

# POMT1 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP12380b

#### **Product Information**

**Application** IHC-P, WB, FC, E

Primary Accession Q9Y6A1

Other Accession NP 001070833.1, NP 001129586.1

Reactivity Human
Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Clone Names RB31305
Calculated MW 84881
Antigen Region 706-735

#### **Additional Information**

**Gene ID** 10585

Other Names Protein O-mannosyl-transferase 1, Dolichyl-phosphate-mannose--protein

mannosyltransferase 1, POMT1

Target/Specificity This POMT1 antibody is generated from rabbits immunized with a KLH

conjugated synthetic peptide between 706-735 amino acids from the

C-terminal region of human POMT1.

**Dilution** IHC-P~1:100~500 WB~~1:1000 FC~~1:10~50 E~~Use at an assay dependent

concentration.

**Format** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is purified through a protein A column, followed by peptide

affinity purification.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** POMT1 Antibody (C-term) is for research use only and not for use in

diagnostic or therapeutic procedures.

#### **Protein Information**

Name POMT1

**Function** Transfers mannosyl residues to the hydroxyl group of serine or threonine

residues. Coexpression of both POMT1 and POMT2 is necessary for enzyme

activity, expression of either POMT1 or POMT2 alone is insufficient (PubMed:12369018, PubMed:14699049, PubMed:28512129). Essentially dedicated to O-mannosylation of alpha- DAG1 and few other proteins but not of cadherins and protocaherins (PubMed:28512129).

**Cellular Location** Endoplasmic reticulum membrane; Multi-pass membrane protein

**Tissue Location** Widely expressed. Highly expressed in testis, heart and pancreas. Detected at

lower levels in kidney, skeletal muscle, brain, placenta, lung and liver

## **Background**

The protein encoded by this gene is an O-mannosyltransferase that requires interaction with the product of the POMT2 gene for enzymatic function. The encoded protein is found in the membrane of the endoplasmic reticulum. Defects in this gene are a cause of Walker-Warburg syndrome (WWS) and limb-girdle muscular dystrophy type 2K (LGMD2K). Several transcript variants encoding different isoforms have been found for this gene.

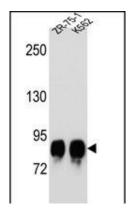
#### References

Manya, H., et al. J. Biochem. 147(3):337-344(2010) Jugessur, A., et al. PLoS ONE 5 (7), E11493 (2010): Cotarelo, R.P., et al. Clin. Genet. 76(1):108-112(2009) Mercuri, E., et al. Neurology 72(21):1802-1809(2009) Judas, M., et al. Neuropediatrics 40(1):6-14(2009)

### **Images**

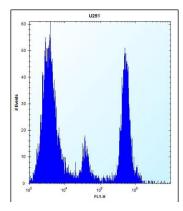


POMT1 Antibody (C-term) (Cat. #AP12380b)immunohistochemistry analysis in formalin fixed and paraffin embedded human testis tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of POMT1 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.



POMT1 Antibody (C-term) (Cat. #AP12380b) western blot analysis in ZR-75-1,K562 cell line lysates (35ug/lane). This demonstrates the POMT1 antibody detected the POMT1 protein (arrow).

POMT1 Antibody (C-term) (Cat. #AP12380b) flow cytometric analysis of U251 cells (right histogram)



compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.