



RTN2 Monoclonal Antibody

Catalog No	BYmab-07319
lsotype	IgG
Reactivity	Human;Mouse
Applications	WB
Gene Name	RTN2 NSPL1
Protein Name	Reticulon-2 (Neuroendocrine-specific protein-like 1) (NSP-like protein 1) (Neuroendocrine-specific protein-like I) (NSP-like protein I) (NSPLI)
Immunogen	Synthesized peptide derived from human protein . at AA range: 391-440
Specificity	RTN2 Monoclonal Antibody detects endogenous levels of protein.
Formulation	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	
Observed Band	59kD
Cell Pathway	Endoplasmic reticulum membrane ; Multi-pass membrane protein . Sarcoplasmic reticulum membrane ; Multi-pass membrane protein . Cell membrane ; Multi-pass membrane protein . Cell membrane protein . Cell membrane, sarcolemma ; Multi-pass membrane protein . Cell membrane, sarcolemma, T-tubule ; Multi-pass membrane protein . Cytoplasm, myofibril, sarcomere, Z line . Cytoplasm, cytoskeleton . Localizes to intermediate filaments in mononucleated myoblasts and to Z lines in mature myotubes.
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Tissue Specificity	[Isoform RTN2-C]: Highly expressed in skeletal muscle.
Tissue Specificity Function	miscellaneous: The sequence shown here is derived from an EMBL/GenBank/DDBJ third party annotation (TPA) entry., similarity: Contains 1 reticulon domain., tissue specificity: Isoform RTN2-C is highly expressed in skeletal muscle.,
	miscellaneous:The sequence shown here is derived from an EMBL/GenBank/DDBJ third party annotation (TPA) entry.,similarity:Contains 1 reticulon domain.,tissue specificity:Isoform RTN2-C is highly expressed in skeletal

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Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.	
matters needing attention	Avoid repeated freezing and thawing!	
	role in intracellular vesicular transport. Alternatively spliced transcript variants encoding different isoforms have been identified. Mutations at this locus have been associated with autosomal dominant spastic paraplegia-12. [provided by RefSeq, Apr 2012],	

Nanjing BYabscience technology Co.,Ltd