

Rabbit Anti-SPECC1L antibody

SL7773R

Product Name:	SPECC1L SPECC1
Chinese Name:	细胞质和纺锤体机化蛋白A抗体
Alias:	Cytokinesis and spindle organization A; Cytospin-A; CYTSA; CYTSA_HUMAN; Renal carcinoma antigen NY-REN-22; SPECC1-like protein; specc1l; Sperm antigen with calponin homology and coiled-coil domains 1-like.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Chicken, Dog, Pig, Cow, Horse, Rabbit, 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:	125kDa
Cellular localization:	cytoplasmicThe cell membrane
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human SPECC1L:161-260/1117
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 coiled-coil domain containing protein. The encoded protein may play a critical role in actin-cytoskeletal reorganization during facial morphogenesis. Mutations in this gene are a cause of oblique facial clefting-1. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. A readthrough transcript composed of SPECC1L (sperm antigen with calponin homology and

coiled-coil domains 1-like) and the downstream ADORA2A (adenosine A2a receptor) gene sequence has been identified, but it is thought to be non-coding. [provided by RefSeq, Jun 2013]

Function:

Involved in cytokinesis and spindle organization. May play a role in actin cytoskeleton organization and microtubule stabilization and hence required for proper cell adhesion and migration.

Subunit:

May interact with both microtubules and actin cytoskeleton.

Subcellular Location:

Cytoplasm, cytoskeleton. Cytoplasm, cytoskeleton, spindle. Cell junction, gap junction. Note=Colocalizes with acetylated alpha-tubulin, gamma-tubulin and F-actin. Also observed in a ring around gamma-tubulin containing centrioles possibly in the microtubule organizing center.

DISEASE:

Defects in SPECC1L are the cause of facial clefting oblique type 1 (OBLFC1) [MIM:600251]. A rare form of facial clefting. A facial cleft is any of the fissures between the embryonic prominences that normally unite to form the face.

Similarity:

Belongs to the cytospin-A family.

Contains 1 CH (calponin-homology) domain.

SWISS:

Q69YQ0

Gene ID:

23384

Database links:

Entrez Gene: 23384 Human

Entrez Gene: 74392 Mouse

SwissProt: Q69YQ0 Human

SwissProt: Q2KN98 Mouse

Unigene: 474384 Human

	Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
Picture:	245— 180— 135— —————————————————————————————————
	Sample: HL60(Human) Cell Lysate at 30 ug
	NIH/3T3(Mouse) Cell Lysate at 30 ug
	U2OS(Human) Cell Lysate at 30 ug
	HL60 (Human) Cell Lysate at 30 ug
	Primary: Anti- SPECC1L (SL7773R) at 1/1000 dilution
	Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution
	Predicted band size: 125 kD

Observed band size: 125 kD

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