

Rabbit Anti-Synaptophysin antibody

SL23504R

Product Name:	Synaptophysin
Chinese Name:	突触小泡蛋白P38抗体
Alias:	Major synaptic vesicle protein P38; MRXSYP; Syn p38; Synaptophysin; SYP; SYPH; SYPH_HUMAN; SypI.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,
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:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human Synaptophysin :201- 300/313 <cytoplasmic></cytoplasmic>
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 encodes an integral membrane protein of small synaptic vesicles in brain and endocrine cells. The protein also binds cholesterol and is thought to direct targeting of vesicle-associated membrane protein 2 (synaptobrevin) to intracellular compartments. Mutations in this gene are associated with X-linked mental retardation (XLMR). [provided by RefSeq, Aug 2011]

Function:

Possibly involved in structural functions as organizing other membrane components or in targeting the vesicles to the plasma membrane. Involved in the regulation of short-term and long-term synaptic plasticity.

Subcellular Location:

Cytoplasmic vesicle > secretory vesicle > synaptic vesicle membrane. Cell junction > synapse > synaptosome.

Tissue Specificity:

Characteristic of a type of small (30-80 nm) neurosecretory vesicles, including presynaptic vesicles, but also vesicles of various neuroendocrine cells of both neuronal and epithelial phenotype.

Post-translational modifications:

Ubiquitinated; mediated by SIAH1 or SIAH2 and leading to its subsequent proteasomal degradation.

DISEASE:

Defects in SYP are the cause of mental retardation X-linked SYP-related (MRXSYP) [MIM:300802]. Mental retardation is characterized by significantly below average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period.

Similarity:

Belongs to the synaptophysin/synaptobrevin family. Contains 1 MARVEL domain.

SWISS: P08247

Gene ID: 6855

Database links:

Entrez Gene: 280937 Cow

Entrez Gene: 6855 Human

Entrez Gene: 20977 Mouse

Entrez Gene: 24804 Rat

<u>Omim: 313475</u> Human

SwissProt: P20488 Cow
<u>SwissProt: P08247</u> Human
SwissProt: Q62277 Mouse
SwissProt: P07825 Rat
Unigene: 632804 Human
Unigene: 223674 Mouse
Unigene: 11067 Rat
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