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PMM2 Antibody (monoclonal) (M02)

Mouse monoclonal antibody raised against a full-length recombinant PMM2.
Catalog # AT3356a

Specification

PMM2 Antibody (monoclonal) (M02) - Product Information

Application	**WB**
Primary Accession	O15305
Other Accession	BC008310
Reactivity	**Human**
Host	**mouse**
Clonality	**Monoclonal**
Isotype	**IgG2b Kappa**
Calculated MW	**28082**

PMM2 Antibody (monoclonal) (M02) - Additional Information

Gene ID 5373

Other Names
Phosphomannomutase 2, PMM 2, PMM2

Target/Specificity
PMM2 (AAH08310, 1 a.a. ~ 246 a.a) full-length recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Dilution
WB~~1:500~1000

Format
Clear, colorless solution in phosphate buffered saline, pH 7.2 .

Storage
Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Precautions
PMM2 Antibody (monoclonal) (M02) is for research use only and not for use in diagnostic or therapeutic procedures.

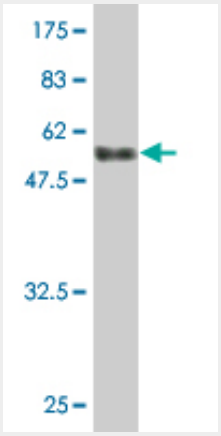
PMM2 Antibody (monoclonal) (M02) - Protocols

Provided below are standard protocols that you may find useful for product applications.

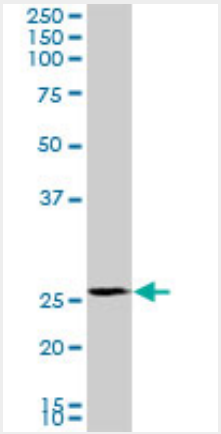
- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry

- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

PMM2 Antibody (monoclonal) (M02) - Images



Antibody Reactive Against Recombinant Protein.Western Blot detection against Immunogen (52.8 KDa) .



PMM2 monoclonal antibody (M02), clone 2A5. Western Blot analysis of PMM2 expression in K-562 ((Cat # AT3356a)

PMM2 Antibody (monoclonal) (M02) - Background

The protein encoded by this gene catalyzes the isomerization of mannose 6-phosphate to mannose 1-phosphate, which is a precursor to GDP-mannose necessary for the synthesis of dolichol-P-oligosaccharides. Mutations in this gene have been shown to cause defects in glycoprotein biosynthesis, which manifests as carbohydrate-deficient glycoprotein syndrome type I.

PMM2 Antibody (monoclonal) (M02) - References

Functional analysis of three splicing mutations identified in the PMM2 gene: toward a new therapy for congenital disorder of glycosylation type Ia. Vega AI, et al. Hum Mutat, 2009 May. PMID 19235233.Primary skeletal dysplasia as a major manifesting feature in an infant with congenital disorder of glycosylation type Ia. Coman D, et al. Am J Med Genet A, 2008 Feb 1. PMID 18203160.Cerebellar ataxia and congenital disorder of glycosylation Ia (CDG-Ia) with normal routine CDG screening. Vermeer S, et al. J Neurol, 2007 Oct. PMID 17694350.Congenital disorder of

glycosylation type 1a: three siblings with a mild neurological phenotype. Coman D, et al. J Clin Neurosci, 2007 Jul. PMID 17451957.Characterization of two unusual truncating PMM2 mutations in two CDG-Ia patients. Schollen E, et al. Mol Genet Metab, 2007 Apr. PMID 17307006.