



PDHA1/2 (Phospho-Ser293/291) rabbit pAb

Catalog No	YP-Ab-10469
Isotype	IgG
Reactivity	Human; Mouse; Rat
Applications	WB
Gene Name	PDHA1 PHE1A
Immunogen	Synthesized peptide derived from human PDHA1/2 (Phospho-Ser293/291)
Specificity	This antibody detects endogenous levels of PDHA1/2 (Phospho-Ser293/291) at Human, Mouse, Rat
Formulation	Liquid in PBS containing 50% glycerol, and 0.124% 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	ODPA_HUMAN; PDH; PDHA; PDHA1; PDHCE1A; PDHE1 A type I; PDHE1-A type I; PHE1A; Pyruvate Dehydrogenase (lipoamide) alpha 1; Pyruvate dehydrogenase complex; E1 alpha polypeptide 1; Pyruvate Dehydrogenase E1 alpha; Pyruvate dehydrogenase E1 component subunit alpha; somatic form; mitochondrial; EC 1.2.4.1; MGC149517; MGC149518; ODPAT_HUMAN; Pdha2; PDHAL; PDHE1 A type II; PDHE1-A type II; Pyruvate dehydrogenase (lipoamide) alpha 2; testis specific form; Pyruvate dehydrogenase; E1 alpha polypeptide; testis spec; testis-specific form; PDHA1/2; PDHA1 PHE1A; PDHA1/2 (Phospho-Ser293/291); mitochondrial (EC 1.2.4.1) (PDHE1-A type I)
Observed Band	43kD
Calculated Molecular Weight	43kD
Cell Pathway	Mitochondrion matrix.
Tissue Specificity	Ubiquitous.
Function	catalytic activity: Pyruvate + [dihydrolipoyllysine-residue acetyltransferase] lipoyllysine = [dihydrolipoyllysine-residue acetyltransferase] S-acetyldihydrolipoyllysine + CO(2)., cofactor: Thiamine pyrophosphate., disease: Defects in PDHA1 are a cause of pyruvate decarboxylase E1



component deficiency (PDHE1 deficiency) [MIM:312170]. PDHE1 deficiency is the most common enzyme defect in patients with primary lactic acidosis. It is associated with variable clinical phenotypes ranging from neonatal death to prolonged survival complicated by developmental delay, seizures, ataxia, apnea, and in some cases to an X-linked form of Leigh syndrome (LS) (Leigh encephalomyelopathy).,disease:Defects in PDHA1 are the cause of X-linked Leigh syndrome (LS) [MIM:308930]. LS is an early-onset progressive neurodegenerative disorder with a characteristic neuropathology consisting of focal, bilateral lesions in o

Background

The pyruvate dehydrogenase (PDH) complex is a nuclear-encoded mitochondrial multienzyme complex that catalyzes the overall conversion of pyruvate to acetyl-CoA and CO(2), and provides the primary link between glycolysis and the tricarboxylic acid (TCA) cycle. The PDH complex is composed of multiple copies of three enzymatic components: pyruvate dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and lipoamide dehydrogenase (E3). The E1 enzyme is a heterotetramer of two alpha and two beta subunits. This gene encodes the E1 alpha 1 subunit containing the E1 active site, and plays a key role in the function of the PDH complex. Mutations in this gene are associated with pyruvate dehydrogenase E1-alpha deficiency and X-linked Leigh syndrome. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Mar 2010],

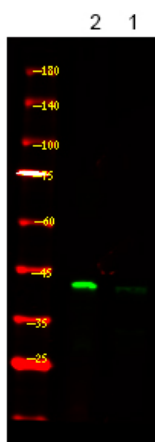
matters needing attention

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 1 HepG2 cell 2, LPS 100ng/mL 30min treated ,using primary antibody at 1:1000 dilution. Secondary antibody(catalog#:RS23920) was diluted at 1:10000