antibodies -online.com







anti-HCV 1b NS5B antibody (AA 92-105)

Images

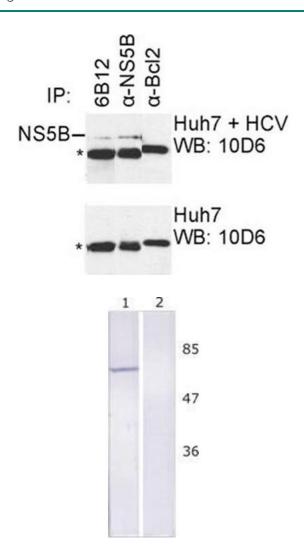


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| 100 μg | |
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| HCV 1b NS5B (HCV NS5b) | |
| AA 92-105 | |
| Hepatitis C Virus (HCV) | |
| Mouse | |
| Monoclonal | |
| This HCV 1b NS5B antibody is un-conjugated | |
| Western Blotting (WB), ELISA, Immunoprecipitation (IP) | |
| | |
| Recombinant Hepatitis C Virus subtype 1b nonstructuralprotein 5B (NS5B) RNA-dependent | |
| RNA polymerase (RdRp) | |
| TPPHSARSKF GYGA | |
| 6B12 | |
| IgG1 | |
| Human, HCV subtype 1b NS5B, | |
| purified by protein G affinity chromatography | |
| | |
| HCV 1b NS5B (HCV NS5b) | |
| | |

Target Details

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| Abstract: | HCV NS5b Products | |
| Target Type: | Viral Protein | |
| Background: | Non-structural protein 5B (NS5B) represents the RNA-dependent RNA polymerase (RdRp) of Hepatitis C Virus, which is a small positive strand RNA virus in the family Flaviviridae. HCV is a major causative agent of acute and chronic hepatitis, hepatocellular carcinoma and liver cirrhosis. The single subunit RNA-dependent RNA polymerase is absolutely essential for the viral replication. | |
| UniProt: | Q9WMX2 | |
| Application Details | | |
| Application Notes: | ELISA titer 1:1000, Western immunoblotting, reduced conditions, dilution 1:1000, Immunoprecipitation. Optimal conditions for IP should be determined for each particular application. | |
| Restrictions: | For Research Use only | |
| Handling | | |
| Format: | Liquid | |
| Concentration: | 1 mg/ml | |
| Buffer: | PBS pH 7.4, with 0.1% | |
| Preservative: | Sodium azide | |
| Precaution of Use: | This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. | |
| Handling Advice: | Divide antibodyinto aliquots prior usage. Avoid multiple freeze-thaw cycles as productdegradation may result. | |
| Storage: | -20 °C | |
| Storage Comment: | This product is shipped in non-frozen liquid form in ambient conditions | |



Western Blotting

Image 1. IP was carried out with NS5B specific mAb 6B12 using the lysates of Huh7 cells harboring selectable subgenomic HCV RNA replicon (upper panel) or plain Huh7 cells (lower panel). NS5B polyclonal antibodies (alpha-NS5B) and alpha-Bcl2 mAb, directed against cellular protein, were used as positive and negative controls respectively. Asterisk indicates immunoglobulin heavy chain; protein blots were probed with NS5B specific mAb 10D6.

Western Blotting

Image 2. WB analysis of Huh-7 cell line expressing NS5B protein. Line 1. Huh-7 transfected with HCV replicon, Line 2. Huh-7 non-transfected cells.