Package Insert



Mouse monoclonal antibody to S-Adenosylhomocysteine (Clone 301-3) (50µl)

Catalog Number:

SAH13-A50

For Research Use Only. Not for use in diagnostic procedures. v. 1.0

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Properties

- Form: Liquid
- Storage instructions: Store at 4°C, -20°C for long term storage
- **Storage buffer:** PBS 10mM pH 7.4 (NaCl 150mM), Sodium azide 0.02%, BSA 10mg/ml or PBS 10mM pH 7.4 (NaCl 150mM), Sodium azide 0.02%, Glycerol 50%, BSA 10mg/ml
- Purity: >95% Purified from mouse ascites fluid by affinity chromatography
- Clonality: Monoclonal
- Clone number: 301-3
- Immunoglobin isotype: IgG3
- Affinity: Ka = 8.32 x 10⁸L/mol (1.20 x 10⁻⁹M)
- **Specificity:** Shows the following reactivities with related compounds: SAdenosylhomocysteine: 100%, S-Adenosylmethionine: < 3%, Adenosine: <1%, Homocysteine: <1%,L-Cysteine: <1%, Glutathione: <1%, L-Cystathionine: <1%.
- Immunogen: S-Adenosylhomocysteine conjugated to BSA

Applications

The application notes include recommended starting dilutions. Optimal dilutions/concentrations should be determined by the end user. Higher dilution than suggested maybe used in IHC and IF. The product may be used in other not-yet-tested applications.

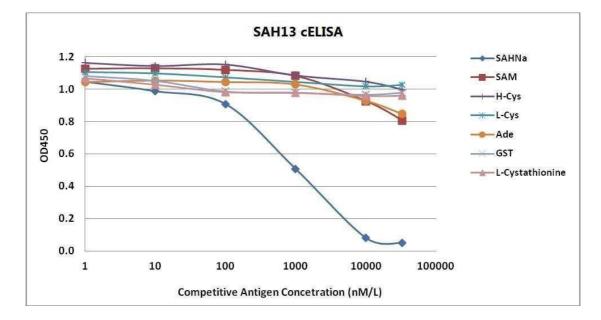
Application	Notes
cELISA	1:4000/8000
FCM	1:200
ІНС	1:200

Target

S-adenosylhomocysteine is a competitive inhibitor of S-adenosylmethionine-dependant methyl transferase reactions. Therefore, it plays a key role in the control of methylation via regulation of the intracellular concentration of S-adenosylhomocysteine.

Cellular localization Cytoplasm, nuclear





Anti-Adenosylhomocysteine antibody [301-3]

Figure 1 Competitive ELISA using anti-S-Adenosymethionine monoclonal antibody [301-3]

The 0.5 µg/ml of SAH-BSA was coated into 96 wells. Serial dilution of SAH standard (SAHNa), SAdenosylmethionine, Homocysteine (H-Cys), L-Cysteine (L-Cys), Adenosine (Ade), Glutathione (GST), L- Cystathionine (L-CTT) and properly diluted MA00303 were added. HRP conjugated Goat anti-Mouse IgG antibody was used to develop the color. OD450 value was measured on each well.

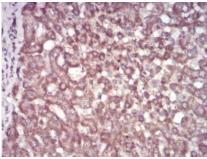


Figure 2 Immunohistochemistry staining was performed using the antibody with benign liver tissue adjacent to carcinoma. Brown areas indicated strong positive staining in nuclear and cytoplasmic areas.

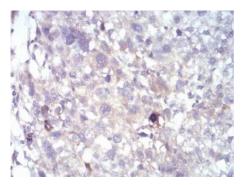


Figure 3 The same samples as in Figure 2 from liver cancer area. Cytoplasmic and nuclear areas showed negative staining.

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