

## Product datasheet for **TP720223L**

### **PMM2 (NM\_000303) Human Recombinant Protein**

#### **Product data:**

<b>Product Type:</b>	Recombinant Proteins
<b>Description:</b>	Recombinant protein of human phosphomannomutase 2 (PMM2)
<b>Species:</b>	Human
<b>Expression Host:</b>	E. coli
<b>Expression cDNA Clone or AA Sequence:</b>	Met1-Ser246
<b>Tag:</b>	C-His
<b>Predicted MW:</b>	29.1 kDa
<b>Concentration:</b>	lot specific
<b>Purity:</b>	>95% as determined by SDS-PAGE and Coomassie blue staining
<b>Buffer:</b>	Provided lyophilized from a 0.2 µm filtered solution of 20 mM Tris-HCl, 150 mM NaCl
<b>Endotoxin:</b>	< 0.1 EU per µg protein as determined by LAL test
<b>Storage:</b>	Store at -80°C.
<b>Stability:</b>	Stable for at least 3 months from date of receipt under proper storage and handling conditions.
<b>RefSeq:</b>	<a href="#">NP_000294</a>
<b>Locus ID:</b>	5373
<b>UniProt ID:</b>	<a href="#">O15305</a> , <a href="#">A0A0S2Z4J6</a> , <a href="#">Q59F02</a>
<b>Cytogenetics:</b>	16p13.2
<b>Synonyms:</b>	CDG1; CDG1a; CDGS; PMI; PMI1; PMM 2
<b>Summary:</b>	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. [provided by RefSeq, Jul 2008]
<b>Protein Families:</b>	Druggable Genome



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**Protein Pathways:** Amino sugar and nucleotide sugar metabolism, Fructose and mannose metabolism, Metabolic pathways

**Product images:**

