



## DHE3 rabbit pAb

Catalog No	BYab-11722
Isotype	IgG
Reactivity	Human; Mouse;Rat
Applications	WB
Gene Name	GLUD1 GLUD
Protein Name	DHE3
Immunogen	Synthesized peptide derived from human DHE3
Specificity	This antibody detects endogenous levels of DHE3 at Human/Mouse/Rat
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
Dilution	WB 1: 500-2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	
Observed Band	
Cell Pathway	Mitochondrion . Endoplasmic reticulum . Mostly translocates into the mitochondria, only a small amount of the protein localizes to the endoplasmic reticulum
Tissue Specificity	
Function	catalytic activity:L-glutamate + H(2)O + NAD(P)(+) = 2-oxoglutarate + NH(3) +

NAD(P)H., disease Defects in GLUD1 are the cause of

NAD(P)H.,disease:Defects in GLUD1 are the cause of hyperinsulinism-hyperammonemia syndrome (HHS) [MIM:606762]. Elevated oxidation rate of glutamate to alpha-ketoglutarate stimulates insulin secretion in the pancreatic beta cells, while they impair detoxification of ammonium in the liver.,enzyme regulation:Subject to allosteric regulation. Activated by ADP. Inhibited by GTP and ATP. ADP can occupy the NADH binding site and activate the enzyme.,function:May be involved in learning and memory reactions by increasing the turnover of the excitatory neurotransmitter glutamate.,online information:Glutamate dehydrogenase 1 entry,similarity:Belongs to the Glu/Leu/Phe/Val dehydrogenases family.,subunit:Homohexamer.,

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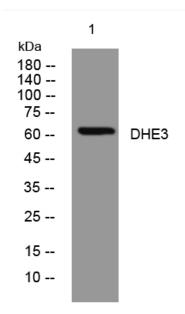


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Background	This gene encodes glutamate dehydrogenase, which is a mitochondrial matrix enzyme that catalyzes the oxidative deamination of glutamate to alpha-ketoglutarate and ammonia. This enzyme has an important role in regulating amino acid-induced insulin secretion. It is allosterically activated by ADP and inhibited by GTP and ATP. Activating mutations in this gene are a common cause of congenital hyperinsulinism. Alternative splicing of this gene results in multiple transcript variants. 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. Related pseudogenes have been identified on chromosomes 10, 18 and X. [provided by RefSeq, Jan 2016],
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## **Products Images**



Western blot analysis of lysates from MDA-MB cells, primary antibody was diluted at 1:1000, 4° over night

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