

Product datasheet for TP762613

OriGene Technologies, Inc.

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SPG7 (NM 003119) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Human spastic paraplegia 7 (pure and complicated

autosomal recessive) (SPG7), nuclear gene encoding mitochondrial protein, transcript variant

1, full length, with N-terminal GST and C-terminal His tag, expressed in E.coli, 50ug

Species: Human

Expression Host: E. coli

Expression cDNA Clone

or AA Sequence:

A DNA sequence encoding the region full length of SPG7

Tag: N-GST and C-HIS

Predicted MW: 88.2 kDa

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

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

Buffer: 50 mM Tris-HCl, pH 8.0, 8 M urea

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

some loss of protein during the filtration process.

Store at -80°C after receiving vials.

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 003110

Locus ID: 6687

UniProt ID: Q9UQ90

RefSeq Size: 3102

Cytogenetics: 16q24.3

RefSeq ORF: 2385

Synonyms: CAR; CMAR; PGN; SPG5C





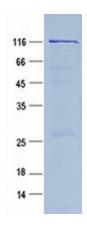
Summary:

This gene encodes a mitochondrial metalloprotease protein that is a member of the AAA family. Members of this protein family share an ATPase domain and have roles in diverse cellular processes including membrane trafficking, intracellular motility, organelle biogenesis, protein folding, and proteolysis. Mutations in this gene cause autosomal recessive spastic paraplegia 7. Two transcript variants encoding distinct isoforms have been identified. [provided by RefSeq, Mar 2014]

Protein Families:

Protease, Transmembrane

Product images:



Purified recombinant protein SPG7 was analyzed by SDS-PAGE gel and Coomossie Blue Staining.