

Anti-pre-S antibodies in different groups of patients with hepatitis B virus infection

<https://www.researchgate.net/publication/20665080>

DOI: 10.1111/j.1440-1746.1989.tb00803.x · Source: PubMed

Cited in:

Journal of Gastroenterology and Hepatology 4(1):25-32, 1989

Anti-pre-S antibodies in different groups of patients with hepatitis B virus infection

Mohammad Irshad, B. M. Gandhi, Subrat Acharya, B N Tandon Department of Gastroenterology, All India Institute of Medical Sciences

Abstract

The anti-pre-S antibody in the samples of sera from normal healthy persons and patients with different clinical types of liver diseases due to hepatitis B virus (HBV) infection was detected by a newly established enzyme-linked immunosorbent assay technique. This test is a blocking assay where anti-pre-S antibody in the patient's serum blocks subsequent addition of horse radish peroxidase-labelled polymerized human serum albumin (pHSA) to the pHSA-receptor site of HBsAg molecules fixed on a solid surface. Anti-pre-S activity was not detected in any from 95 healthy persons who were negative for all HBV-markers or from 105 healthy HBV carriers. In 12 sera from HBV vaccine recipients, anti-pre-S activity was noted in higher proportions compared with anti-HBs, after both the second and third doses of vaccine. Anti-pre-S activity was detected in small proportions of HBsAg positive sera from acute viral hepatitis (4.2%) and chronic active hepatitis (10%). In subacute viral hepatitis patients, the anti-pre-S antibody was totally absent. However, anti-pre-S activity was recorded in high proportions of HBsAg-positive sera from patients with cirrhosis of liver (57.2%) and fulminant hepatitis (41.6%). The anti-pre-S antibodies were assumed to be implicated in the clearance of HBV particles from circulation without causing tissue damage.

Reference

M Irshad, B M Gandhi, S K Acharya, Y K Joshi, B N Tandon, An enzyme-linked immunosorbent assay (ELISA) for the detection of IgG and IgM anti-idiotypes directed against anti-HBs molecules. Journal of Immunological Methods 96(2):211-7, 1987