

Product datasheet for TP311947M

C1orf69 (IBA57) (NM_001010867) Human Recombinant Protein

Product data:

| | |
|---------------------------------------|--|
| Product Type: | Recombinant Proteins |
| Description: | Recombinant protein of human chromosome 1 open reading frame 69 (C1orf69), 100 µg |
| Species: | Human |
| Expression Host: | HEK293T |
| Expression cDNA Clone or AA Sequence: | >RC211947 representing NM_001010867 Red =Cloning site Green =Tags(s) |
| | MATAALLRGATPGRGGPVWRWRLRAAPRCRLAHSSCSPGGDPTAGAAWACFRLDGRTLLRVRGPDAAPF L LGLLTNELPLPSPAAAGAPPAARAGYAHFLNVQGRTLYDVILYGLQEHSEVSGFLLCEDSSVQGALQKHL ALYRIRRKVTVEPHPELRVWAVLPSSPEACGAASLQERAGAAAAILRDPRTARMGWRLLTQDEGPALVPG GRLGDLWDYHQHRYLQGVPEGVRLPPGVALPLESNLAFMNGVSFTKGCYIGQELTARTHMGVIRKRL F PVRFLDPLPTSGITPGATVLTASGQTVGKFRAGQGNVGLALLWSEKIKGPLHIRASEGAQVALAASVPDW WPTVSK |
| | TRTRPLEQKLISEEDLAANDILDYKDDDDKV |
| Tag: | C-Myc/DDK |
| Predicted MW: | 38 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. |



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RefSeq: [NP_001010867](#)

Locus ID: 200205

UniProt ID: [Q5T440](#)

RefSeq Size: 7817

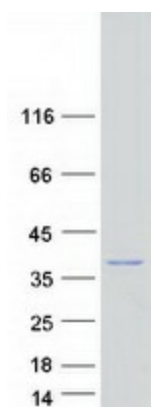
Cytogenetics: 1q42.13

RefSeq ORF: 1068

Synonyms: C1orf69; MMDS3; SPG74

Summary: The protein encoded by this gene localizes to the mitochondrion and is part of the iron-sulfur cluster assembly pathway. The encoded protein functions late in the biosynthesis of mitochondrial 4Fe-4S proteins. Defects in this gene have been associated with autosomal recessive spastic paraplegia-74 and with multiple mitochondrial dysfunctions syndrome-3. Two transcript variants encoding different isoforms have been found for this gene. The smaller isoform is not likely to be localized to the mitochondrion since it lacks the amino-terminal transit peptide. [provided by RefSeq, Jul 2015]

Product images:



Coomassie blue staining of purified IBA57 protein (Cat# [TP311947]). The protein was produced from HEK293T cells transfected with IBA57 cDNA clone (Cat# [RC211947]) using MegaTran 2.0 (Cat# [TT210002]).