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Product Information

HCV Core Antigen recombinant, expressed in *E. coli*

Catalog Number **H9034** Storage Temperature –20 °C

Product Description

This product is the hepatitis C virus (HCV) core antigen, amino acids 2–192 of the HCV polyprotein, which has been expressed in *E. coli* as a β -galactosidase tagged protein. The 2–192 region represents the viral precursor polyprotein and HCV core nucleocapsid immunodominant region. The HCV core protein is co-translationally inserted into the endoplasmic reticulum membrane.

The HCV core protein is highly basic and acts as a carrier/chaperone for RNA.^{1,2} It contains three major domains with particular functions:²

- The N-terminal hydrophilic domain of ~120 amino acids (domain D1), a highly basic region with many positively charged amino acids that is involved mainly in RNA binding
- A hydrophobic domain of ~50 amino acids (domain D2), involved in core association with endoplasmic reticulum membranes and with lipid droplets in mammalian cells, and core folding and oligomerization³
- A signal peptide, containing the final 20 amino acids, for the downstream protein E1

This product of the HCV core antigen fused to the N-terminal of β -galactosidase is a construct of ~136 kDa. It reacts positively with human HCV serum and may be used as a positive control for HCV antibodies in various immunoassay procedures.

This product is supplied as a solution at a concentration of ~1 mg/ml in 20 mM Tris-HCl, pH 8.0, containing 8 M urea and 10 mM β -mercaptoethanol.

Purity: ≥95% (SDS-PAGE)

Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

Storage/Stability

This product ships on dry ice and is stable at -20 °C for at least a year. Upon thawing, store at 2-4 °C for one month or as frozen aliquots at -20 °C. Avoid repeated freeze-thaw cycles.

References

- Cristofari, G. et al., Nucleic Acids Res., 32(8), 2623-2531 (2004).
- 2. Boulant, S. *et al.*, *J. Virol.*, **79(17)**, 11353-11365 (2005).
- 3. Kunkel, M., and Watowich, S.J., *FEBS Lett.*, **557(1-3)**, 174-180 (2004).

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