



MAN1B1 Monoclonal Antibody

Catalog No	YP-mAb-02666
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	MAN1B1
Immunogen	Synthesized peptide derived from MAN1B1 . at AA range: 100-180
Specificity	MAN1B1 Monoclonal Antibody detects endogenous levels of MAN1B1 protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA 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 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	EC:3.2.1.113;Endoplasmic reticulum mannosyl-oligosaccharide 1;2-alpha-mannosidase;ER alpha 1;2 mannosidase;ER alpha-1;2-mannosidase;ER mannosidase 1;MAN1B1;ERMan1;Man9GlcNAc2-specific-processing alpha-mannosidase;Mannosidase alpha class 1B member 1
Observed Band	80kD
Calculated Molecular Weight	80kD
Cell Pathway	Endoplasmic reticulum membrane ; Single-pass type II membrane protein .
Tissue Specificity	Widely expressed.
Function	catalytic activity:Hydrolysis of the terminal (1->2)-linked alpha-D-mannose residues in the oligo-mannose oligosaccharide Man(9)(GlcNAc)(2)., cofactor:Calcium.,enzyme regulation:Inhibited by both 1-deoxymannojirimycin and kifunensine.,function:Involved in the maturation of Asn-linked oligosaccharides. Trim a single alpha-1,2-linked mannose residue from Man(9) GlcNAc(2) to produce Man(8)GlcNAc(2). The only product is the Man(8) GlcNAc(2) isomer B, the form lacking the middle-arm terminal alpha 1,2-mannose. It may be involved in glycoprotein quality control since it is important to target misfolded glycoproteins for degradation.,pathway:Protein modification; protein glycosylation.,similarity:Belongs to the glycosyl hydrolase 47 family.,



tissue specificity: Widely expressed.,

Background

This gene encodes an enzyme belonging to the glycosyl hydrolase 47 family. This enzyme functions in N-glycan biosynthesis, and is a class I alpha-1,2-mannosidase that specifically converts Man9GlcNAc to Man8GlcNAc isomer B. It is required for N-glycan trimming to Man5-6GlcNAc2 in the endoplasmic-reticulum-associated degradation pathway. Mutations in this gene cause autosomal-recessive intellectual disability. Alternative splicing results in multiple transcript variants. A related pseudogene has been identified on chromosome 11. [provided by RefSeq, Dec 2011],

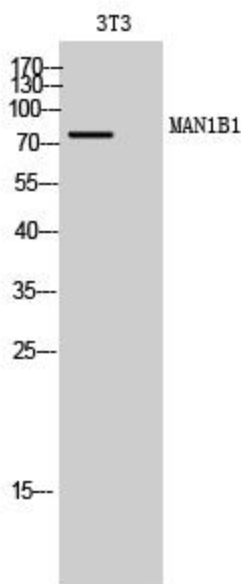
matters needing attention

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 cells using MAN1B1 Monoclonal Antibody