

Rabbit Anti-Transaldolase 1 antibody

SL4041R

Product Name:	Transaldolase 1
Chinese Name:	转醛醇酶/EPS8L2抗体
Alias:	Dihydroxyacetone transferase; EPS8L2; Glycerone transferase; TAL; TAL H; TALDO; TALDO1; TALDOR; TALH; Transaldolase 1; Transaldolase1; Transaldolase-1; EC 2.2.1.2; TALDO_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Dog, Cow, Horse, Sheep,
Applications:	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.
Molecular weight:	37kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	lmg/ml
immunogen:	KLH conjugated synthetic peptide derived from human Transaldolase 1:145-250/337
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:	<u>PubMed</u>
Product Detail:	Transaldolase 1 is a key enzyme of the nonoxidative pentose phosphate pathway providing ribose-5-phosphate for nucleic acid synthesis and NADPH for lipid biosynthesis. This pathway can also maintain glutathione at a reduced state and thus protect sulfhydryl groups and cellular integrity from oxygen radicals. The functional gene of transaldolase 1 is located on chromosome 11 and a pseudogene is identified on

chromosome 1 but there are conflicting map locations. The second and third exon of this gene were developed by insertion of a retrotransposable element. This gene is thought to be involved in multiple sclerosis.

Function:

Transaldolase is important for the balance of metabolites in the pentose-phosphate pathway.

Subunit:

Homodimer.

DISEASE:

Transaldolase 1 deficiency (TALDO1 deficiency) [MIM:606003]: Results in telangiectases of the skin, hepatosplenomegaly, and enlarged clitoris. Note=The disease is caused by mutations affecting the gene represented in this entry.

SWISS:

Q9H6S3

Gene ID:

64787

Database links:

Entrez Gene: 64787 Human

Entrez Gene: 98845 Mouse

Omim: 614988 Human

SwissProt: Q9H6S3 Human

SwissProt: Q99K30 Mouse

Unigene: 55016 Human

Unigene: 27451 Mouse

Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

Transaldolase 1称醛羧移转酶

或转二羟丙酮基酶, 经研究该蛋白与多发性硬化症有关。