

GLUD1 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP11701b

Product Information

Application	WB, IHC-P, E
Primary Accession	<u>P00367</u>
Other Accession	<u>P49448</u> , <u>P10860</u> , <u>P26443</u> , <u>P00368</u> , <u>P00366</u> , <u>NP_005262.1</u>
Reactivity	Human, Mouse
Predicted	Bovine, Chicken, Rat
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Clone Names	RB20073
Calculated MW	61398
Antigen Region	438-465

Additional Information

Gene ID	2746
Other Names	Glutamate dehydrogenase 1, mitochondrial, GDH 1, GLUD1, GLUD
Target/Specificity	This GLUD1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 438-465 amino acids from the C-terminal region of human GLUD1.
Dilution	WB~~1:1000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.
Format	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	GLUD1 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	GLUD1
Synonyms	GLUD
Function	Mitochondrial glutamate dehydrogenase that catalyzes the conversion of

L-glutamate into alpha-ketoglutarate. Plays a key role in glutamine
anaplerosis by producing alpha-ketoglutarate, an important intermediate in
the tricarboxylic acid cycle (PubMed:<u>11032875</u>, PubMed:<u>11254391</u>,
PubMed:<u>16023112</u>, PubMed:<u>16959573</u>). Plays a role in insulin homeostasis
(PubMed:<u>11297618</u>, PubMed:<u>9571255</u>). May be involved in learning and
memory reactions by increasing the turnover of the excitatory
neurotransmitter glutamate (By similarity).Cellular LocationMitochondrion. Endoplasmic reticulum. Note=Mostly translocates into the
mitochondria, only a small amount of the protein localizes to the endoplasmic
reticulum.

Background

This gene encodes glutamate dehydrogenase protein; a mitochondrial matrix enzyme that catalyzes the oxidative deamination of glutamate to alpha-ketoglutarate and ammonia. This enzyme has an important role in regulating amino acid induced insulin secretion and activating mutations in this gene are a common cause of congenital hyperinsulinism. This enzyme is allosterically activated by ADP and inhibited by GTP and ATP. The related glutamate dehydrogenase 2 gene on the human X-chromosome originated from this gene via retrotransposition and encodes a soluble form of glutamate dehydrogenase. Multiple pseudogenes of this gene are present in humans.

References

Martins-de-Souza, D., et al. J Psychiatr Res 44(14):989-991(2010) Jia, P., et al. Schizophr. Res. 122 (1-3), 38-42 (2010) : Joslyn, G., et al. Alcohol. Clin. Exp. Res. 34(5):800-812(2010) Flanagan, S.E., et al. Eur. J. Endocrinol. 162(5):987-992(2010) Bao, X., et al. J. Neurosci. 29(44):13929-13944(2009)

Images





GLUD1 Antibody (C-term) (Cat.

#AP11701b)immunohistochemistry analysis in formalin fixed and paraffin embedded human liver tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.This data demonstrates the use of GLUD1 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.

Citations

- Antrodia cinnamomea Inhibits Migration in Human Hepatocellular Carcinoma Cells.
- NOX1 Supports the Metabolic Remodeling of HepG2 Cells.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.