

LIPT1 Rabbit pAb

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Product Information

Application IHC-P, IHC-F, IF

Primary Accession <u>Q9Y234</u>

Reactivity Rat, Mouse, Rabbit, Dog, Horse

Host Rabbit
Clonality Polyclonal
Calculated MW 42479
Physical State Liquid

Immunogen KLH conjugated synthetic peptide derived from human LIPT1

Epitope Specificity 101-200/373

Isotype IgG

Purity affinity purified by Protein A

Buffer 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

SUBCELLULAR LOCATION Mitochondrion.

SIMILARITY Belongs to the IpIA family.

Important Note This product as supplied is intended for research use only, not for use in

human, therapeutic or diagnostic applications.

Background Descriptions The process of transferring lipoic acid to proteins is a two-step process. The

first step is the activation of lipoic acid by lipoate-activating enzyme to form lipoyl-AMP. For the second step, the protein encoded by this gene transfers the lipoyl moiety to apoproteins. Alternative splicing results in multiple

transcript variants. A related pseudogene has been identified on chromosome 13. Read-through transcription also exists between this gene and the

neighboring downstream mitochondrial ribosomal protein L30 (MRPL30)

gene. [provided by RefSeq, Mar 2011]

Additional Information

Gene ID 51601

Other Names Lipoyl amidotransferase LIPT1, mitochondrial, 2.3.1.200, Lipoate biosynthesis

protein, Lipoate-protein ligase, Lipoyl ligase, Lipoyltransferase 1, 2.3.1.-, LIPT1

(HGNC:29569)

Target/Specificity Highly expressed in skeletal muscle and heart, moderately in kidney and

pancreas, and detected at lower levels in liver, brain, placenta and lung.

Dilution IHC-P=1:100-500,IHC-F=1:100-500,ICC/IF=1:100-500,IF=1:100-500

Storage Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. 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.

Protein Information

Name LIPT1 (HGNC:29569)

Function Lipoyl amidotransferase that catalyzes the transfer of lipoyl moieties from

lipoyl-protein H of the glycine cleavage system (lipoyl- GCSH) to E2 subunits of the pyruvate dehydrogenase complex (PDCE2) (PubMed: 29987032). Unable to

catalyze the transfer of octanoyl from octanoyl-GCSH to PDCE2

(PubMed:<u>29987032</u>). In vitro, it is also able to catalyze the transfer of the lipoyl group from lipoyl-AMP to the specific lysine residue of lipoyl domains of lipoate-dependent enzymes but this reaction may not be physiologically

relevant (Probable).

Cellular Location Mitochondrion {ECO:0000250 | UniProtKB:O46419}.

Tissue Location Highly expressed in skeletal muscle and heart, moderately in kidney and

pancreas, and detected at lower levels in liver, brain, placenta and lung.

Background

The process of transferring lipoic acid to proteins is a two-step process. The first step is the activation of lipoic acid by lipoate-activating enzyme to form lipoyl-AMP. For the second step, the protein encoded by this gene transfers the lipoyl moiety to apoproteins. Alternative splicing results in multiple transcript variants. A related pseudogene has been identified on chromosome 13. Read-through transcription also exists between this gene and the neighboring downstream mitochondrial ribosomal protein L30 (MRPL30) gene. [provided by RefSeq, Mar 2011]

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.