A Reliable Research Partner in Life Science and Medicine

HMGCS2 Polyclonal Antibody

catalog number: E-AB-64961

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human; Mouse; Rat

Immunogen Recombinant fusion protein of human HMGCS2 (NP 005509.1).

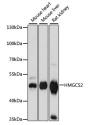
Host Rabbit Isotype IgG

Purification Affinity purification

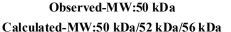
Buffer Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

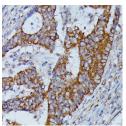
Applications Recommended Dilution 1:500-1:2000 WB 1:50-1:100 IHC IF 1:50-1:100

Data

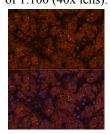


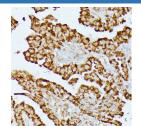
Western blot analysis of extracts of various cell lines using HMGCS2 Polyclonal Antibody at dilution of 1:1000.



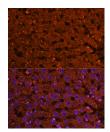


Immunohistochemistry of paraffin-embedded Human colon carcinoma using HMGCS2 Polyclonal Antibody at dilution of 1:100 (40x lens).

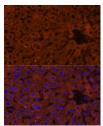




Immunohistochemistry of paraffin-embedded Human thyroid cancer using HMGCS2 Polyclonal Antibody at dilution of 1:100 (40x lens).



Immunofluorescence analysis of Rat liver using HMGCS2 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



For Research Use Only

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Immunofluorescence analysis of Human liver using HMGCS2 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

Immunofluorescence analysis of Mouse liver using HMGCS2 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

Preparation & Storage

Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.

Shipping The product is shipped with ice pack,upon receipt, store it immediately at the

temperature recommended.

Background

The protein encoded by this gene belongs to the HMG-CoA synthase family. It is a mitochondrial enzyme that catalyzes the first reaction of ketogenesis, a metabolic pathway that provides lipid-derived energy for various organs during times of carbohydrate deprivation, such as fasting. Mutations in this gene are associated with HMG-CoA synthase deficiency. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

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