAYA	2(0.9%)	1(33.3%)	68(22%)	34(24%)	3(7.9%)	10(29.4%)	0
TRIPURA	40(18%)	0	45(14.6%)	20(14%)	8(21%)	14(41.1%)	5(100%)
ARUNACHAL	26(11.8%)	0	34(11%)	21(14.8%)	5(13.1%)	1(2.9%)	0
SIKKIM	47(21.3%)	1(33.3%)	39(12.6%)	6(4.2%)	2(5.2%)	1(2.9%)	0
MANIPUR	35(15.9%)	1(33.3%)	43(14%)	9(6.4%)	8(21%)	0	0

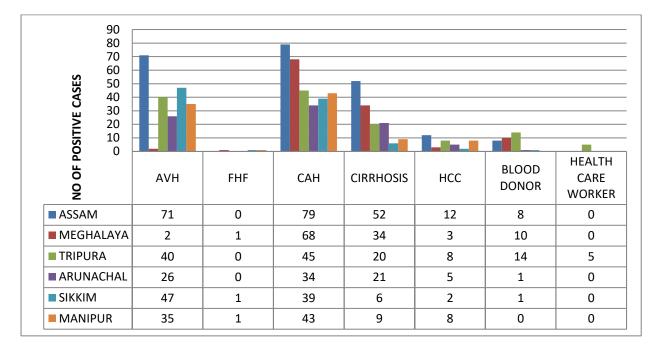


Figure4: Distribution of different stages of HBV in different study centres

Table4: Unpaired t test results for comparing the different study groups with their biochemical profiles.

	Male (n=556); 74%	Female(n=1	P value
		95),26%	
Age:<20yr	23	2	0.2
21-30	162	119	
31-40	274	45	
41-50	68	21	
51-60	21	7	
>60	8	1	
ALT	238	89	0.04
<40 U/L	318	106	
>40 U/L			
AST	291	78	0.016
<30U/L	265	117	
>30U/L			
Viral load (IU/Ml)	31	6	0.04
<1000 IU/Ml	114	30	
1000-10000 IU/MI	189	49	
10000-100000 IU/MI	142	86	
100000-1000000 IU/MI	80	24	
>1000000 IU/Ml			

Hb(%)	386	115	0.2
>12	170	80	
<12			
TotalBilirubin (mg/dl)	245	78	0.04
0.03-1.9	311	117	
>1.9			
DirectBilirubin (mg/dl)	257	68	0.03
<0.3	299	127	
>0.3			
Prothrombintime (INR)	95	48	0.13
<0.5	310	79	
0.5-1.5	151	68	
>1.5			
Serumcreatinine (mg%)	286	87	<0.05
<1.1-1.3	270	108	
>1.1-1.3			
Na(mEq/L)	74	39	0.16
<135	169	69	
135-145	313	87	
>145			
K+(mEq/L)	55	38	0.15
<3.7	219	96	
3.7-5.2	282	61	
>5.2			
HBeAg status	231	166	0.09
Present (n=397)	198	159	
Absent (n=357)			

Additionally HBe antigen is used as an indicator of virological response when treating patients with chronic hepatitis B. The HBeAg positive/negative was studied among all the study groups and it was found that HBeAg is not highly associated with a specific phase of HBV DNA positive liver disease (p>0.05) presented in the table5.

Table5: HBeAg profile among the different stages of HBV DNA positive liver disease.

	HBeAg positive (N=397);52.87%	HBeAgnegative(N =354);47.1%	P value
AVH(n=221)	103(25.96%)	118(33.33%)	0.46
CLD(N=308)	210(52.90%)	98(27.70%)	0.07
CIRRHOSI (N=142)	52(13%)	90(25.42%)	0.77
HCC(N=38)	20(5.04%)	18(5.08%)	0.37
FHF(N=3)	1(0.25%)	18(5.08%)	0.67
BLOOD DONOR (N=34)	10(2.6%)	24(6.78%)	0.92

HEALTH	1(0.25%)	4(1.13%)	Ref
CARE (N=5)			

Discussion:

To the best of our knowledge, this was the first study to assess the molecular and epidemiology study of HBV disease in the Northeast India. Different factors associated with HBV in different part of the Northeast India. We examined the relationship of different biochemical factors in different types of liver diseases associated with HBV. The present study found the incident rate of HBV among found illiterate and ever drinkers (current and former drinkers) are independently associated with increased risk of HCC. Illiteracy limits a person's ability to comprehend verbal and written health care information which adversely affect the awareness about healthy life style and precaution measure of various health issues. In the North-eastern region of India, preparation and consumption of traditional homemade alcohol is a social tradition of most of the tribe. Possibly folks with low accomplishment level ineffectual to spot the chance of excess consumption of any alcohol liquid, which make them vulnerable to liver disease. It has also been documented that one of the main cause of chronic liver disease and HCC is alcohol consumption chronic liver disease and HCC is alcohol consumption (10,11). Better approaches of large scale of HBV vaccination, identification and treatment of chronic HBV, alcohol reducing campaigns in high risk area are needed to prevent this important form of cancer.

Conclusion:

In conclusion, the findings of this study is first ever biochemical factors among HBV and Blood donor and health care workers patients in this increasing liver cancer incidence states of India. This study also demonstrates a protective relationship of HBV with consumption of different food habbits. Prevention of primary infection by vaccination is a crucial strategy to decrease the chance of chronic HBV infection and its later complications. The burden of adverse outcomes related to hepatitis B on individuals and communities, particularly in high-prevalence populations, is increasingly recognized. Although ensuring the high coverage of infant vaccination will have a profound impact on this burden in coming decades, attention must be given to comprehensive policy responses now. Understanding the epidemiology of HBV infection will enable evidence-based and cost-effective public health and clinical interventions within countries and at the global level.

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Conflict of interest

None.

References:

- 1. Hepatitis B: World Health Organization Fact Sheet 204. World Health Organization; [2000].
- 2. 1.Hou, J., Liu, Z., & Gu, F. (2005). Epidemiology and Prevention of Hepatitis B Virus Infection. International journal of medical sciences, 2(1), 50-57.
- 3. Hou, J., Liu, Z., & Gu, F. (2005). Epidemiology and Prevention of Hepatitis B Virus Infection. International journal of medical sciences, 2(1), 50-57.
- 4. Margolis HS, Alter MJ, Hadler SC. Hepatitis B: evolving epidemiology and implications for control. Semin Liver Dis. 1991;11:84–92.
- 5. Alter M. Epidemiology of hepatitis B in Europe and worldwide. J Hepatol. 2003;39:S64–S69
- 6. Toukan A. Strategy for the control of hepatitis B virus infection in the Middle East and North Africa. Vaccine. 1990;8(suppl):S117–S121.
- 7. McQuillan GM, Townsend TR, Fields HA, Carroll M, Leahy M, Polk BF. Seroepidemiology of hepatitis B virus infection in the United States. Am J Med. 1989;87(suppl 3A):5S–10
- 8. Scott RM, Snitbhan R, Bancroft WH, Alter HJ, Tingpalapong M. Experimental transmission of hepatitis B virus by semen and saliva. J Infect Dis. 1980;142:67–71.
- 9. Hyams KC. Risks of chronicity following acute hepatitis B virus infection: a review. Clin Infect Dis. 1995;20:992–1000.
- 10. S. Sidharthan, S. Kottilil, Mechanisms of alcohol-induced hepatocellular carcinoma, Hepatol. Int. 8 (2) (2014) 452.
- 11. G. Testino, S. Leone, P. Borro, Alcohol and hepatocellular carcinoma: a review and a point of view, World J. Gastroenterol. 20 (43) (2014) 15943.