

Rabbit Anti-LDHA antibody

SL1810R

Product Name:	LDHA U
Chinese Name:	乳酸脱氢酶抗体
Alias:	Lactate Dehydrogenase; Lactate Dehydrogenase Isoenzyme V; Lactate Dehydrogenase isozyme H4; L lactate dehydrogenase A chain; Lactate dehydrogenase A; Lactate dehydrogenase A chain; LDH A; LDH heart subunit; LDH M; LDHM; LDH muscle subunit; LDH1; LDHA; PIG 19; PIG19; Proliferation inducing gene 19 protein; TRG 5; TRG5; LDHA_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Dog, Horse,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow- Cyt=0.2µg /testIF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	37kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human LDHA:261-361/361
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized 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:	PubMed
Product Detail:	The protein encoded by this gene catalyzes the conversion of L-lactate and NAD to pyruvate and NADH in the final step of anaerobic glycolysis. The protein is found predominantly in muscle tissue and belongs to the lactate dehydrogenase family.

Mutations in this gene have been linked to exertional myoglobinuria. Multiple transcript variants encoding different isoforms have been found for this gene. The human genome contains several non-transcribed pseudogenes of this gene. [provided by RefSeq].

Subunit:

Homotetramer.

Subcellular Location: Cytoplasm.

Post-translational modifications: ISGylated.

DISEASE:

Glycogen storage disease 11 (GSD11) [MIM:612933]: A metabolic disorder that results in exertional myoglobinuria, pain, cramps and easy fatigue. Note=The disease is caused by mutations affecting the gene represented in this entry.

Similarity: Belongs to the LDH/MDH superfamily. LDH family.

SWISS: P00338

Gene ID: 3939

Database links:

Entrez Gene: 3939Human

Entrez Gene: 16828 Mouse

Entrez Gene: 24533Rat

Omim: 150000Human

SwissProt: P00338Human

SwissProt: P06151Mouse

SwissProt: P04642Rat

Unigene: 2795Human

Unigene: 29324Mouse

Unigene: 107896Rat



