

Rabbit Anti-SLC27A4 antibody

SL11535R

Product Name:	SLC27A4
Chinese Name:	长链 脂肪酸 Transporter4 抗体
Alias:	ACSVL 4; ACSVL-4; ACSVL4; EC 6.2.1; FATP 4; FATP4; Fatty acid transport protein 4; Fatty acid transport protein 4; IPS; Long chain fatty acid transport protein 4; Long chain fatty acid transport protein 4; OTTHUMP00000022264; S27A4; SLC27 A4; SLC27A 4; Solute carrier family 27 (fatty acid transporter) member 4; Solute carrier family 27 member 4; Solute carrier family 27 member 4; S27A4 HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Chicken, Pig, Cow, Horse,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow-Cyt=1μg/TestICC=1:100-500IF=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:	72kDa
Cellular localization:	The cell membrane
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human SLC27A4:288-360/643
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:	This gene encodes a member of a family of fatty acid transport proteins, which are involved in translocation of long-chain fatty acids cross the plasma membrane. This protein is expressed at high levels on the apical side of mature enterocytes in the small

intestine, and appears to be the principal fatty acid transporter in enterocytes. Clinical studies suggest this gene as a candidate gene for the insulin resistance syndrome. Mutations in this gene have been associated with ichthyosis prematurity syndrome. [provided by RefSeq, Apr 2010]

Function:

nvolved in translocation of long-chain fatty acids (LFCA) across the plasma membrane. Appears to be the principal fatty acid transporter in small intestinal enterocytes. Plays a role in the formation of the epidermal barrier. Required for fat absorption in early embryogenesis. Has acyl-CoA ligase activity for long-chain and very-long-chain fatty acids

Subcellular Location:

Membrane; Multi-pass membrane protein

Tissue Specificity:

Expressed at highest levels in brain, testis, colon and kidney. Expressed at medium levels in heart and liver, small intestine and stomach. Expressed at low levels in peripheral leukocytes, bone marrow, skeletal muscle and aorta. Expressed in adipose tissue.

DISEASE:

Defects in SLC27A4 are the cause of ichthyosis prematurity syndrome (IPS) [MIM:608649]. A keratinization disorder characterized by complications in the second trimester of pregnancy resulting from polyhydramnion, with premature birth of a child with thick caseous desquamating epidermis, respiratory complications and transient eosinophilia. After recovery during the first months of life, the symptoms are relatively benign and the patients suffer from a lifelong non-scaly ichthyosis with atopic manifestations.

Similarity:

Belongs to the ATP-dependent AMP-binding enzyme family.

SWISS:

O6P1M0

Gene ID:

10999

Database links:

Entrez Gene: 10999 Human

Entrez Gene: 26569 Mouse

Omim: 604194 Human

SwissProt: Q6P1M0 Human

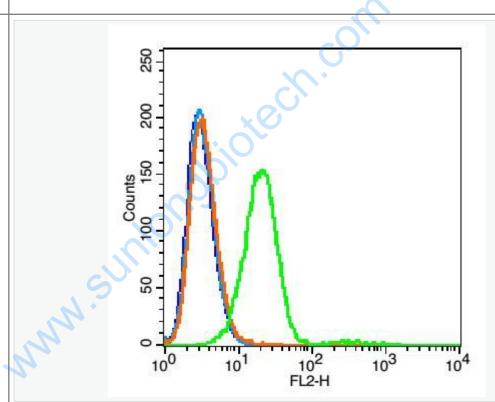
SwissProt: Q91VE0 Mouse

Unigene: 656699 Human

Unigene: 330113 Mouse

Important Note:

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



Picture:

Blank control: RSC96 cells(blue).

Primary Antibody: Rabbit Anti-SLC27A4 antibody(SL11535R), Dilution: 1µg in

100 μL 1X PBS containing 0.5% BSA;

Isotype Control Antibody: Rabbit IgG (orange) ,used under the same conditions.

Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X

PBS containing 0.5% BSA.

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