Hepatitis B: transmission and natural history

R P Perrillo

Abstract

This paper reviews the transmission and natural history of hepatitis B viral infection. Means of transmission are compared by geographical region, and the association with risk factors is described. Long term outcome and overall survival are documented, and the need for universal screening programmes is discussed.

(Gut 1993; supplement: S48–S49)

Gastroenterology Section, Veterans Administration Medical Center, Washington University, St Louis, USA R P Perrillo

Correspondence to: Dr R P Perrillo, Veterans Administration Medical Center (111JC), 915 North Grand Boulevard, St Louis, Missouri 63106, USA.

Epidemiology and transmission

The prevalence of hepatitis B virus (HBV) infection is highest in developing countries in Asia, Africa, and the Pacific Islands and lowest in developed countries in North America, western Europe, and Australasia (Table). Methods of transmission vary geographically

TABLE Worldwide distribution of hepatitis B virus infection¹

	Endemic status		
	Low	Intermediate	High
Prevalence;			
Chronic infection	<2%	2-7%	8-15%
Total infection	<20%	2060%	>60%
Distribution	North America	Eastern Europe	Southeast Asia
		Southern Europe	China
	Western Europe	Soviet Union	Philippines
	-	Central Asia	Indonesia
	Australia	Japan	Middle East
	New Zealand	Israel	Africa
	South America	South America	Amazon Basin
	(southern)	(northern)	Pacific Islands
			Arctic (Eskimo)

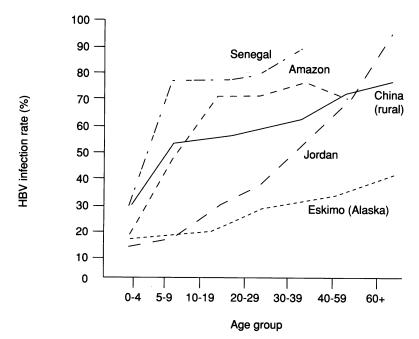


Figure Age specific seroprevalence of hepatitis B virus (HBV) infection in selected populations with a high endemicity of infection. \(^1\)

and are generally related to the incidence of infection. In areas of high HBV incidence, transmission is usually vertical from infected mother to child, or horizontal within families (Figure). In intermediate areas of prevalence, HBV is spread horizontally, with the highest rate of infection occurring among older children, adolescents, and adults. In areas of low prevalence, HBV is primarily a disease of adolescents and young adults and is transmitted sexually or parenterally.

HBV transmission through exchange of body fluids and blood contact, occurs in the following exposure settings: (a) perinatal infection (pregnancy or childbirth); (b) sexual contact (homosexual or heterosexual); (c) inadvertent transmission of contaminated blood via exchange of blood products (for example, transfusion, dialysis), shared needles in intravenous drug abusers, and needlestick type and other open wound injuries (for example, accidents, violence).

Nearly 300 000 cases of acute HBV infection occur in the United States each year. While most of these are short lived, chronic infection occurs in approximately 2-5% of instances. Hepatitis B is seldom a fulminant disease, but when this occurs mortality rates are high unless liver transplantation is done. The epidemiology of acute HBV infection in the United States has changed over the past decade, with a major decline observed in male homosexuals and health care workers and an increase in heterosexually promiscuous and intravenous drug user populations. Irrespective of the exposure setting, recipients of HBV antigen (HBeAg) reactive donors are three to fourfold more likely to develop infection. The risk of persistent HBV infection in children is inversely related to the age of the child at initial infection. Therefore, the greatest impact on prevention of hepatitis B will result from universal screening of pregnant women and immunoprophylaxis of exposed infants, and the integration of HBV vaccine into current childhood immunisation schedules.

Long term outcome

Worldwide, it has been estimated that over 250 000 people die annually from HBV associated acute and chronic liver disease. In a study correlating outcome with the initial histological diagnosis, the estimated five year survival rate was 97% for chronic persistent hepatitis, 86% for chronic active hepatitis, and 55% for chronic active hepatitis with cirrhosis. Recently, investigators in the Netherlands have described an overall five year survival of 14% in patients with decompensated chronic

hepatitis B.3 Prospective studies in Asia have shown that HBV carriers have at least a 200 fold greater risk of developing hepatocellular carcinoma than those who do not carry the virus.4 Moreover, it has been estimated that the lifetime risk of death from hepatocellular carcinoma or cirrhosis, or both is between 40 and 50% for men, and the risk of death from liver disease is approximately 15% for women. The exact frequency of transformation to cirrhosis is not known, but this generally occurs insidiously and without a noticeable change in symptoms. Several studies have shown that even mild forms of histological injury, such as chronic persistent hepatitis, can progress to

chronic active hepatitis and cirrhosis in as many as one third of patients when viral replication is sustained.

- 1 Margolis HS, Alter MJ, Hadler SC. Hepatitis B: Evolving epidemiology and implications for control Sem Liver Dis 1991; 11: 84-92.
- 1991; 11: 84-92.
 Weissberg J, Andres LL, Smith CI, et al. An analysis of 379 patients. Ann Intern Med 1984; 101: 613-16.
 de Jongh FE, Janssen HLA, de Man RA, Hop WCJ, Schalm SW, van Blankenstein M. Survival and prognostic indicators in HBsAg-positive cirrhosis of the liver. The role of HBeAg seroconversion. Gastroenterology 1992; 103: 1630-5. 103: 1630-5
- 4 Beasley RP, Hwang LY. Epidemiology of hepatocellular carcinoma. In: Vyas GN, Dienstag JL, Hoofnagle JH, eds. Viral hepatitis and liver disease. New York: Grune and Stratton, 1983: 209-24.