

Hepatitis D Is a Forgotten Problem in Hemodialysis Patients in the World

Seyed-Moayed Alavian

Baqiyatallah Research Center for Gastroenterology and Liver Disease (BRCGL),
Baqiyatallah University of Medical Sciences, Founder of Asian Hepatitis Network & Tehran Hepatitis Center,
Tehran, Iran
editor@hepmon.ir

Hepatitis Delta Virus (HDV) is a small defective virus that replicates only in the presence of Hepatitis B surface Antigen (HBsAg) ⁽¹⁾. The epidemiology of HDV infection is similar to HBV but with notable exceptions. Evidence of HDV infection was found all around the world. It is estimated that approximately 5% of HBsAg carriers are infected with HDV infection all around the world. HDV infection occurs worldwide but incidence and prevalence data are limited due to inaccurate reporting and delayed detection. It is more difficult to determine the number of cases of acute or fulminant hepatitis related to HDV infection, as the incidence varies among continents, countries, and regions.

In general, three epidemiological patterns of HDV infection can be identified. They include the endemic pattern (such as what occurs in southern Italy and Greece), the epidemic pattern (epidemics have been reported in the Amazon Basin of Venezuela), and the occurrence of HDV infection among high-risk groups such as intravenous drug users (in developed Western countries). The epidemiology of HDV infection seems to be changing in some regions. Vaccination against HBV, decrease in HBV infection and thus in the pool of HBsAg carriers who may be infected with HDV are responsible for this decrease. Immigration patterns can be expected to have an impact on HDV infection. Nonetheless, it continues to represent a public health problem in some parts of the world yet ⁽²⁻⁵⁾.

HBV infection in dialysis patients varies among

different localities and correlates with the endemicity in the general population of the region. HBV prevalence has decreased in many countries in general population and dialysis patients. Improvement of people's knowledge about risk factors, national vaccination programs for neonates and vaccination of high risk groups are responsible for these decrease in prevalence of HBV infection in general population ⁽⁶⁾. The overall incidence and prevalence of HBV infection in dialysis patients has decreased over the years as a result of routine screening of blood products for HBsAg, the advent of recombinant human erythropoietin, HBV vaccination and the implementation of infection control measures ^(7, 8). However, the prevalence and incidence rates of HBsAg positivity are still high among patients undergoing maintenance hemodialysis in the less developed countries ⁽⁹⁾.

Transmission of HDV is similar to HBV, via blood and blood fluids containing the virus, and infection occurs by parenteral routes ⁽¹⁰⁾. Dialysis patients may acquire HDV infection as they are at risk of hepatitis B infection. The information on the epidemiology of HDV infection in the dialysis patients is limited. This may be in part related to limited use or availability of delta testing. In European countries, such as France, Sweden, and the USA, HDV infection is restricted to high risk group of drug addicts and has decreased during recent years ⁽¹¹⁾. There are some reports of acute and fulminant hepatitis or symptom-free transmission in dialysis patients with HBV and HDV infections ⁽¹²⁻¹⁴⁾. The prevalence was different from zero to 44.5% in

hemodialysis patients (15, 16). The prevalence of HDV in different groups is related to routes of transmission. In Iran, the main route for HBV transmission was vertical in past (6, 17) and the difference between the prevalence of HDV infection in hemodialysis patients (44.5%) and asymptomatic carriers (2.5%) is meaningful (15).

Delta virus is of particular potential concern in hemodialysis units where segregation of HBsAg positive patients to minimize hepatitis B transmission to susceptible patients may facilitate the transmission of delta agent (18). HDV infection is not important in developed countries, but may be a major risk for fulminant hepatitis in hemodialysis patients with HBV infection in developing countries. The importance of HDV is rising due to immigration phenomenon from area of developing countries. I recommend periodic testing for HDV infection by anti-HDV antibody in HBsAg positive carriers on chronic hemodialysis treatment.

References

1. Taylor JM. Genetic organization and replication strategy of hepatitis delta virus. *Semin Virol* 1993; **4**: 313-7.
2. Hadziyannis SJ. Review: hepatitis delta. *J Gastroenterol Hepatol* 1997; **12**: 289-98.
3. London WT, Evans AA. The epidemiology of hepatitis viruses B, C, and D. *Clin Lab Med* 1996; **16**: 251-71.
4. Ponzetto A, Forzani B, Parravicini PP, Hele C, Zanetti A, Rizzetto M. Epidemiology of hepatitis delta virus (HDV) infection. *Eur J Epidemiol* 1985; **1**: 257-63.
5. Rizzetto M. Delta virus hepatitis. *Adv Exp Med Biol* 1989; **257**: 205-9.
6. Alavian S, M, Fallahian F, Bagheri-Lankarani K. The Changing Epidemiology of Viral Hepatitis B in Iran. *J Gastrointest Liver Dis* 2007; **16**: 403-6.
7. Wong PN, Fung TT, Mak SK, Lo KY, Tong GM, Wong Y, et al. Hepatitis B virus infection in dialysis patients. *J Gastroenterol Hepatol* 2005; **20**: 1641-51.
8. Alavian SM, Mahdavi-Mazdeh M, Bagheri-Lankarani K. Hepatitis B and C in dialysis units in Iran, Changing the epidemiology. *Hemodial Int* 2008 (In Press).
9. Fabrizi F, Martin P, Lunghi G, Ponticelli C. Novel evidence on hepatitis B virus infection in dialysis. *Int J Artif Organs* 2001; **24**: 8-16.
10. Rizzetto M, Verme G. Delta hepatitis-present status. *J Hepatol* 1985; **1**: 187-93.
11. Roggendorf M, Gmelin K, Zoulek G, Wolf P, Schlipkoter U, Jilg W, et al. Epidemiological studies on the prevalence of hepatitis Delta virus infections in the Federal Republic of Germany. *J Hepatol* 1986; **2**: 230-6.
12. Marinucci G, Di Giacomo C, Orchi N, Iannicelli G, Ferrazzi M, De Paolis P, et al. HBV and HDV infection in chronic hemodialysis treatment patients (in Italian). *Riv Eur Sci Med Farmacol* 1987; **9**: 313-6.
13. Kharsa G, Degott C, Degos F, Carnot F, Potent F, Kreis H. Fulminant hepatitis in renal transplant recipients. The role of the delta agent. *Transplantation* 1987; **44**: 221-3.
14. Gmelin K, Roggendorf M, Schlipkoter U. Delta infection in a hemodialyzed patient. *J Infect Dis* 1985; **151**: 374.
15. Rezvan H, Ahmadi J, Farhadi M. A preliminary study on the prevalence of anti-HCV amongst healthy blood donors in Iran. *Vox Sang* 1994; **67**: A100.
16. Pol S, Dubois F, Mattlinger B, Carnot F, Legendre C, Brechot C, et al. Absence of hepatitis delta virus infection in chronic hemodialysis and kidney transplant patients in France. *Transplantation* 1992; **54**: 1096-7.
17. Alavian SM, Mostajabi P, Malekzadeh R, Azimi K, Vosough H, Sarrafi M, et al. Evaluation of Hepatitis B Transmission Risk Factors in Tehran Blood Donors (in Persian). *Govareh* 2004; **3**: 169-75.
18. Fabrizi F, Lunghi G, Martin P. Epidemiology of hepatitis delta virus (HDV) infection in the dialysis population. *Int J Artif Organ* 2002; **25**: 8-17.