



CSTN2 mouse mAb

Catalog No	BYmab-08120
Isotype	lgG
Reactivity	Human; Mouse;Rat
Applications	WB
Gene Name	CLSTN2 CS2
Protein Name	CSTN2
Immunogen	Synthesized peptide derived from human CSTN2 AA range: 494-544
Specificity	This antibody detects endogenous levels of CSTN2 at Human/Mouse/Rat
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.235% 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	Calsyntenin-2 (Alcadein-gamma) (Alc-gamma)
Observed Band	105kD
Cell Pathway	Cell membrane ; Single-pass type I membrane protein . Endoplasmic reticulum membrane . Golgi apparatus membrane . Cell junction, synapse, postsynapse . Cell projection, dendrite . Most prominent in the postsynaptic specializations of asymmetric (type I) synapses with both axodendritic and axospinous localization.
Tissue Specificity	Restricted to the brain.
Function	domain:Binds synaptic Ca(2+) with its cytoplasmic domain.,function:May modulate calcium-mediated postsynaptic signals.,PTM:Proteolytically processed under normal cellular conditions. A primary zeta-cleavage generates a large extracellular (soluble) N-terminal domain (sAlc) and a short C-terminal transmembrane fragment (CTF1). A secondary cleavage catalyzed by gamma-secretase within the transmembrane domain releases the beta-Alc-gamma chain in the extracellular milieu and produces an intracellular fragment (AlcICD). This processing is strongly suppressed in the tripartite complex formed with APBA2 and APP, which seems to prevent the association with PSEN1.,similarity:Contains 2 cadherin domains.,subcellular location:Most
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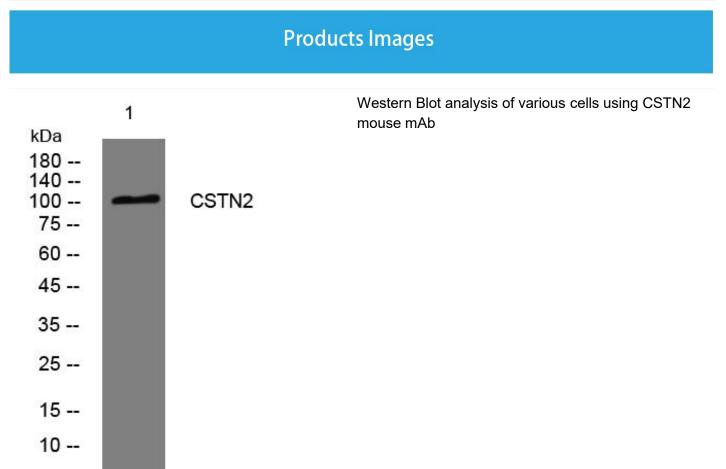


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Background	
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.



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