



MYO1A Polyclonal Antibody

Catalog No	YP-Ab-05776
Isotype	IgG
Reactivity	Human;Rat;Mouse
Applications	WB;ELISA
Gene Name	MYO1A MYHL
Immunogen	Synthesized peptide derived from human protein . at AA range: 370-450
Specificity	MYO1A Polyclonal Antibody detects endogenous levels of protein.
Formulation	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000 ELISA 1:5000-20000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	MIHC;DFNA48;Brush border myosin I;BBM-I;BBMI;MYO1A;MYO1A MYHL; Unconventional myosin-Ia (Brush border myosin I) (BBM-I) (BBMI) (Myosin I heavy chain) (MIHC)
Observed Band	114kD
Calculated Molecular Weight	114kD
Cell Pathway	cytoplasm,microvillus,brush border,basal plasma membrane,basolateral plasma membrane,apical plasma membrane,lateral plasma membrane,myosin complex, cortical actin cytoskeleton,filamentous actin,plasma membrane raft,
Tissue Specificity	Intestine,Jejunum,Placenta,
Function	disease:Defects in MYO1A are the cause of non-syndromic sensorineural deafness autosomal dominant type 48 (DFNA48) [MIM:607841]. DFNA48 is a form of sensorineural hearing loss. Sensorineural deafness results from damage to the neural receptors of the inner ear, the nerve pathways to the brain, or the area of the brain that receives sound information.,function:Involved in directing the movement of organelles along actin filaments .,similarity:Contains 1 myosin head-like domain.,similarity:Contains 3 IQ domains.,



Background

This gene encodes a member of the myosin superfamily. The protein represents an unconventional myosin; it should not be confused with the conventional skeletal muscle myosin-1 (MYH1). Unconventional myosins contain the basic domains characteristic of conventional myosins and are further distinguished from class members by their tail domains. They function as actin-based molecular motors. Mutations in this gene have been associated with autosomal dominant deafness. Alternatively spliced variants have been found for this gene. [provided by RefSeq, Dec 2011],

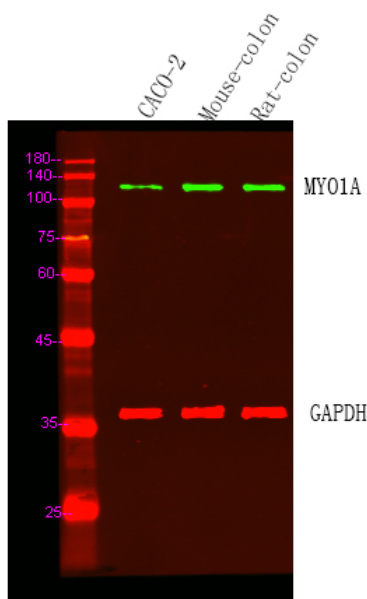
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cell lysis. Primary Antibody was diluted at 1:1000. Secondary antibody(catalog#:RS23920) was diluted at 1:10000