

# **Product datasheet for TP318781M**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

#### PMP22 (NM\_153322) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human peripheral myelin protein 22 (PMP22), transcript variant 3, 100

μg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC218781 representing NM\_153322 or AA Sequence: Red=Cloning site Green=Tags(s)

MLLLLLSIIVLHVAVLVLLFVSTIVSQWIVGNGHATDLWQNCSTSSSGNVHHCFSSSPNEWLQSVQATMI LSIIFSILSLFLFFCQLFTLTKGGRFYITGIFQILAGLCVMSAAAIYTVRHPEWHLNSDYSYGFAYILAW

VAFPLALLSGVIYVILRKRE

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK
Predicted MW: 17.7 kDa

Concentration: >0.05 µg/µL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 696997

Locus ID: 5376

**UniProt ID:** Q01453, Q6FH25



#### PMP22 (NM\_153322) Human Recombinant Protein - TP318781M

RefSeq Size: 1669

Cytogenetics: 17p12 RefSeq ORF: 480

Synonyms: CIDP; CMT1A; CMT1E; DSS; GAS-3; GAS3; HMSNIA; HNPP; Sp110

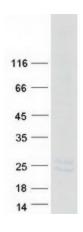
**Summary:** This gene encodes an integral membrane protein that is a major component of myelin in the

> peripheral nervous system. Studies suggest two alternately used promoters drive tissuespecific expression. Various mutations of this gene are causes of Charcot-Marie-Tooth disease Type IA, Dejerine-Sottas syndrome, and hereditary neuropathy with liability to pressure palsies. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul

2013]

**Protein Families:** Transmembrane

### **Product images:**



Coomassie blue staining of purified PMP22 protein (Cat# [TP318781]). The protein was produced from HEK293T cells transfected with PMP22 cDNA clone (Cat# [RC218781]) using

MegaTran 2.0 (Cat# [TT210002]).