Prevalence, SeroconversionAnd Risk Factors Of Hepatitis B And C Infection In Haemodialysis Patients At District Hospital Of Mehsana Komal Patel*, Gaurishankar Shrimali**

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Abstracts:<u>Background& Objective</u>: Hepatitis B and Hepatitis C are significant problems in the management of haemodialysis patients.We aimed to investigate the incidence and prevalence of HBV and HCV infection in the hemodialysis patients as well as risk factors for infection.<u>Methodology</u>:All adult patients receiving maintenance hemodialysis (n=150) in District hospital, Mehsana were studied between June 2014 to October 2015. Testing for Hepatitis B surface antigen (HBsAg) and anti-HCV antibodies was performed at initiation of dialysis and every 3 monthly thereafter. Patients who were sero-negative for HBV and HCV were followed up for 1 year to detect sero-conversions.<u>Results</u>: The prevalence and seroconversion rates were 11.33% and 4.66% in HBV patients and 14% and 6% in HCV patients respectively. There was a significant correlation of the prevalence and seroconversion of HCV and HBV with number of blood transfusion and duration of heamodialysis.<u>Conclusion</u>:Patients on maintenance hemodialysis have a high incidence and prevalence of HCV infection and lower rates of HBV infection in this study. The factors associated with HBV and HCV infection are highly suggestive of nosocomial transmission withinhemodialysis units. Strict infection control measures are required.[Patel K NJIRM 2015; 6(6):19-23] **Key Words**: Haemodialysis, Hepatitis B, Hepatitis C, Incidence, Nosocomial infection

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Introduction: Infections with hepatitis B virus (HBV) and hepatitis C virus (HCV) are well-known and important causes of liver disease in end-stage renal failure patients on hemodialysis^{1,2,3,4}. Patients receiving maintenance haemodialysis (HD) therapy are at increased risk for acquiring these infections and have a higher prevalence of HBV and HCV than the general population^{5,6}. The introduction of blood donor screening and a reduction in blood transfusions due to the availability of recombinant erythropoietin has significantly reduced the incidence of new HCV infections among HD patients in many countries^{7,8,9}. HBV infection is usually due to patient to patient transmission within HD units¹⁰. Prior to effective screening of blood donations, HCV infection was associated with blood transfusions needed to correct the anaemia associated with kidney disease^{11,12} but patient to patient transmission in HD units is also reported^{13,14}. Recognition of the risk of nosocomial infection has resulted in recommendations that strict infection control procedures should be followed on HD units; patients with blood-borne virus infections should be isolated from sero-negative patients during dialysis and patients as well as staff should be vaccinated against hepatitis B.This study aimed to investigate for the first time the incidence and prevalence of HBV and HCV infection in the entire HD population at district hospital of Mehsana.

Material and Methods:The study was performed in haemodialysis unit of general hospital, Mehsana after

permission of respective authorities.A total170 patients were enrolled in this study from june 2014 to october 2015. 150 patients gave informed consent and thus 92.5% of the whole patient population were investigated. The patients who were on hemodialysis for a minimum period of 1 month and were likely to be available for follow-up for at least 6 months were included in the study. The patients who had acute renal failure were not included in the study. The patients were monitored forseroconversion every 3 months during their follow-up visit using HbsAg ELISA and Anti-HCV ELISA tests.

All tests were carried out and interpreted strictly in accordance with the manufacturer's instructions. Allsero-conversions were recorded and included even if patients was transplanted or died afterwards. All patients were frequently monitored during the study period to assure the inclusion of every new seroconversion.All new sero-conversions were retested to confirm positive result. All samples were also tested for SGPT and SGOT.

Results:Total 150 patients were observed at least for a period of 6months for seroconversion.Among them 85 were males(56.66%) and 65 were females(43.33%).Out of total 150 patients, 17(11.33%) were reactive to HbsAg ELISA test and 24(14%) were reactive to anti-HCV ELISA test.

Patient Status	Prevalence	Seroconversion
HbsAg reactive	17(11.33%)	7(4.66%)
HbsAg non-	133(88.6%)	-
reactive		
HCV reactive	24(14%)	9(6%)
HCV NON-	126(84%)	-
reactive		

Maximum number of patients underwent seroconversionbetween 6month to 1 year after the start of heamodialysis.7 patients (4.66%) who were initially non-reactive became HbsAg reactive during follow-up. Of these one had previously been vaccinated against hepatitis B, four had not been vaccinated and in two vaccination status was not documented. 9 patients (6.00%) who were initially non-reactive became HCV reactive during follow-up.

Out of 17 HbsAgractive 11(64.70%) received blood transfusion and out of 133 HbsAg nonreactive 34(25.56%) received blood transfusion.Out of 24 HCVractive 17(70.83%) received blood transfusion and out of 126 HCV nonreactive 35(27.77%) received blood transfusion. In our study 20 (60.60%) HCV reactive patients had history of blood transfusion more than 10times and 12 (36.36%) HCV reactive patients had history of blood transfusion more than 5times. only 1 (3.03%) HCV reactive patient had no history of blood transfusion. In our study 10(41.66) HbsAg reactive patients had history of blood transfusion more than 10times and 8 (33.33%) HbsAg reactive patients had history of blood transfusion more than 5times. 6 (25%) HbsAg reactive patient had no history of blood transfusion.

In our study 25 patients were Hcv reactive having history of heamodialysis more than 6 months. Among HCV reactive patients 8 were on heamodialysis every 7th day and 12 HCV reactive patients on heamodialysis every 3rd day. We did not detected HCV infection in patients who were on heamodilysis every 20th day.

Most common cause for end stage renal disease was diabetes mellitus (45%). 2nd most common cause was hypertension (22%).We found that 7% patients were HCV reactive, having history of major invasive procedure and 3% had history of minor invasive procedure.

2(8.33%) HbsAgractivepatients,had history ofi.v. drug abuse while 8 (6.01%) HbsAg non-reactive patients, had history of i.v. drug abuse. Similarly 4(12.12%) anti-HCV reactive patients,had history of i.v. drug abuse while 5 (3.96%) anti-HCVnon-reactive patients, had history of i.v. drug abuse.Hepatitis B vaccine had been administered in 133 of 150 patients but antibody levels were not checked post vaccination. Vaccination status was not documented in remaining 17(11.33%) patients.

Mean values for Alanine Aminotransferase and Aspartate Aminotransferase were higher in seropositive patients despite being within the normal range.

Discussion:Hepatitis B and C infections are serious problem in chronic haemodialysis patients. This study was conducted to determine the prevalence and seroconversion characteristics of HbsAg and anti-HCV in patients with CRF, with the tests done at baseline and repeated after every 3 months. The prevalence and seroconversion rates were 16%% and 6% respectively for HCV. The prevalence of anti-HCV in patients on heamodialysis from india is reported to be in the range of 3% to 45%¹⁵. The prevalence of HCV infection is known to very widely in different regions of the world. The reason for variation in prevalence of HCV in heamodialysis patients is largely unknown¹⁶. Implemention of universal precaution in HD unit, method of HCV testing, blood transfusionand variable policy isolation could be the potential of reason.Prevalence of HCV in our study is relatively higher then other study from india. Possible reason for that are lack of knowledge of nursing staff and high prevalence of hepatitis C in general population.

The prevalence and seroconversion rates were 11.33% and 4.66% respectively for HBV. In India, prevalence in HD patients ranged from 3.4% to 45% which is clearly in excess to the prevalence of 4.7% in the general populatio¹⁷. The study conducted by Bhowmiket all in HD patients, shows 5.5% seroconversion for hepatitis B infection¹⁸. Our study shows relatively higher prevalence of hepatitis B than other study. Possible reason is we receive patients at a relatively later stage of CKD and effectivity of hepatitis B vaccination decreases as renal function declines, making patients prone to hepatitis B infection in spite of vaccinationUse of HBV vaccine, use of dedicated dialysis machines and regular surveillance are the

20

possible reason for less prevalence of HBV than HCV in HD unit¹⁹.

As per our study seroconversion rate for hepatitis B and hepatitis C were 4.66% and 6% respectively. In study conducted by VikasMakkar et al seroconversion rate for hepatitis B and Hepatitis C were 4.8% and 6.8% respectively²⁰. In study conducted by Kumar etalseroconversion rate for hepatitis C was 7.44%¹⁶.

A positive history of blood transfusions as well as the number of blood transfusions was strongly associated with HBV or HCV infection in our study.Several study have shown that the risk of acquiring the HCV infection increase with an increase in the number of units of blood which were transfused^{21,22,23}.

On the other hand, the prevalence and incidence of HBV or HCVsero-positivity was significantly related to the length of time on HD. This is consistent with nosocomial transmission related to dialysis since longer duration of dialysis represents a longer period at risk of acquiring an infection. Similar observations have been reported by other authors^{24,25,26}. Prevention of nosocomial transmission is of vital importance as HCV antiviral treatment is expensive and its availability is limited to only a few centres.

There was no significant differences in HCV infection among patients undergone for major and minor invasive procedure. So irrespective of type of invasive procedure breach in body tissue will increase the risk of HCV infection. Few case series also observed high prevalence of HCV infection in IV drug abuse^{27,28}.

Mean values for Alanine Aminotransferase and Aspartate Aminotransferase were higher in seropositive patients despite being within the normal range in this study.Various investigators confirm the lack of sensitivity or specificity of ALT and AST as a surrogate marker for chronic liver disease. mainly in heamodialysis patients^{29,30,31}. There is no clear explanation for this. But possible reason is thought that uremia suppresses serum transferases. Therefore, HCV RNA and liver histology rather than serum transferases should probably be the means to determine the presence or absence of liver disease in hemodialysis patients, since these two parameters correlate relatively well in the majority of patients. HCV infected heamodialysis patients may develop liver damage despite normal ALT levels.

Our study had several limitations. First, we did not use HCV RNA due to financial limitation which is recomondedtest for HCV in heamodialysis patients in areas of higher prevalence. Secondly, we did not evaluate efficasy of HBV vaccine in our patients and whether a relatively higher rate of seroconversion was due to inadequate antibody titre. Thirdly, we did not analyse data regarding patients receiving dialysis at other centres and its contribution to seroconversion.

Conclusion: In conclusion, patients on maintenance HD in District hospital Mehsana have a higher incidence and prevalence of HCV infection than HBV infection. The factors associated with HBV and HCV infection are highly suggestive of nosocomial transmission. Urgent action is required to improve infection control measures in HD centre and to reduce dependence on blood transfusions for the treatment of anaemia.

References:

- Alavian SM, Bagheri-Lankarani K, Mahdavi-Mazdeh M, Nourozi S. Hepatitis B and C in dialysis units in Iran: changing the epidemiology. Hemodial Int. 2008;12(3):378–82.
- Fabrizi F, Messa PG, Lunghi G, Aucella F, Bisegna S, Mangano S, Villa M, Barbisoni F, Rusconi E, Martin P. Occult hepatitis B virus infection in dialysis patients: a multicentre survey. Aliment PharmacolTher. 2005;21(11):1341–7.
- Reddy GA, Dakshinamurthy KV, Neelaprasad P, Gangadhar T, Lakshmi V. Prevalence of HBV and HCV dual infection in patients on haemodialysis. Indian J Med Microbiol. 2005;23(1):41–3.
- Wong PN, Fung TT, Mak SK, Lo KY, Tong GM, Wong Y, Loo CK, Lam EK, Wong AK. Hepatitis B virus infection in dialysis patients. J Gastroenterol Hepatol.2005;20(11):1641–51.
- 5. Fabrizi F, PoordadFF, Martin P: Hepatitis C infection and the patient with end-stage renal disease.*Hepatology* 2002, 36(1):3-10.
- 6. Fabrizi F, Lunghi G, Martin P: Hepatitis B virus infection in hemodialysis: recent discoveries. *JNephrol* 2002, 15(5):463-468
- Mohamed WZ: Prevention of hepatitis C virus in hemodialysis patients: five years experience from a single center.*Saudi J Kidney Dis Transpl* 2010, 21(3):548-554
- 8. Patel PR, Thompson ND, KallenAJ, ArduinoMJ: Epidemiology, surveillance, and prevention of hepatitis C virus infections in

NJIRM 2015; Vol. 6(6) Nov – Dec

21

hemodialysis patients.*Am J Kidney Dis* 2010, 56(2):371-378

- Saune K, Kamar N, Miedouge M, Weclawiak H, Dubois M, Izopet J, Rostaing L: Decreased prevalence and incidence of HCV markers in haemodialysis units: a multicentric French survey.*Nephrol Dial Transplant* 2010, 26(7):2309-2316
- Ozer A, Yakupogullari Y, Beytur A, Beytur L, Koroglu M, Salman F, Aydogan F: Risk factors of hepatitis B virus infection in turkey: a population-based, case– control study: risk factors for HBV infection.*Hepat Mon* 2011, 11(4):263-268
- 11. Taal MW, van Zyl-Smit R: Hepatitis C virus infection in chronic haemodialysis patients—relationship to blood transfusions and dialyser re-use.*SAfr Med* J 2000, 90(6):621-625
- Knudsen F, Wantzin P, Rasmussen K, Ladefoged SD, Lokkegaard N, Rasmussen LS, Lassen A, Krogsgaard K: Hepatitis C in dialysis patients: relationship to blood transfusions, dialysis and liver disease.*KidneyInt* 1993, 43(6):1353-1356.
- Allander T, Medin C, Jacobson SH, Grillner L, Persson MA: Hepatitis C transmission in a hemodialysis unit: molecular evidence for spread of virus among patients not sharing equipment. JMVirol 1994, 43(4):415-419
- Le Pogam S, Le Chapois D, Christen R, Dubois F, Barin F, GoudeauA: Hepatitis C in a hemodialysis unit: molecular evidence for nosocomial transmission.*JClinMicrobiol* 1998, 36(10):3040-3043
- 15. Agarwal SK, Dash S, Irshad M. Hepatitis C cirus infection during hemodialysis in India. J Assoc. Physic India. 1999,47:1139-43.
- 16. Surendra Kumar P, Venu G, MadhusudhanaRao A et al. Prevalence andrisk factors of hepatitis C among maintenance hemodialysis patients at a tertiary care hospital in Coimbatore, India. Journal Of Clinical and Diagnostic Resarch 2011; 5 (4): 725-728.
- 17. Chawla NS, Sajiv CT, Pawar G, Pawar G. Hepatitis B and C Virus infections associated with renal replacement therapy in patients with end stage renal disease in a tertiary care hospital in India – prevalence, risk factors and outcome.Indian J Nephrol.2005; 15: 205–213.
- 18. Bhaumik P, Debnath K. Prevalence of hepatitis B and C amonghaemodialysis patients of Tripura,

India. Euroasian Journal ofHepato-Gastroenterology 2012; 2 (1): 10–13

- 19. Mittal G, Gupta P, Thakuria P et al. Profile of Hepatitis B virus, Hepatitis C Virus, Hepatitis D Virus and HIV Infections in HemodialysisPatients of a Tertiary Care Hospital in Uttarakhand. J ClinExpHepatol 2013; 3: 24–28.
- VikasMakkar,DineshGupta,KanishBansal,N.S.Khaira Prevalence, seroconversion and risk factors of hepatitis B and C infection in patients on maintenance hemodialysis. Journal of evolution of medical and dental sciences.2014;3(50):11790-11798.
- 21. Dentico P, Buogiorno R; Volpe A. Prevalence and incidence of hepatitisC virus (HCV) in hemodialysis patients: study of risk factors. ClinNephrol 1992; 38: 49-52.
- Saab S, Martion P, Brezina M, Gitrich G et al. Serum alanine aminotransferase in hepatitis C screening of the patients onhemodialysis. Am J Kidney Dis 2001; 37: 308-15.
- 23. Alfurayh O, Sobh M, Buali A, Ali MA, Barri Y, Qunibi W, et al. Hepatitis C virus infection in chronic hemodialysis patients: A Clinicopathologicalstudy.Nephrol Dialysis Transplant 1992; 7: 327.

gMuller GY, Zabaleta ME, Arminio A, Colmenares CJ, Capriles FI, Bianco NE, Machado IV.Risk factors for dialysis-associated hepatitis C in Venezuela. Kidney Int. 1992 Apr; 41(4):1055-8.

- Ayoola EA, Huraib S, Arif M, al-FalehFZ, al-Rashed R, Ramia S, al-Mofleh IA, Abu-Aisha H. Prevalence and significance of antibodies to hepatitis C virus among Saudi haemodialysis patients. J Med Virol. 1991 Nov; 35(3):155-9.
- 25. Hardy NM, Sandroni S, Danielson S, Wilson WJ. Antibody to hepatitis C virus increases with time on hemodialysis. Clin Nephrol. 1992 Jul; 38(1):44-8
- Kalantar-Zadeh K, Kilpatrick RD, McAllister CJ, Miller LG, Daar ES, GjertsonDW, Kopple JD, Greenland S: Hepatitis C virus and death risk in hemodialysis patients. J Am SocNephrol 18: 1584– 1593, 2007.
- AlterMJ, Kruszon-Moran D, NainanOV, McQuillan GM, Gao F, Moyer LA, Kaslow RA, Margolis HS. The prevalence of hepatitis C virus infection in the United States.NEngl J Med.1999;341(8):556-62.
- 28. Al Traif, Huraib S, Abdullah A etal.AntiHCV positive hemodialysispatients:clinical,biochemical,histologi cal and virologic study and a proposed

management scheme. Annals of Saudi medicine.2000;20(3)

- 29. Silini E, Bono F, Cerino A, Piazza V,Solcia E, Mondelli MU. Virological features of hepatitis C virus infection in heamodialysis patients. J Clin Microbiology.1993;31(11):2913-7
- 30. DuBios DB. Gretch D. Dela Rosa C,LeeW,FineJ,BlaggCR,CoreyL. Quantitation of heapatitisCviral RNA in sera of hemodialysispatients: Gender related differences in viral load. Am J Kidney Dis.1994;24(5):795-801

Conflict of interest: None

Funding: None

Cite this Article as:Patel K, Shrimali G.Seroconversion of Hepatitis B and C infection in haemodialysis patients. Natl J Integr Res Med 2015; 6(6): 19-23