

Rabbit Anti-ZNF379 antibody

SL6200R

Product Name:	ZNF379
Chinese Name:	Zinc finger protein379抗体
Alias:	CXorf11; DHHC9; Palmitoyltransferase ZDHHC9; ZDHHC 9; ZDHHC10; Zinc finger DHHC domain containing protein 9; Zinc finger protein 379; ZNF379; ZNF380; ZDHC9_HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Chicken, Dog, Pig, Cow, Horse, Rabbit,
Applications:	IHC-P=1:400-800IHC-F=1:400-800 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	40kDa
Cellular localization:	cytoplasmicThe cell membrane
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human ZNF379/ZDHHC9:118-155/364
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 an integral membrane protein that is a member of the zinc finger DHHC domain-containing protein family. The encoded protein forms a complex with golgin subfamily A member 7 and functions as a palmitoyltransferase. This protein specifically palmitoylates HRAS and NRAS. Mutations in this gene are associated with X-linked mental retardation. Alternate splicing results in multiple transcript variants that encode the same protein. [provided by RefSeq, May 2010].

Function:

The ZDHHC9-GOLGA7 complex is a palmitoyltransferase specific for HRAS and NRAS.

Subunit:

Interacts with GOLGA7.

Subcellular Location:

Endoplasmic reticulum membrane; Multi-pass membrane protein. Golgi apparatus membrane; Multi-pass membrane protein.

Tissue Specificity:

Highly expressed in kidney, skeletal muscle, brain, lung and liver. Absent in thymus, spleen and leukocytes.

DISEASE:

Defects in ZDHHC9 are the cause of mental retardation syndromic X-linked ZDHHC9-related (MRXSZ) [MIM:300799]. A disorder characterized by significantly sub-average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. Some patients have marfanoid habitus as an additional feature.

Similarity:

Belongs to the DHHC palmitoyltransferase family. ERF2/ZDHHC9 subfamily. Contains 1 DHHC-type zinc finger.

SWISS:

Q9Y397

Gene ID:

51114

Database links:

Entrez Gene: 51114Human

Entrez Gene: 208884Mouse

Entrez Gene: 302808Rat

Omim: 300646Human

SwissProt: Q9Y397Human

SwissProt: P59268Mouse

Unigene: 193566Human

Unigene: 207367 Mouse
Important Note:
This product as supplied is intended for research use only, not for use in human,
therapeutic or diagnostic applications.

