Anti-Hepatitis C Virus (HCV) core protein antibody, monoclonal (H6-29)

65-052  100 ug

Hepatitis C virus (HCV) is a small (55-65 nm in size), enveloped, positive sense single-stranded RNA virus in the family Flaviviridae and the principal cause of parenteral non-A, non-B hepatitis. The virus genome consists of a single open reading frame of approximately 9.4 kb which encodes a single polyprotein of about 3,010 amino acids (1, 2, 3). The polyprotein is processed by host cell and viral proteases into four structural proteins (core, envelope 1 and 2, and p7) and six non-structural proteins (NS2, 3, 4a, 4b, 5a, and 5b) necessary for viral replication. HCV core protein is not only a component of nucleocapsid but also has multiple functions and is thought to be a pathogenic factor for hepatitis. It also participates in some cellular processes, including transcriptional regulation and cellular transduction. HCV core antigen is used as diagnostic marker for HCV infection.

Applications

1. Western blotting (1/1,000~1/2,000 dilution)
2. Immunohistochemistry (1/100~1/500 dilution)
3. Immunofluorescence staining (1/100~1/500 dilution)
4. ELISA (assay dependent)

Immunogen: A part of the core region (nucleotides 369-704, amino acids 13-124) of HCV genotype 1b expressed in E. coli (the nucleotide sequence is shown in ref.3)

Isotype: Mouse IgG2a kappa

Form: Purified monoclonal antibody (IgG) 1mg/ml in PBS, 50% glycerol, filter-sterilized

Specificity: Specific to human HCV core antigen of genotype 1b. Not tested in other genotypes

Storage: -20°C

Data Link: Swiss Prot HCV protein

References: This antibody is used in ref.4 and 5.


Related products: #65-057 anti-HCV NS4a antibody  #65-062 anti-HCV NS5a antibody  #65-067 anti-HCV NS5b antibody

to be continued...
Fig. 1  Western blotting of HCV core protein.
Chimp liver cells were infected with recombinant vaccinia virus containing a HCV genome cDNA and were subjected to Western blotting using this antibody. The core protein is detected as a 22-kDa band.

Fig. 2  Detection of HCV core protein by immunofluorescence antibody staining.
Chimp liver cells were infected with recombinant vaccinia virus containing a HCV genome cDNA. After incubation for 48 hr, the cells were fixed with acetone and HCV core protein was detected by indirect immunofluorescence staining using this antibody.

Fig. 3  Immunohistochemical detection of HCV core protein.
Tissue section from a patient with chronic hepatitis C was immunostained to reveal cells expressing HCV core antigen, which are scattered in the lobules (indirect immuno-histochemical method, counterstained with Mayer’s hematoxylin).