



Determinants of Hepatitis B Virus (HBV) Infection Among University Students in Central Bangladesh

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Abstract

This study aimed to determine the seroprevalence and determinants of hepatitis B virus (HBV) infection among university students in Bangladesh. This cross-sectional study was conducted among 614 students from five universities in central Bangladesh. Data were collected on demographic information, immunization history, medical and blood transfusion history through the face-to-face interview. Blood samples were collected and screened for anti-HBsAg using ELISA, HBsAg Rapid Test-cassette, and immune chromatographic test. The overall seroprevalence of HBV infection was 5.0%, and vaccination coverage was 19.2% among the participants. Students having a history of surgery (OR 11.004, 95% CI 3.211–37.707), blood transfusion (OR 5.651, 95% CI 0.965–33.068), being married (OR 4.776, 95% CI 1.508–15.127), and not being vaccinated (OR 9.825, 95% CI 1.130–85.367) were at higher risk of being infected by HBV. This study showed the endemicity of HBV infection among the Bangladeshi population. Marriage, surgical or blood transfusion history, not being vaccinated were the determinants of HBV infection within the study population. Public health initiatives for preventing HBV infection at the university levels should be envisaged.

Keywords Hepatitis B virus · Infection · Risk factors · Students · Bangladesh

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Introduction

Hepatitis B virus (HBV) infection is one of the most common and severe infectious diseases responsible for significant morbidity and mortality [1]. Worldwide, around 257 million people have been chronically infected with HBV, and 887,000 deaths have resulted from HBV infection, including cirrhosis and hepatocellular carcinoma in 2015 [2]. Almost half of all cases of hepatocellular carcinoma and one-third of all cases of cirrhosis are accountable for HBV infections [3, 4]. A high prevalence (HBV infection affects more than 8.0% of the population) of chronic HBV infection has been found in the Asia Pacific and sub-Saharan African regions of the world [5]. HBV seroprevalence is classified as intermediate (2–7%) in North Africa and the Middle East, Eastern and Southern Europe, Latin America, and South Asia [5]. In South-West Asia, the Russian Federation, Eastern, and Southern Europe, most of central and South America, the seroprevalence of chronic HBV was > 2 to < 8%. The seroprevalence of chronic HBV infection was low (< 2% of the population HBsAg-positive) in Northern and Western Europe, Australia, New Zealand, and North America [6]. Almost three-quarters of HBV infection was found in Asia, the Middle East, and Africa [7]. HBV infection has been identified as an emerging future health problem in many developing countries in Asia [8–11]. Substantial morbidity and mortality occur due to severe infection and chronic sequelae, including chronic hepatitis, liver cirrhosis, liver cancer, and hepatocellular carcinoma [12, 13]. Among the 1.34 million deaths caused by viral hepatitis, most of the deaths were due to chronic liver cirrhosis (720,000 deaths) and hepatocellular carcinoma (47,000 deaths) in 2015 [14].

In the Indian sub-continent, the chronic HBV infection carriage rate ranged within 2–5%, which is recognized as intermediate endemicity of HBV infection [15, 16]. The HBsAg prevalence of Bangladesh is within the range of 2–7%, which was reported by previous studies among the different selective populations of this country [17]. In developed countries, HBV infection is mostly infected people through injecting the drug, sexual activity, or occupational exposure. Other reasons for infections can be included in developing countries, such as household contact, vertical transmission via hemodialysis, transmission from a surgeon [18], and the receipt of blood or bloody fluids or organs [19]. The sustainable development goals (SDGs) have been set to target 3.3.4 to combat HBV incidence per 100,000 populations [20]. Younger people are at more risk of being infected by HBV [21]. However, most previous studies were conducted among selected populations such as drug addicts, blood donors, hospitalized patients, and commercial sex workers [22–26]. In Bangladesh, there is a scarcity of information about HBV infection among young individuals. This

study aimed at assessing the seroprevalence of HBV infection and determinants among university students in Central Bangladesh.

Methods

Study Design and Population

This study was planned as a descriptive cross-sectional study among students from five Bangladeshi universities (Jahangirnagar University, Daffodil International University, City University, Manarat International University, and Eastern University, Dhaka). Pre-advertising was done using a poster, brochure, and university Facebook page/group after receiving approval from each university. At each university's medical center, a data and sample collection area was developed. From 10 a.m. to 3 p.m., one research assistant (data collector) and one nurse were stationed in each corner to collect data and blood samples. The information was gathered between July and November 2018. Students who expressed an interest were freely questioned, and a blood sample was taken. The sample was taken to the Thalassemia Hospital and Institute's laboratory and analyzed daily. The report was sent to the pupils at no cost to them.

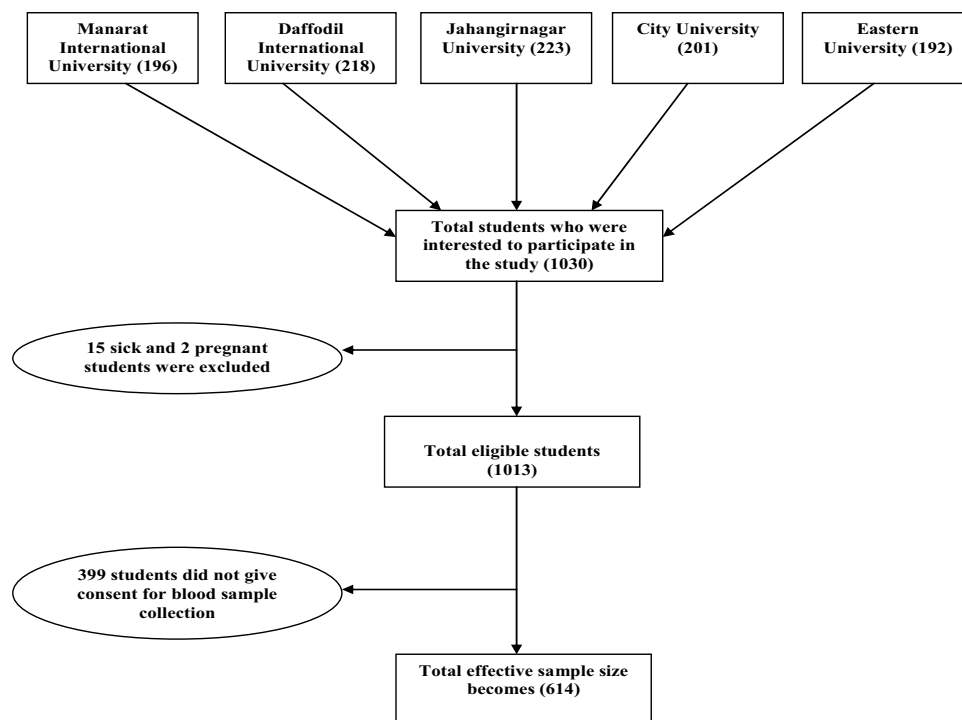
Sample Size and Data Collection

A total of 1030 students from five universities expressed an interest in participating in this study. 424 male and 207 female students approved blood testing. Of them, 15 were not included in the study because they were unwell, and two were pregnant. In the end, 416 students were removed, and 614 students were enrolled in the study out of 1030. A structured questionnaire was utilized to capture their socio-demographic information using a face-to-face interview technique.

Blood Sample Collection and Lab Test

A nurse took blood samples from the middle cubital vein after taking adequate aseptic precautions and storing them in an EDTA tube on the spot. The blood samples were subsequently delivered to the Thalassemia Hospital and Institute's laboratory. To avoid errors and biases, the samples were labeled differently. To reduce inaccuracies, data about the responders were assigned the same credential number as their blood samples. The test tubes containing blood samples were centrifuged at 3000 rotations per minute for 8–10 min at room temperature, and blood plasma was extracted. Plasma samples were tested for the presence of HBsAg using a commercial test kit (chromatographic immunoassay), the HBsAg Rapid Test-cassette

Fig. 1 Selection of study sample



(CTK, bioscience, Inc. USA). The test was repeated once if any of the test 1 results were positive. A medical technologist performed the test, which a physician later validated. The ELISA test was used to confirm positive instances (Fig. 1).

Data Analysis

The data were analyzed using Stata version 14.2 SE (Stata Corp 2015; Statistical Software: Release 14. College Station, TX: Stata Corp LP). The age of the students was changed from a continuous to a categorical variable. Using descriptive statistics, the frequency and proportion of socio-demographic and background variables, HBV infection, immunization status, and clinical exposure of university students were investigated. The correlation between HBV infection and demographic factors was investigated using chi-square tests. Multiple logistic regressions were used to examine the relationships between marital status and HBV infection, with marital status as an explanatory variable and HBV infection status as an outcome variable. To find the determinants of HBV infection, researchers used multiple logistic regressions with odds ratios (OR). We used HBV infection as an outcome variable in numerous logistic regressions, age, level of study, gender, marital status, division, history of surgery, HBV vaccination status, and history of blood transfusion as explanatory variables. Multiple logistic regression models were used to control the influence of all confounders when they were connected with the key exposure variable and the result of interest in this article. A covariate was considered a confounder if it was significantly

related to the exposure variable and the result of interest in the binary logistic regression.

Results

Socio-Demographic, Infection and Vaccination Status of the Students

A total of 614 students were tested for HBV infection, with 92.7% of those tested being under the age of 25. 79.8% of the study participants were undergraduate students, the majority (66.9%) were men, and 93.5% of the respondents were single. Only 19.2% of the 614 students tested positive for HBsAg, and only 31 (5.0%) were HBV vaccinated. Only 2.0% of all responders had ever received a blood transfusion, and only 3.9 percent had ever had surgery (Table 1).

Seroprevalence of Hepatitis B Infection by Demographic Characteristics

The prevalence of HBV infection was substantially greater among respondents aged 25 years or less ($P = 0.008$), and there was also a significant link between HBV infection and level of study ($P < 0.05$); however, undergraduates reported a higher proportion of HBV infection. Gender was not shown to be significantly linked ($P = 0.096$) with HBV infection in this study, while male students had a greater frequency of HBV infection (6.1%) than female

Table 1 Socio-demographic, infection and vaccination status of the students (N = 614)

Variables	Frequency	Percentage
Age (years)		
≤ 25	569	92.7
> 25	45	7.3
Level of study		
Undergraduate	490	79.8
Graduate	124	20.2
Gender		
Male	411	66.9
Female	203	33.1
Marital status		
Married	40	6.5
Unmarried	574	93.5
Hepatitis B virus infection		
Yes	31	5.0
No	583	95.0
Hepatitis B virus vaccination status		
Yes	118	19.2
No	496	80.8
History of blood transfusion		
Yes	12	2.0
No	602	98.0
History of surgery		
Yes	24	3.9
No	590	96.1

students (3%), ($P = 0.010$). The prevalence of married students (22.5%) was higher than that of unmarried students (3.8%) ($P = 0.001$). Respondents who had previously undergone surgery had a higher prevalence (37.5%) than those who had not (3.7%) ($P = 0.001$). The frequency of HBV infection was higher among students (50.0%) who had previously received blood transfusions than among others (4.2%) ($P = 0.001$) (Table 2).

Determinants for HBV Infection Among University Students

A binomial logistic regression analysis was done to identify the determinants of HBV infection. HBV infection status was the dependent variable, while age, education level, gender, marital status, surgery history, hepatitis vaccine status, and blood transfusion history were the independent variables. The risk of HBV infection was greater (OR 1.293, 95% CI 0.335–4.990) among students aged 25 and under. There was no significant relationship between the students' ages and the risk of HBV infection. The level of

Table 2 Distribution of hepatitis infection by demographic characteristics (N = 614)

Characteristics	N	Hepatitis		P value
		N	%	
Age (years)				
≤ 25	569	25	4.4	0.008
> 25	45	6	13.3	
Level of study				
Undergraduate	490	20	4.1	0.030
Graduate	124	11	8.9	
Gender				
Male	411	25	6.1	0.096
Female	203	6	3.0	
Marital status				
Married	40	9	22.5	0.000
Unmarried	574	22	3.8	
History of surgery				
Yes	24	9	37.5	0.000
No	590	22	3.7	
Hepatitis B virus vaccination status				
Yes	118	1	0.8	0.020
No	496	30	6.0	
History of blood transfusion				
Yes	12	6	50.0	0.000
No	602	25	4.2	

study of the participants did not affect the likelihood of having HBV infection (OR 1.691, 95% CI 0.594–4.816). Married students had a greater risk of HBV infection (OR 4.776, 95% CI 1.508–15.127) than unmarried students. HBV infection was 11 times more likely among students who had previously undergone surgery (OR 11.004, 95% CI 3.211–37.707) than students who had never undergone surgery. Students who were not immunized (OR 9.825, 95% CI 1.130–85.367) had an almost ninefold higher chance of contracting HBV than those who were. HBV infection was linked to a history of blood transfusion (OR 5.651, 95% CI 0.965–33.068), with individuals who had blood transfusions being more than 5 times more likely to contract the virus (Table 3).

Discussion

To the best of our knowledge, this is the first study to investigate the seroprevalence of HBV infection, vaccination status, and HBV infection determinants among university students in Bangladesh. In this study, the seroprevalence of HBV infection was around 5.0%, which is similar to 4.3% in a similar survey conducted in Bangladesh [27]. The findings

Table 3 Determinants for HBV infection among university students (N=614)

Factors	Crude OR	CI (95%)		P value	Adjusted OR	CI (95%)		P value
		Lower	Upper			Lower	Upper	
Age (years)								
≤25	Ref							
>25	3.347	1.296	8.642	0.013	1.293	0.335	4.990	0.709
Level of study								
Undergraduate	Ref							
Graduate	2.287	1.065	4.910	0.034	1.691	0.594	4.816	0.325
Marital status								
Married	7.284	3.095	17.141	0.000	4.776	1.508	15.127	0.008
Unmarried	Ref							
History of surgery								
Yes	15.490	6.113	39.249	0.000	11.004	3.211	37.707	0.000
No	Ref							
HBV vaccination status								
Yes	Ref							
No	7.532	1.016	55.802	0.04	9.825	1.130	85.367	0.038
History of blood transfusion								
Yes	23.08	6.949	76.653	0.000	5.651	0.965	33.068	0.055
No	Ref							

of our investigation are validated by Ashraf et al., [28], who found a 6.5% prevalence rate of HBsAg. The prevalence of HBsAg in our study is within 2–7%, which has been documented in earlier studies among Bangladesh's various selective populations [17]. In 2017, the Directorate General of Health Services (DGHS) of Bangladesh's Ministry of Health and Family Welfare recorded 546 cases of HBV infection per 100,000 people [29]. Marital status was one of the most critical determinants for HBsAg positive among university students. Married students were more likely than unmarried pupils to test positive for HBsAg.

HBV infection can be spread through sexual activity, and unsafe sexual practices can lead to HBsAg infection [30–34]. One probable explanation is that married people are more likely to become infected with HBsAg through sexual activity. Pathogen transmission from the hospital environment to the population, particularly HBV infection, is a significant source of pathogen transmission from the hospital environment to the community, especially in developing countries like Bangladesh. The majority of HBV infection transmissions happen during invasive or obstetric procedures [35]. This study indicated that persons with any surgical history were at risk for HBV infection, consistent with prior findings. During surgery, HBV infection can be spread in three ways: from the surgeon to the patient, from contaminated surgical tools to the patient, and from an HBsAg positive patient to another in the same hospital room [36–38]. To prevent all healthcare-related transmissions of HBV infection, a comprehensive approach is required, which includes

ensuring hygiene and cleanliness for health workers, hospital rooms and medical equipment to prevent nosocomial infections, administering HBV vaccination to health workers, and taking measures to reduce blood exposure [39–41]. Another key risk factor for HBV infection, according to this study, is a history of blood transfusion. Recent research has proven that blood transfusion is one of the critical risk factors exacerbated by blood donors' occult HBV infection (OBI) [16, 42]. The previous investigation found that the seroprevalence of HBsAg was high among commercial blood donors (18–29%) in Bangladesh [22, 43]. This higher prevalence occurs partly due to previous contamination of blood transfusion.

The presence of HBV immunization was a significant factor in HBsAg infection. Immunization against HBV can prevent mild to chronic infection, as well as complications such as cirrhosis and hepatocellular cancer [35]. The introduction of HBV vaccination resulted in a considerable decrease in HBV seroprevalence and HBV-related illnesses [35, 44, 45]. The government of Bangladesh began including the HBV vaccine in the expanded program of immunization (EPI) in 2003 to meet the World Health Organization's objective of eradicating HBV infection by 2030 [46, 47]. The immunization coverage program, however, is just for infants. It is highly anticipated that a program for HBV infection vaccination coverage among young adults will be implemented [48]. Unlike other studies, ours found gender and location to be unimportant [42, 49].

There were a few flaws in this study as well. For starters, due to a lack of time and funds, only ELISA tests were used

to detect positive instances of HBV infection using the ICT method. Second, because medical records were not used to verify vaccination status for HBV infection, self-reported vaccination status could be underreported due to recollection bias. Finally, the researchers were unable to determine the precise path of HBV infection transmission among university students. Fourth, we could not determine whether the risk factors and behaviors were adopted after or before the entrance to the institution. Finally, we included the key risk factors for HBV infection; nevertheless, other factors may have been overlooked in our analysis.

Conclusion

In this study, university students in Bangladesh had a high seroprevalence of HBV infection. HBV infection was more likely in married students, had a surgical history, had a history of blood transfusion, and had not been vaccinated. To prevent HBV infection among married students, sexual education and awareness should be widely disseminated. To prevent HBV infection resulting from a blood transfusion or surgery, healthcare facilities should maintain good cleanliness and sanitary procedures. Young people should be included in the vaccination program. To prevent HBV infection among university students in Bangladesh, public health activities are required.

Declarations

Ethical Approval The Bangladesh Institute of Allergy and Clinical Immunology's ethical review committee approved the study (IACIB). Each participant signed a written informed consent form. They were informed about the study's nature and goal and the fact that their information would be kept private. In addition, all volunteers received free laboratory tests, interviews, and results from notification. Each participant's test results were sealed, encased, and personally delivered. As a thank you for taking part in the study, each participant received a modest gift.

References

1. Franco, E., Meleleo, C., Serino, L., Sorbara, D., & Zaratti, L. (2012). Hepatitis A: Epidemiology and prevention in developing countries. *World Journal of Hepatology*, 4(3), 68–73.
2. World Health Organization (WHO). (2020). Hepatitis B. Accessed August 20, 2020, from <https://www.who.int/news-room/factsheets/detail/hepatitis-b>
3. Parkin, D. M. (2006). The global health burden of infection-associated cancers in the year 2002. *International Journal of Cancer*, 118(12), 3030–3044.
4. Ghouri, Y. A., Mian, I., & Rowe, J. H. (2017). Review of hepatocellular carcinoma: Epidemiology, etiology, and carcinogenesis. *Journal of Carcinogenesis*. https://doi.org/10.4103/jcar.JCar_9_16
5. MacLachlan, J. H., & Cowie, B. C. (2015). Hepatitis B virus epidemiology. *Cold Spring Harbor Perspectives in Medicine*, 5(5), a021410. <https://doi.org/10.1101/cshperspect.a021410>
6. World Health Organization (WHO). (2004). Hepatitis B vaccines. *Weekly Epidemiological Record*, 79(28), 255–263.
7. André, F. (2000). Hepatitis B epidemiology in Asia, the Middle East and Africa. *Vaccine*, 18(Suppl 1), S20–S22. [https://doi.org/10.1016/S0264-410X\(99\)00456-9](https://doi.org/10.1016/S0264-410X(99)00456-9)
8. Maynard, J. E. (1990). Hepatitis B: global importance and need for control. *Vaccine*, 8(1), S18–S20. [https://doi.org/10.1016/0264-410X\(90\)90209-5](https://doi.org/10.1016/0264-410X(90)90209-5)
9. Romano, L., Paladini, S., Van Damme, P., & Zanetti, A. R. (2011). The worldwide impact of vaccination on the control and protection of viral hepatitis B. *Digestive and Liver Diseases*, 43, S2–S7. [https://doi.org/10.1016/S1590-8658\(10\)60685-8](https://doi.org/10.1016/S1590-8658(10)60685-8)
10. Sung, J. L., & The Asian Regional Study Group. (1990). Hepatitis B virus eradication strategy for Asia. *Vaccine*, 8(Suppl), S95–S99. [https://doi.org/10.1016/0264-410X\(90\)90227-d.PMID2139290](https://doi.org/10.1016/0264-410X(90)90227-d.PMID2139290)
11. Qua, C. S., & Goh, K. L. (2011). Liver cirrhosis in Malaysia: Peculiar epidemiology in a multiracial Asian country. *Journal of Gastroenterology and Hepatology*, 26(8), 1333–1337. <https://doi.org/10.1111/j.1440-1746.2011.06732.x>
12. Wisnom, C., & Siegel, M. A. (2003). Advances in the diagnosis and management of human viral hepatitis. *Dental Clinics of North America*, 47(3), 431–447. [https://doi.org/10.1016/S0011-8532\(03\)00021-1](https://doi.org/10.1016/S0011-8532(03)00021-1)
13. Reichen, J., & Grob, P. (2002). Hepatitis B virus infection: Diagnosis, clinical sequelae, therapy and prevention. *Praxis*, 91(8), 307–319.
14. Mushtaq, M. U., Gull, S., Khurshid, U., Shahid, U., Shad, M. A., & Siddiqui, A. M. (2011). Prevalence and socio-demographic correlates of stunting and thinness among Pakistani primary school children. *BMC Public Health*, 11(11), 790. <https://doi.org/10.1186/1471-2458-11-790.PMID:21988799;PMCID:PMC3209698>
15. Mahtab, M. A., Rahman, S., Karim, M. F., Khan, M., Foster, G., Solaiman, S., & Afroz, S. (2008). Epidemiology of hepatitis B virus in Bangladeshi general population. *Hepatobiliary & Pancreatic Diseases International*, 7(6), 595–600.
16. Uz-Zaman, M. H., Rahman, A., & Yasmin, M. (2018). Epidemiology of hepatitis B virus infection in Bangladesh: Prevalence among general population, risk groups and genotype distribution. *Genes*, 9(11), 541. <https://doi.org/10.3390/genes9110541>
17. Zaki, H., Darmstadt, G. L., Baten, A., Ahsan, C. R., & Saha, S. K. (2003). Seroepidemiology of hepatitis B and delta virus infections in Bangladesh. *Journal of Tropical Pediatrics*, 49(6), 371–374. <https://doi.org/10.1093/tropej/49.6.371>
18. Harpaz, R., Von Seidlein, L., Averhoff, F. M., Tormey, M. P., Sinha, S. D., Kotsopoulou, K., Lambert, S. B., Robertson, B. H., Cherry, J. D., & Shapiro, C. N. (1996). Transmission of hepatitis B virus to multiple patients from a surgeon without evidence of inadequate infection control. *New England Journal of Medicine*, 334(9), 549–554. <https://doi.org/10.1056/NEJM199602293340901>
19. Lee, W. M. (1997). Hepatitis B virus infection. *New England Journal of Medicine*, 337(24), 1733–1745.
20. United Nations Initiative on Global Geospatial Information Management (UN-GGIM). (2016). Report of the Inter-Agency and Expert Group on sustainable development goal indicators (E/CN.3/2016/2/Rev.1). Accessed December 5, 2018, from <http://ggim.un.org/knowledgebase/KnowledgebaseArticle51479.aspx>

21. Yogambigai, R., Niazzlin, M. T., Mudatsir, M., Harapan, H., Abram, L. W., Subramaniam, M., Rahim, K. A., & Radam, A. (2020). Risk behaviours related to hepatitis B virus infection among adults in Malaysia: A cross-sectional household survey. *Clinical Epidemiology and Global Health*, 8(1), 76–82. <https://doi.org/10.1016/j.cegh.2019.04.011>
22. Islam, M., Islam, K., & Islam, N. (1984). Hepatitis-B virus infection in Dhaka, Bangladesh. *Bangladesh Medical Research Council bulletin*, 10(1), 1–6.
23. Rumi, M. A., Begum, K., Hassan, M. S., Hasan, S. M., Azam, M. G., Hasan, K. N., & Khan, A. K. (1998). Detection of hepatitis B surface antigen in pregnant women attending a public hospital for delivery: Implication for vaccination strategy in Bangladesh. *The American Journal of Tropical Medicine and Hygiene*, 59(2), 318–322. <https://doi.org/10.4269/ajtmh.1998.59.318>
24. Mustafa, M., Islam, M. N., Rahman, M., & Salauddin, A. K. (1989). Prevalence of hepatitis B surface antigen (HBsAg) among parenteral drug abusers at Dhaka. *Bangladesh Medical Research Council Bulletin*, 15(1), 1–7.
25. Ahmad, Q., Chowdhury, S. G., Islam, M. N., Khan, F. D., Alam, M. R., & Miah, A. H. (1991). HBsAg amongstunscreened operated patients. *Bangladesh Medical Research Council Bulletin*, 17(1), 11–16.
26. Sattar, H., & Islam, M. (1996). Hepatitis B virus markers among the prostitutes of Dhaka. *Bangladesh Medical Research Council Bulletin*, 22(1), 8–11.
27. Islam, M. M., Islam, S., Azad, A. K., Alam, M., & Mia, Z. (2016). Sero-prevalence of hepatitis b virus(HBV) infection among students of a public university, Dhaka, Bangladesh. *Jagannath University Journal of Life and Earth Sciences*, 2, 1–6.
28. Ashraf, H., Alam, N. H., Rothermundt, C., Brooks, A., Bardhan, P., Hossain, L., Salam, M. A., Hassan, M. S., Beglinger, C., & Gyr, N. (2010). Prevalence and risk factors of hepatitis B and C virusinfections in an impoverished urban community in Dhaka, Bangladesh. *BMC Infectious Diseases*, 10, 1–8. <https://doi.org/10.1186/1471-2334-10-208>
29. Health Bulletin. (2018). Ministry of health and family welfare. Government of People's Republic of Bangladesh. Accessed July 8, 2019, from <https://dghs.gov.bd/images/docs/Publicaations/HB%202018%20final.pdf>
30. Piot, P., Goilav, C., & Kegels, E. (1990). Hepatitis B: Transmission by sexual contact and needle sharing. *Vaccine*, 8, S37–S40.
31. Atkins, M., & Nolan, M. (2005). Sexual transmission of hepatitis B. *Current Opinion in Infectious Diseases*, 18(1), 67–72.
32. Gorgos, L. (2013). Sexual transmission of viral hepatitis. *Infectious Disease Clinics*, 27(4), 811–836.
33. Lichtenhan, J. (1995). Sexual activity and recommendations for the hepatitis B vaccine. *American Family Physician*, 52(5), 1277–1278.
34. Walling, A. (1998). Role of semen in the sexual transfer of hepatitis viruses. *American Family Physician*, 58(2), 509–509.
35. Shepard, C. W., Simard, E. P., Finelli, L., Fiore, A. E., & Bell, B. P. (2006). Hepatitis B virus infection: Epidemiology and vaccination. *Epidemiologic Reviews*, 28, 112–125. <https://doi.org/10.1093/epirev/mxj009>
36. Spijkerman, I. J., van Doorn, L. J., Janssen, M. H., Wijkman, C. J., Bilkert-Mooiman, M. A., Coutinho, R. A., & Weers-Pathoff, G. (2002). Transmission of hepatitis B virus from a surgeon to hispatients during high-risk and low-risk surgical procedures during 4 years. *Infection Control and Hospital Epidemiology*, 23(6), 306–312. <https://doi.org/10.1086/502056>
37. Allos, B. M., & Schaffner, W. (2007). Transmission of hepatitis B in the health care setting: The elephant in the room ... or the mouse? *Journal Infectious Diseases*, 195(9), 1245–1247. <https://doi.org/10.1086/513436>
38. Gershon, R. R., Sherman, M., Mitchell, C., Vlahov, D., Erwin, M. J., Lears, M. K., Felknor, S., Lubelczyk, R. A., & Alter, M. J. (2007). Prevalence and risk factors for bloodborne exposure andinfection in correctional healthcare workers. *Infection Control and Hospital Epidemiology*, 28(1), 24–30. <https://doi.org/10.1086/510813>
39. Rumi, N. A., Sultana, R., Luby, S. P., Islam, M. S., Uddin, M., Hossain, M. J., Zaman, R. U., Nahar, N., & Gurley, E. S. (2014). Infrastructure and contamination of the physical environment in three Bangladeshi hospitals: Putting infection control into context. *PLoS ONE*, 9(2), e89085. <https://doi.org/10.1371/journal.pone.0089085>
40. Chiarello, L. A., & Cardo, D. M. (2002). Preventing transmission of hepatitis B virus from surgeons to patients. *Infection Control & Hospital Epidemiology*, 23(6), 301–302.
41. Saag, M. S., Squires, K. E., Aberg, J. A., & Bardeguez, A. (2010). Letter in response to the new SHEA guideline for healthcare workers with hepatitis B virus, hepatitis C virus, and/or human immunodeficiency virus. *Infection Control Hospital Epidemiology*, 31(10), 1092–1093. <https://doi.org/10.1086/656559>
42. Li, X., Zheng, Y., Liau, A., Cai, B., Ye, D., Huang, F., Sheng, X., Ge, F., Xuan, L., Li, S., & Li, J. (2012). Hepatitis B virus infections and risk factors among the general population in Anhui province China: An epidemiological study. *BMC Public Health*, 12, 272. <https://doi.org/10.1186/1471-2458-12-272>
43. Khan, M., & Ahmad, N. (1996). Seroepidemiology of HBV and HCV in Bangladesh. *International Hepatology Communications*, 5(1), 27–29.
44. Lu, F. M., & Zhuang, H. (2009). Prevention of hepatitis B in China: Achievements and challenges. *Chinese Medical Journal*, 122(24), 2925–2927.
45. Wait, S., & Chen, D. S. (2012). Towards the eradication of hepatitis B in Taiwan. *The Kaohsiung Journal of Medical Sciences*, 28(1), 1–9.
46. World Health Organization (WHO). (2016). Combating hepatitis B and C to reach elimination by 2030. Accessed August 6, 2020, from <https://www.who.int/hepatitis/publications/hep43elimination-by-2030-brief/en/>
47. Sarkar, P., Sarker, N., Doulah, S., & Bari, T. I. (2017). Expanded programme on immunization in Bangladesh: A success story. *Bangladesh Journal of Child Health*, 39(2), 93–98. <https://doi.org/10.3329/bjch.v39i2.31540>
48. Lu, P. J., O'Halloran, A. C., Williams, W. W., & Nelson, N. P. (2018). Hepatitis B vaccination coverage among adults aged ≥ 18 years traveling to a country of high or intermediate endemicity, United States, 2015. *Vaccine*, 36(18), 2471–2479. <https://doi.org/10.1016/j.vaccine.2018.03.030>
49. Lewis-Ximenez, L. L., do O, K. M., Ginuino, C. F., Silva, J. C., Schatzmayr, H. G., Stuver, S., & Yoshida, C. F. (2002). Risk factors for hepatitis B virusinfection in Rio de Janeiro, Brazil. *BMC Public Health*, 2, 26. <https://doi.org/10.1186/1471-2458-2-26>

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