A Reliable Research Partner in Life Science and Medicine

(KO Validated) NSDHL Polyclonal Antibody

Catalog No. E-AB-91649

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

Description

Reactivity Human, Mouse, Rat

Immunogen Recombinant fusion protein of human NSDHL

Host Rabbit Isotype IgG

Purification Affinity purification

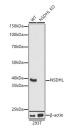
Conjugation Unconjugated

Buffer PBS with 0.01% thiomersal,50% glycerol,pH7.3.

Applications Recommended Dilution

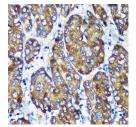
WB 1:500-1:2000 IHC 1:50-1:200 IF 1:50-1:200

Data

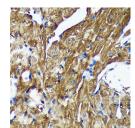


Western blot analysis of extracts from wild type(WT) and NSDHL knockout (KO) 293T(KO) cells, using NSDHL Polyclonal Antibody at 1:1000 dilution.

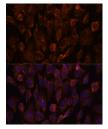
Observed Mw:40KDa Calculated Mw:41kDa



Immunohistochemistry of paraffin-embedded Human liver cancer using [KO Validated] NSDHL Polyclonal Antibody at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.



Immunohistochemistry of paraffin-embedded Mouse heart using [KO Validated] NSDHL Polyclonal Antibody at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.



Immunofluorescence analysis of C6 cells using NSDHL Polyclonal Antibody at dilution of 1:100. Blue: DAPI for nuclear staining.

For Research Use Only

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017

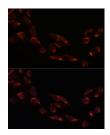
Email: techsupport@elabscience.com

Web: www.elabscience.com

Elabscience Bionovation Inc.



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Immunofluorescence analysis of U2OS cells using NSDHL Polyclonal Antibody at dilution of 1:100. Blue: DAPI for nuclear staining.

Preparation & Storage

Storage

Store at -20°C. Avoid freeze/thaw cycles.

Background

The protein encoded by this gene is localized in the endoplasmic reticulum and is involved in cholesterol biosynthesis. Mutations in this gene are associated with CHILD syndrome, which is a X-linked dominant disorder of lipid metabolism with disturbed cholesterol biosynthesis, and typically lethal in males. Alternatively spliced transcript variants with differing 5' UTR have been found for this gene.

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