Original Article

Shima Soleimani Amiri (MD)¹ Mohammad Jafar Soleimani Amiri (PhD)² Mohammad Reza Hasanjani Roushan (MD)^{3*}

 Pathology Resident, Shahid Beheshti Hospital, Babol University of Medical Sciences, Babol, Iran.
 Razi Laboratory Medicine, Babol, Iran.
 Infectious Diseases & Tropical Medicine Research Center, Babol University of Medical Sciences, Babol, Iran.

* Correspondence:

Mohammad Reza Hasanjani Roushan, Infectious Diseases & Tropical Medicine Research Center, Babol University of Medical Sciences, Babol, Iran. **Post Code:** 46917-14141 **E-mail:** hagar2q@yahoo.ca **Tel:** +98-111-2207924 **Fax:** +98-1112207918

 Received:
 18 March 2010

 Revised:
 10 Apr 2010

 Accepted:
 30 May 2010

Possible sources for transmission of Hepatitis B Virus Infection in 80 Children in Babol, North of Iran

Abstract

Background: To determine the possible sources for the transmission of Hepatitis B Virus (HBV) infection to children in Babol, North of Iran.

Methods: Forty-five boys and 35 girls with the mean age of 9.2 ± 5.1 years were evaluated from 1993 to 2004. Hepatitis B viral markers in all the family members were assessed. The risk factors for other sources were also determined.

Results: Sixty-five (81.3%) of them were HBeAg positive. Chronic HBV infection was found in the family members of 53 (66.2%) infected children. Chronic HBV infection was seen in 27 (33.8%) cases without any evidence of chronic HBV infection in their family members. Chronic HBV infected mothers were the most probable source of infection in 54.3% of the girls and in 24.4% of the boys (p=0.006). There was no evidence of chronic HBV infected girls (p=0.002).

Conclusion: The results of this study show that more than one-third of children acquired *HBV* infection in the society.

Key words: Hepatitis B virus, Transmission, Children, Iran.

Casp J Intern Med 2010; 1(2): 50-52.

A pproximately, 5% of the world's population has chronic HBV infection, which is the leading cause of chronic hepatitis, cirrhosis, and hepatocellular carcinoma (1). HBV is transmitted by percutaneous or permucosal exposure to infectious body fluids, sexual contact with an infected person, and perinatally from an infected mother to her infant (1, 2). The frequency of HBV infection and patterns of HBV transmission vary markedly in different parts of the world (3-7). Five to 20% of infants born from HBsAg-positive and HBeAg negative mothers become infected at birth. Infants of HBsAg-positive women who are not infected at birth are at increasing risk of HBV infection during early childhood because of household contact with infected persons (8). Intra-household horizontal personto-person transmission is an important route of HBV transmission in some parts of the world (9). Children under four years of age seemed to become carriers more easily than older children (10). Strategies for the control of hepatitis B virus (HBV) infection rely on the information about the modes of its spread and the numbers of individuals who are at risk in particular community subgroups. The aim of the present study was to determine the possible routes of transmission of Hepatitis B Virus to children in Babol, Iran.

Methods

This prospective study was performed on 80 children <15 years with chronic HBV infection at the Department of Infectious Diseases, Babol University Medical Sciences, from 1993 to 2004. Children with previous history of Hepatitis B vaccination were excluded from this study.

Most of these cases were diagnosed during the screening of children of chronic HBV infected parents. The purpose of the study was explained and the written informed consent for the study was obtained from the parents of these children. A 5 ml blood sample was taken from each of these children born before 1993 and tested in Elisa for HBsAg, HBeAg, HBsAb, (HBsAg, from Bio Merieux, the Netherlands; anti-HBs from Radim Italy, HBeAg from Dia.Pro Diagnostic BioProbes, Italy). In Iran, the universal neonatal vaccination against HBV started in 1993 according to WHO recommendations. HBV markers in all family members of these children were assessed and the probable sources for acquiring of the infection in these children were determined. Categorical variables were tested by the two-tailed Chi-square test and Fisher exact test.

Results

Fourty-five boys and 35 girls with the mean age of 9.2 ± 5.1 years were evaluated. The mean age of the girls was lower than the boys. Sixty-five (81.3%) of them were HBeAg positive. Chronic HBV infection was found in the family members of 53 (66.2%) infected children. Chronic HBV infection was seen in 27 (33.8%) cases without any evidence of chronic HBV infection in their family members. Chronic HBV infected mothers were the most probable source of infection in 54.3% of the girls and in 24.4% of the boys (p=0.006).

There was not any evidence of chronic HBV infection in the family members of 48.9% of the infected boys and in 14.5% of the infected girls (p=0.002). Boys acquired HBV infection more often than the girl in the society (48.3% versus 14.3%, p=0.002). The most probable sources for the transmission of HBV are shown in table 1.

Table 1. Possible source for HBV transmission to 80children in Babol, northern Iran

Possible source for	Boy	Girl	Total
HBV transmission	No (%)	No (%)	
Mother	11(36.7)	19 (63.3)	30 (100)
Father	5(41.7)	7 (58.3)	12 (100)
Brother or sister	7(77.8)	2 (22.2)	9 (100)
Both father and mother	0(0)	2 (100)	2 (100)
Society	22(81.5)	5 (18.5)	27 (100)
Total	45(56.3)	35 (43.7)	80 (100)

Discussion

In this study, the most probable source for HBV transmission in 80 HBV infected children were from their mother (37.5%), father (15%), and the society (33.8%)(table 1). HBsAg-positive mothers who are anti-HBe positive transfer the infection from 5 to 20 percent to their infants while the infection transfer occurs in 70 to 90% infants of HBs and HBeAg positive mothers if not given immunoprophylaxis (11).

A study from Singapore showed that the prevalence of HBV infection in children of infected mothers and fathers was almost equal, 48% and 44%, respectively (12). Our results are consistent with the results of a study from Saudi Arabia showing that the prevalence of HBV infection in siblings of families where the mother is a carrier is higher than the families where the father is the carrier (13). In that study, 17.9% of siblings were carriers and 34.8% had a resolved HBV infection. A number of studies have reported that mother–to-child transmission is a main route of HBV transmission (14-19).

In this study, most of these mothers were HBsAg and anti-HBe positive. Consequently, HBV infection likely occurred only in a small percentage of children during the prenatal period and most of the children were infected during early childhood. In one study, all babies born from carrier mothers became HBsAg positive at one year of age (20). Several studies from other countries where the prevalence of HBV is considered intermediate, reported relatively high transmission rates from infected mothers or fathers to their children (21-24).

However, one study in the Middle East showed that 21% of the children born from HBsAg positive mothers became HBV carriers. One factor accounting for this may be the low prevalence of Hepatitis B e antigen in HBsAg positive mothers in this region (14). In conclusion, the results of this study show that more than one-third of children acquired HBV infection in the society.

Acknowledgement

We would like to thank the personnel of the Department of Infectious Diseases for helping us in this study and Dr. Mahmood Hajiahmadi in the statistical analysis of the data.

References

- Mahoney FJ. Update on Diagnosis, Management, and Prevention of hepatitis B virus infection. Clin Microbiol 1999; 357-66.
- Hoofnagle J. Chronic hepatitis B (editorial) New Engl J Med 1990; 323: 337-8.
- 3.Hsu SC, Chang MH, Ni YH, Hsu HY, Lee CY. Horizontal transmission of hepatitis B virus in children. J Pediatr Gastroentrol Nutr 1993; 17: 346-7.
- Bisharat N, Elias M, Raz R, Flatau E. Familial pattern of infection with hepatitis B virus among Ethiopian Jews in Israel. Eur J Epidemiol 1998; 14: 89-91.
- Ni YH, Chang MH, Huang LM, et al. Hepatitis B virus infection in children and adolescents in a hyperendemic area: 15 years after mass hepatitis B vaccination. Ann Intern Med 2001; 135: 835-6.
- Aweis D, Brabin BJ, Beeching NJ, et al. Hepatitis B prevalence and risk factors for HBsAg carriage amongst Somali households in Liverpool. Common Dis Public Health 2001; 4: 247-52.
- Kashiwagi S, Hayashi J, Nomura H, Ikematsu H, Kajiyama W. Large-scale survey of hepatitis B virus infection in families. Microbiol Immunol 1985; 29: 951-8.
- Beasley RP, Hwang LY. Postnatal infectivity of hepatitis B surface antigen-carrier mothers. J Infect Dis 1983; 147:185-90.
- Abdool Karim SS, Thejpal R, Coovadia HM. Household clustering and intra-household transmission patterns of hepatitis B virus infection in South Africa. Int J Epidemiol 1991; 20:495-503.
- Kashiwagi S, Hayashi J, Ikematsu H, et al. Transmission of hepatitis B virus among siblings. Am Epidemiol 1984; 120: 617-25.
- Koziel MJ, Siddiqui A. Hepatitis B virus and hepatitis B virus. In: Mandell GL, Bennett JE, Dolin E, eds. Principles and Practice of Infectious diseases.6th edn. Philadelphia: Chuchill Livingstone, 2005:1864-90
- Tan CC, Guan R, Yap I, Tay NH, Kang JY. Horizontal or vertical transmission of hepatitis B virus. A serological survey in family members of hepatitis B carriers in Singapore. Trans R Soc Trop Med Hyg 1991; 85; 656-9.

- Ramia S. Intrafamilial clustering of hepatitis B virus (HBV) infection: study of 10 Saudi families. Ann Trop Med Parasitol 1990; 84: 623-7.
- Toukan A. Strategy for the control of hepatitis B virus infection in the Middle East and North Africa. The Middle East regional study group. Vaccine 1990; 8: S117-21.
- Hurie MB, Mast EE, Davis JP. Horizontal transmission of hepatitis B virus infection to United State-born children of Hmong refugees. Pediatrics 1992; 89: 269-73.
- 16.Soltani MS, Bchir A, Slimane MN, et al. Mother-child transmission of hepatitis B virus in the Tunisian Sahel. Rev Epidemiol Sante Publique 1994; 42: 529-32.
- William SJ, Craig PI, and Liddle C, et al. Hepatitis B in Australia: determinants of intrafamily spread. Aust N Z J Med 1987; 17: 220-7.
- Kashiwagi S, Hayashi J, Nomura H, et al. Changing pattern of intrafamilial transmission of hepatitis B virus in Okinawa, Japan. Am J Epidemiol 1988; 127: 783-7.
- Lok AS, Lai CL, Wu PC, et al. Hepatitis B virus infection in Chinese families in Hong Kong.Am J Epidemiol 1987; 126: 492-9.
- 20. Kotkat A, Farghaly AG, el-Zaiadi AR. Intrafamilial spread of asymptomatic hepatitis B in Alexandria. J Egypt public Health Assoc 1990; 63: 401-10.
- 21.Stevens CE, Neurath RA, Beasley RP, Szmuness W. HBeAg and anti-HBe detection with radio immunoassay: correlation with vertical transmission of hepatitis B virus in Taiwan. J Med virol 1979; 3: 237-41.
- 22. Xu ZY, Liu CB, Francis DB, et al. Prevention of perinatal acquisition of hepatitis B virus carriage using vaccine: preliminary report of a randomized doubleblind placebo-controlled and comparative trial. Pediatrics 1985; 76:713-8.
- 23. Beasley RP, Hwang LY. Postnatal infectivity of hepatitis B surface antigen-carrier mothers. J I Dis 1983; 147:185-90.
- Ordog K, Szendroi A, Szarka K, et al. Perinatal and intrafamily transmission of hepatitis B virus in three generations of a low- prevalence. J Med Virol 2003; 70:194-204.

This document was created with Win2PDF available at http://www.daneprairie.com. The unregistered version of Win2PDF is for evaluation or non-commercial use only.