

Product datasheet for **RG236169**

PMP22 (NM_001281455) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids
Product Name: PMP22 (NM_001281455) Human Tagged ORF Clone
Tag: TurboGFP
Symbol: PMP22
Synonyms: CIDP; CMT1A; CMT1E; DSS; GAS-3; GAS3; HMSNIA; HNPP; Sp110
Mammalian Cell Selection: Neomycin
Vector: pCMV6-AC-GFP (PS100010)
E. coli Selection: Ampicillin (100 ug/mL)
ORF Nucleotide Sequence: >RG236169 representing NM_001281455.
 Blue=ORF Red=Cloning site Green=Tag(s)

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GCTCGTTTAGTGAACCGTCAGAATTTGTAAACGACTCACTATAGGGCGCCGGGAATTCGTCGACTG
GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC
ATGCTCCTCCTGTTGCTGAGTATCATCGTCTCCACGTCGCGGTGCTGGTCTGCTGTTCTGCTCCACG
ATCGTCAGCCAATGGATCGTGGGCAATGGACACGCAACTGATCTCTGGCAGAAGTGTAGCACCTCTTCC
TCAGGAAATGTCCACCACTGTTTCTCATCATCACCAAACGAATGGCTGCAGTCTGTCCAGGCCACCATG
ATCCTGTCGATCATCTTCAGCATTCTGTCTCTGTTCTTCTGCGCAACTCTTCACCCTCACCAAG
GGGGCAGGTTTTACATCACTGGAATCTTCAAATCTTGCTGGTCTGTGCGTGATGAGTCTGCGGCC
ATCTACACGGTGAGGCACCCGGAGTGGCATCTCAACTCGGATTACTCCTACGGTTTCGCCTACATCCTG
GCCTGGGTGGCCTTCCCCTGGCCCTTCTCAGCGGTGTCATCTATGTGATCTTGGGAAACGCGAA
ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAAAC
```

Protein Sequence: >Peptide sequence encoded by RG236169
 Blue=ORF Red=Cloning site Green=Tag(s)

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MLLLLLSIIVLHVAVLVLLFVSTIVSQWIVGNHATDLWQNCSTSSSGNVHHCFS SSPNEWLQSVQATM
ILSIIIFSILSLFFCQLFTLTKGRFYITGIFQILAGLCVMSAAAIYVRHPEWHLNSDYSYGFAYIL
AWVAFPLALLSGVIYVILRKRE
TRTRPLEMESDESGLPAMEIECRITGTLNGVEFELVGGEGTPEQGRMTNKMSTKGALTFSPYLLSHV
MGYGFYHFGTYPYSGYENPFLHAINNGGYNTRIEKYEDGGVLHVSFSYRYEAGRVIGDFKVMGTGFPED
SVIFTDKIIRSNATVEHLHPMGDNDLDGSFTRTFSLRDGGYSSVVD SHMHFSAIHPSILQNGGPMFA
FRRVEEDHSNTELGIVEYQHAFKTPDADAGEERV
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Restriction Sites: Sgfl-MluI



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Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
RefSeq:	<u>NM_001281455.1</u> , <u>NP_001268384.1</u>
RefSeq Size:	1822 bp
RefSeq ORF:	483 bp
Locus ID:	5376
UniProt ID:	<u>Q01453</u>
Cytogenetics:	17p12
Protein Families:	Transmembrane
MW:	17.9 kDa
Gene 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 tissue-specific 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]