

Rabbit Anti-SLC25A38 antibody

SL19815R

Product Name:	SLC25A38
Chinese Name:	溶质载体家族蛋白25成员38抗体
Alias:	FLJ20551; FLJ22703; S2538 HUMAN; slc25a38; Solute carrier family 25 member 38.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Cow,Rabbit,Sheep,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=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:	34kDa
Cellular localization:	The cell membrane
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human SLC25A38:101-200/304
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:	This gene is a member of the mitochondrial carrier family. The encoded protein is required during erythropoiesis and is important for the biosynthesis of heme. Mutations in this gene are the cause of autosomal congenital sideroblastic anemia.[provided by RefSeq, Mar 2010]
	Function: Mitochondrial carrier required during erythropoiesis. Probably involved in the

biosynthesis of heme, possibly by facilitating 5-aminolevulinate (ALA) production. May act by importing glycine into mitochondria or by exchanging glycine for ALA across the mitochondrial inner membrane.

Subcellular Location: Mitochondrion inner membrane.

Tissue Specificity: Preferentially expressed in erythroid cells.

DISEASE:

Defects in SLC25A38 are a cause of anemia sideroblastic pyridoxine-refractory autosomal recessive (PRARSA) [MIM:205950]. A form of sideroblastic anemia not responsive to pyridoxine. Sideroblastic anemia is characterized by anemia of varying severity, hypochromic peripheral erythrocytes, systemic iron overload secondary to chronic ineffective erythropoiesis, and the presence of bone marrow ringed sideroblasts. Sideroblasts are characterized by iron-loaded mitochondria clustered around the nucleus.

Similarity:

Belongs to the mitochondrial carrier family. SLC25A38 subfamily. Contains 3 Solcar repeats.

SWISS: Q96DW6

Gene ID: 54977

Database links:

Entrez Gene: 512325 Cow

Entrez Gene: 54977 Human

Entrez Gene: 208638 Mouse

Entrez Gene: 301067 Rat

Entrez Gene: 100147784 Sheep

<u>Omim: 610819</u> Human

SwissProt: Q5EAC0 Cow

<u>SwissProt: Q96DW6</u> Human

	SwissProt: Q91XD8 Mouse
	<u>SwissProt: Q499U1</u> Rat
	SwissProt: S2538 Sheep
	Unigene: 369615 Human
	Unigene: 236656 Mouse
	<u>Unigene: 21048</u> Rat
	Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
Picture:	100 100 75 63 48 35 SLC25A38 25 20 17 17 100 100 50 50 50 50 50 50 50 50 50
	Sample: bone (Mouse) Lysate at 40 ug
	Primary: Anti-SLC25A38(SL19815R)at 1/300 dilution
	Secondary: IRDye800CW Goat Anti-RabbitIgG at 1/20000 dilution
	Predicted band size: 34kD

