Leader in Biomolecular Solutions for Life Science



Catalog No.: A25585 Recombinant



Basic Information

Observed MW 113kDa

Calculated MW 113kDa

Category Primary antibody

Applications ELISA,WB,IHC-P

Cross-Reactivity Human, Mouse, Rat

CloneNo number ARC66832

Recommended Dilutions

Background

Degradation of glycine is brought about by the glycine cleavage system, which is composed of four mitochondrial protein components: P protein (a pyridoxal phosphate-dependent glycine decarboxylase), H protein (a lipoic acid-containing protein), T protein (a tetrahydrofolate-requiring enzyme), and L protein (a lipoamide dehydrogenase). The protein encoded by this gene is the P protein, which binds to glycine and enables the methylamine group from glycine to be transferred to the T protein. Defects in this gene are a cause of nonketotic hyperglycinemia (NKH).

Immunogen Information

WB	1:1000 - 1:5000	Gene ID	Swiss Prot	
IHC-P	1:50 - 1:200	2731	P23378	
		Immunegen		

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 580-749 of human GLDC (NP_000161.2).

Synonyms

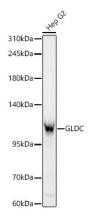
GCE; GCSP; HYGN1

Product Information

Source Rabbit **lsotype** IgG Purification Affinity purification

Storage

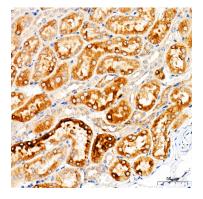
Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.



Western blot analysis of lysates from Hep G2 cells using GLDC Rabbit mAb (A25585) at 1:2000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 5s.



Immunohistochemistry analysis of GLDC in paraffin-embedded Rat liver tissue using GLDC Rabbit mAb (A25585) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of GLDC in paraffin-embedded Mouse kidney tissue using GLDC Rabbit mAb (A25585) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.