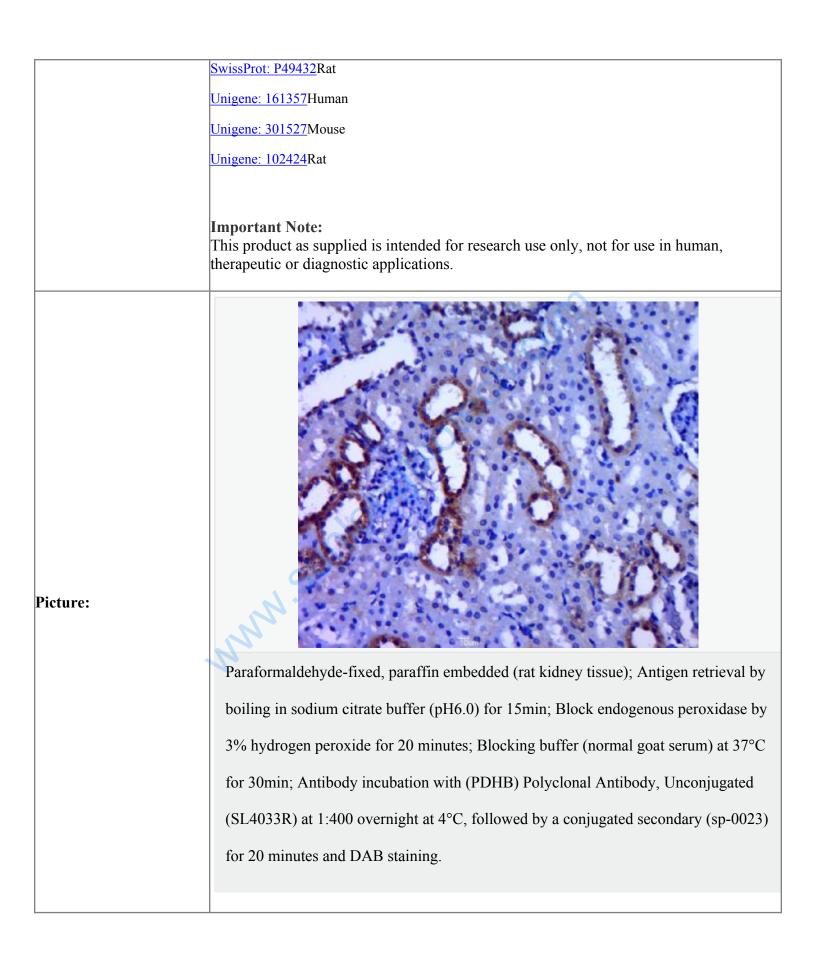


Rabbit Anti-PDHB antibody

SL4033R

PDHB U
丙酮酸脱氢酶E1β亚单位抗体
PDHB; PDHE1 B; PHE1B; Pyruvate dehydrogenase (lipoamide) beta; Pyruvate dehydrogenase E1 beta polypeptide; Pyruvate dehydrogenase E1 component subunit beta; Pyruvate dehydrogenase E1 component subunit beta mitochondrial; DKFZp564K0164; ODPB_HUMAN.
Specific References(1) SL4033R has been referenced in 1 publications.
[IF=2.08]Xing, Wen Min, et al. "Proteomic identification of mitochondrial targets
involved in andrographolide sodium bisulfite-induced nephrotoxicity in a rat
model."Environmental Toxicology and Pharmacology (2015).WB;Rat.
PubMed:26356389
Rabbit
Polyclonal
Human, Mouse, Rat, Dog, Cow, Horse, Rabbit,
WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=1:100- 500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
35kDa
cytoplasmic
Lyophilized or Liquid
1mg/ml
KLH conjugated synthetic peptide derived from human PDHB:51-150/359
IgG
affinity purified by Protein A
0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized

PubMed:	antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C. PubMed
Product Detail:	The pyruvate dehydrogenase complex catalyzes the overall conversion of pyruvate to acetyl-CoA and CO2. It contains multiple copies of three enzymatic components: pyruvate dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and lipoamide dehydrogenase (E3). Function: The pyruvate dehydrogenase complex catalyzes the overall conversion of pyruvate to acetyl-CoA and CO(2). It contains multiple copies of three enzymatic components: pyruvate dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and lipoamide dehydrogenase (E3). Subuni: Tetramer of 2 alpha and 2 beta subunits. Subcellular Location: Mitochondrion matrix. DISEASE: Defects in PDHB are the cause of pyruvate dehydrogenase E1-beta deficiency (PDHBD) [MIM:614111]. An enzymatic defect causing primary lactic acidosis in children. It is associated with a broad elinical spectrum ranging from fatal lactic acidosis in the newborn to chronic neurologic dysfunction with structural abnormalities in the central nervous system without systemic acidosis. SWISS: P11177 Gene ID: S162 Database links: Entrez Gene: 5162Human Entrez Gene: 289950Rat Omim: 179060Human SwissProt: P11177Human SwissProt: P02051Mouse



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