

Original Article

Evaluation of the antibody response against Hepatitis B Virus infection in patients on maintenance hemodialysis: A Pilot Study

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Abstract

Objective: This study was undertaken to evaluate the antibody response of hepatitis B virus infection in patients on maintenance hemodialysis (MHD) by detecting different viral markers. **Method:** Study subjects comprised a total of 88 chronic kidney disease (CKD) patients from Bangladesh Institute of Research and Rehabilitation in Diabetes, Endocrine and Metabolic Disorders (BIRDEM) and Bangabandhu Sheikh Mujib Medical University (BSMMU). Of them 63 patients on MHD and 25 predialysis patients served as cases and controls respectively. Clinical history was taken and serological markers for HBV (HBsAg, Anti-HBs, and Anti-HBc) were determined by using ELISA. **Results:** Hepatitis B virus was positive in 1.6% of maintenance hemodialysis (MHD) patients and in 16% of controls ($p < 0.02$). Anti-HBc antibody was positive in 62% of dialysis patients and 72% of controls ($p = \text{NS}$) and the positivity was significantly associated in dialysis subjects with longer duration of dialysis (18 ± 22 vs. 10 ± 7 , months, $p < 0.04$), multiple units of blood transfusions (22 ± 29 vs. 10 ± 12 , units, $p < 0.04$) and more reuse of dialyzer (3 ± 1 vs. 2 ± 1 , times, $p < 0.03$) than the negative ones. Among MHD patients 84% were vaccinated against HBV with a schedule of 3 (79%) and 4 (21%) doses and protective antibody titer (>10 IU/L) was found in 57%. None of the controls were vaccinated but 66% had protective titer indicating post exposure natural immunity. **Conclusions:** Hepatitis B virus positivity was significantly higher among the predialysis subjects compared to dialysis group.

Key words: Hepatitis B virus, Antibody response, Hemodialysis

Introduction

End-stage renal disease (ESRD) subjects on maintenance hemodialysis are at high risk for hepatitis B virus infection¹. Parenteral route is the major route for HBV transmission². The process of hemodialysis requires vascular access for prolonged period³. Furthermore, hemodialysis patients are immunosuppressed⁴ that increases their susceptibility to infection requiring frequent hospitalization and surgery, which again increases their risk for exposure to

nosocomial infections³. Although vaccination is routinely recommended in ESRD patients, antibody response to vaccination is suppressed and its level rapidly declines among patients on chronic dialysis due to the decreased immunological responses⁵. The prevalence of chronic hepatitis B virus (HBV) infection is high ($>8\%$) in sub-Saharan Africa, most of Asia and the Pacific Islands, intermediate prevalence (2 to 7%) regions include the

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Amazon, southern parts of Eastern and central Europe, the Middle East and the Indian sub-continent, low prevalence (<2%) regions include most of Western Europe and North America⁶. In India, HBV prevalence was 8.8% and 14.2% in predialysis and hemodialysis group respectively^{7,8}. In Turkey, prevalence of HBV was 10.5% and 13.3% in predialysis and hemodialysis patients respectively^{9,10}. In Bangladesh, around 12% of all patients on MHD were serologically positive for hepatitis B virus infection, has been shown in a recent study¹¹. So far no study has been conducted to see the seroprevalence of HBV in CKD (predialysis) patients in Bangladesh. Therefore, this study was undertaken to evaluate the antibody status of HBV in predialysis and dialysis patients followed in two selected tertiary renal care center

Subjects and methods

Study design

This cross sectional study was carried out in the Department of Immunology, BIRDEM, Dhaka and Nephrology Department of BSMMU during the period of June 2006 to June 2007.

Study subjects

Eighty-eight patients were finally included in this study. Of them 63 end stage renal disease (ESRD) patients who were on maintenance hemodialysis for at least 3 months and getting dialysis through

arteriovenous (AV) fistula considered as cases and 25 chronic renal failure (CRF) patients attending Nephrology Out-patients departments of BIRDEM and BSMMU and 'CRF patients follow-up project' who were not on dialysis (predialysis) considered as control group.

Sample collection and preservation

Five milliliter blood was taken from the arterial channel immediately after pricking the fistula during dialysis session in MHD patients and labeled with a known serial number for each patient. In controls fasting samples were taken. Serum samples were preserved at -20°C and assayed within fifteen days of collection.

Laboratory analysis

Serological markers for hepatitis B (HBsAg, Anti-HBc, Anti-HBs) were assessed using commercial third generation enzyme-linked immunosorbent assay kit (Diasorin, Italy). Serum creatinine and alanine aminotransferase were assessed by standard laboratory method (Kinetic method).

Statistical analysis of data

All the relevant data were entered and then analyzed using the statistical package for social science (SPSS) version 13. Results were expressed as mean \pm SD or percentage as appropriate. Level of significance was expressed as 'P' value and $p < 0.05$ was considered as significant.

Results

Table I- Baseline Parameters of Study Subjects

Parameters	Cases (n=63)	Controls (n=25)	P Value
Age (yrs)	54±11	57±10	0.28
M/F	36/27	15/10	0.80
DM Duration(yrs)	12±6	5±2	0.92
CKD Duration(yrs)	6±4	4±3	0.53
S Cr (mg %)	9±2.5	4±2	0.001
ALT (U/L)	25±17	20±16	0.30

DM = Diabetes mellitus, results are expressed in mean ± SD on percentage where suitable, M/F = Male/Female, CKD = Chronic Kidney Disease, S Cr = Serum Creatinine, ALT = Alanine Aminotransferase

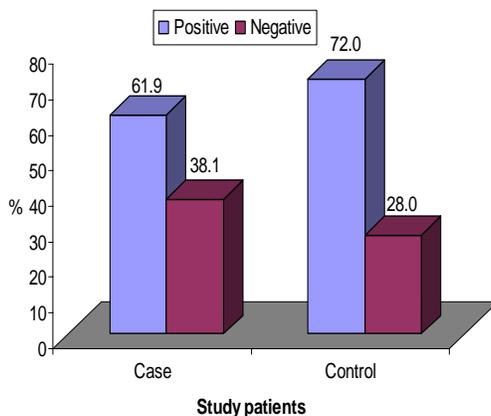
Table II - Distribution of patients by hepatitis B virus infection

HBsAg	Cases	Controls	P Value
	n (%)	n (%)	
Positive	1 (1.6%)	4 (16%)	0.022
Negative	62 (98.4%)	21 (84%)	

HBsAg – Hepatitis B surface antigen

P value reached from chi square test; $p < 0.05$ considered significant

Figure 1- Distribution of patients by anti-HBc total



Note: Figure 2 above indicated that among cases and controls, anti-HBc positive was 61.9% (n=39) vs. 72% (n=18) ($P=NS$) and this was not significantly different between the two groups

Table III- Distribution of Cases (MHD patients) by anti-HBc total

Parameters	Positive (n=39)	Negative (n=24)	P Value
Age (yrs)	52±10	58±11	0.03
CKD Duration(yrs)	6±4	5±2	0.41
DM Duration(yrs)	14±7	13±7	0.28
Dialysis Duration (m)	18± 22	10±7	0.03
BT (total units)	22± 29	10±12	0.03
BT (units/month)	1.4± 1.2	1.1±1.3	0.07
Dialyzer Reuse	3±1	2±1	0.02

CKD- Chronic Kidney Disease, DM – Diabetes mellitus, BT- Blood transfusion, m - months
Results are shown in mean ± SD

Table IV - Level of immunity against hepatitis B virus in cases (MHD)

	Vaccinated n=53, (84%)	Non-vaccinated n=10, (16%)	P Value
Protective (>100IU/l)	11 (19%)	4 (57%)	0.74
Low protective (10-100IU/L)	20 (39%)	-	
Non protective (<10IU/L)	22 (42%)	3 (43%)	

P value reached from chi square test; $p < 0.05$ considered significant

Different laboratory parameters were similar between cases (MHD patients) and controls (predialysis patients). The only difference was in serum creatinine (S. Cr) level and this was higher in MHD patients ($p < 0.001$) (Table I).

The proportion of positive hepatitis B virus infection was found to be higher among the control (16%) compared to case (1.6%) and the difference was statistically significant ($p < 0.02$) (Table II).

Results showed no significant association between positivity of anti-HBc with duration of chronic renal failure and duration of diabetes mellitus (Table III). However, data shows higher preponderance of positive anti-HBc among the patients with prolonged

duration of maintenance haemodialysis ($p < 0.03$) and number of total units of blood transfusion ($p < 0.03$). A statistically significant association was also found between anti-HBc status and number of reuse of dialyser ($p < 0.02$) indicating the positivity of anti-HBc was high among the patients with more reuse of dialyser. Data analysis also indicated that the mean age of the positive anti-HBc total was found to be low (52.21 ± 10.2 years) compared to negative anti-HBc total (57.96 ± 10.8 years) and the difference was significant ($p < 0.05$). Result showed that 84% of the cases (dialysis patients) were vaccinated and 16% of them non-vaccinated ($P < 0.001$). No significant difference was seen in proportion

of the patients with protective and non-protective titers among the vaccinated and non-vaccinated subjects. Vaccination schedule was 3 doses in 79% and 4 doses in 21% of cases (Table IV).

Discussion

Bangladesh has an intermediate prevalence of hepatitis B virus infection with a 4% HBsAg positive population¹². In a previous study 3.5% prevalence rate of hepatitis B virus infection in pregnant woman of Bangladesh was seen¹³. It was observed in the present study that HBsAg was positive in hemodialysis patients in lower frequency (1.6%). Lower prevalence of HBsAg in MHD patients probably due to routine screening for HBsAg before selection of patients for MHD. Furthermore, we selected patients from two hemodialysis units where HBsAg positive patients were not accepted for hemodialysis to minimize the risk of spreading of HBV infection. This notion was similar to¹⁴. Moreover, 84% of our dialysis patients were vaccinated which might also contribute to low number of HBsAg positive cases in MHD group. On the other hand, HBsAg was found in higher frequencies in our predialysis diabetic CKD patients (16%). In diabetic patients HBV infection has been shown about 14% in a study¹⁵ and in predialysis patients it ranged from 8-10% in studies from India and Turkey^{7,9}. Higher prevalence of HbsAg in our predialysis patients may be attributed to the fact that these patients had not been undergone routine screening for HBsAg and vaccination. Although, other factors like, frequent hospitalization, history of injections (including insulin), poor nutritional status (metabolic derangement) leading to suppressed immune response etc. remain common for both groups. Therefore, HBV vaccination alone seems to make the difference of HBsAg status observed in both groups.

We found 62% of the dialysis patients and 72% controls are anti-HBc positive, which is higher than some other reports showing around 40%¹⁶ but similar to one showing 76%¹⁷. Positive anti-HBc total always indicate a remote HBV infection and is the most valuable single serologic marker in diagnosis of HBV infection even when HBsAg remains negative. We found positivity of anti-HBc total higher among the patients with more reuse of dialyzer¹⁸, Qadi *et al.* reported reuse of dialyzer is one of the risk factors for viral transmissions. Similarly increased duration of dialysis and higher number of blood transfusion has been shown to be associated with increased frequency of HCV infection¹⁹, which was seen in our patients too. It is possible that our patients became more anti-HBc positive due to higher blood transfusion and longer duration of dialysis.

Majority of our MHD patients had vaccination (84%) but analysis found that only 19% have protective immunity, 39% low protective immunity and 42% have non protective immunity indicating that approximately half of the dialysis patients had no protection despite vaccination which is probably due to immunosuppression. In present study in all immunity groups, majority of the patients were more than 50 years ages. It has been suggested that advanced age reduces the response against HBV vaccine in hemodialysis patients²⁰. In our study response against hepatitis B vaccine to attain a protective titer was in 58% subjects. This relatively low response may be due to higher age, presence of diabetes and lower doses of vaccination schedule as majority (79%) took three-dose regimen. Some studies showed that vaccine response is 64% with 3 doses whereas 86% with 4 doses²¹.

We can conclude that Hepatitis B virus positivity was significantly higher among the predialysis subjects compared to dialysis group. Anti-HBc antibody was positive in two-third of the study subjects. With a three-dose vaccine schedule only half of the dialysis patients could attain protective antibody titer. According to standard statistical formula, a large sample size should have been taken to reflect the picture

of whole population. However, as this is a Pilot study sample size was confined at 88 subjects.

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References

1. Otedo AE, MC Ligeo SO, Okoth FA, Kayima JK. Seroprevalence of hepatitis B and hepatitis C in maintenance dialysis in a public hospital in a developing country. *South African Medical Journal* 2003; **93**:380-4.
2. Teles SA, Martins RMB, Lopes CLR, Carneiro MA, Souza KP, Yoshida CFT. Immunogenicity of a recombinant hepatitis B vaccine (Euvax-B) in a hemodialysis patients and staff. *European Journal of Epidemiology* 2001; **17**: 145-49.
3. Busek SU, Baba EH, Tavares Filho HA, Pimenta L, Salomao A, Correa-Oliveira R, Oliveira GC. Hepatitis C and hepatitis B virus infection in different hemodialysis units in Belo Horizonte, Minas Gerais, Brazil. *Mem Inst Oswaldo Cruz* 2002; **97**: 775-778.
4. Devesa M, Khudyakov YE, Capriles F, Blitz L, Fields HA, Liprandi F, Pujol FH. Reduced Antibody Reactivity to Hepatitis C Virus Antigens Hemodialysis Patients Coinfected with Hepatitis B Virus. *Clinical and Diagnostic Laboratory Immunology* 1997; **4**: 639-642.
5. Tong NKC, Beran J, Kee SA, Miguel JL, Sanchez C, Bayas JM, Vilella A, De Juanes JR, Arrazola P, Torrecillas FC, De Novales EL, Hamtiaux V, Lievens M, Stoffel M. Immunogenicity and safety of an adjuvanted hepatitis B vaccine in pre-dialysis and hemodialysis patients. *Kidney Int* 2005; **68**: 2298-2303.
6. Jadallah RI, Adwan GM, Hasan NA, Adwan KM. Prevalence of hepatitis B virus markers among high risk groups in Palestine. *Medial Journal Islamic World Academy of Sciences* 2005; **15**: 157-60.
7. Ahmed B, Grover R, Ratho RK, Mahajan RC. Prevalence of hepatitis B virus infection in Chandigarh over a six years period. *Trop Gastroenterol* 2001; **22**:18-19.
8. Chattopadhyay S, Rao S, Das BC, Singh NP, Kar P. Prevalence of transmitted virus infection in patients on maintenance hemodialysis from New Delhi, India. *Hemodial Int* 2005; **9**: 362-6.
9. Sit D, Kadiroglu AK, Kayabasi H, Yilmaz ME, Goral V. Seroprevalence of hepatitis B and C viruses in patients with chronic kidney disease in the predialysis stage at a university hospital in Turkey 2007; **50**: 133-7.
10. Yakaryilmaz F, Gurbuz OZ, Gulter S, Mert A, Songur Y, Karakan T. Prevalence of occult hepatitis B and hepatitis C virus

infection in Turkish hemodialysis patients. *Ren Fail* 2006; **28**: 729-35.

11. Islam MN, Hossain RM, Rahman MH, Mansur MA, Hassan MS, Islam MS, Sultana R, Iqbal MM. Hepatitis B (HBV) and C (HCV) among maintenance hemodialysis patients, family members and dialysis staffs. *Intern Society Hemodial* 2007; **11**: 108.

12. Kane M, 1994. Global Plan of Action for Hepatitis B Immunization: Global Program for Vaccine and Immunization. Expanded Program on Immunization. Geneva: World Health Organization.

13. Rumi MAK, Begum K, Hassan MS, Hasan SMM, Azam MG, Hasan KN, Shirin M, Khan AKA. Detection of hepatitis B surface antigen in pregnant woman attending a public hospital for delivery: Implication for vaccination strategy in Bangladesh. *Am J Trop Med Hyg* 1998; **59**: 318-322.

14. Qadi AA, Tamim H, Ameen G, Bu-Ali A, Al-Arrayed S, Fawaz NA, Almawi WY. Hepatitis B and hepatitis C virus prevalence among dialysis patients in Bahrain and Saudi Arabia: A survey by serologic and molecular methods. *Am J Infect Control* 2004; **32**: 493-5.

15. Chen HF, Li CY, Chen P, See TT, Lee HY. Seroprevalence of Hepatitis B and C in Type 2 Diabetic Patients. *J Chin Med Assoc* 2006; **69**: 146-152.

16. Souza KP, Luj A, Teles SA, Carneiro MAS, Oliveira LA, Gomes AS, Dias MA, Gomes SA, Yoshida CFT. Hepatitis B and C

in the Hemodialysis Unit of Tocantins, Brazil: Serological and Molecular Profiles. *Mem Inst Oswaldo Cruz, Rio de Janeiro* 2003; **98**: 599-603.

17. Ambuhl PM, Binswanger U, Renner EL. Epidemiology of chronic hepatitis B and C among dialysis patients in Switzerland. *Schweiz Med Wochenschr* 2000; **130**: 341-348.

18. Saxena Anil K, Panhotra BR. Impact of dedicated space, dialysis equipment, and nursing staff on the transmission of hepatitis C virus in a hemodialysis unit of the Middle East. *Am J Infect Control* 2003; **31**:26-33.

19. Albuquerque ACCD, Coelho MRCD, Lopes EPA, Lemos MF, Moreira RC. Prevalence and risk factors of hepatitis C virus infection in hemodialysis patients from one center in Recife, Brazil. *Mem inst Oswaldo Cruz Rio de Janeiro* 2005; **100**: 467-70.

20. Taheri Sh, Shahidi Sh, Moghtaderi J, Seirafian Sh, Emami A, Eftekhari SM. Response rate to Hepatitis B Vaccination in Patients with Chronic Renal Failure and End-Stage-disease: Influence of Diabetes Mellitus. *J Res Med Sci* 2005; **10**: 384-390.

21. Centers for Diseases Control and Prevention. Recommendation for preventing transmission of infections among chronic hemodialysis patients. *MMWR* 2001; **50**: 1-43.

“Allah has not sent down a disease except that He also sent down its cure: whoever knows it (the cure), knows it, and whoever is unaware of it (the cure), he is unaware of it.” (the medicine) while those who are ignorant of it are unaware of it.” [An-Nasai’, Ibn Mājah, Al-Hakim and Ibn Hibban]