

OriGene Technologies, Inc.

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Product datasheet for TP302230L

MCFD2 (NM_139279) Human Recombinant Protein

Product data:

| Product Type: | Recombinant Proteins | |
|--|--|--|
| Description: | Recombinant protein of human multiple coagulation factor deficiency 2 (MCFD2), 1 mg | |
| Species: | Human | |
| Expression Host: | HEK293T | |
| Expression cDNA Clone or AA Sequence: | >RC202230 protein sequence Red=Cloning site Green=Tags(s) | |
| | MTMRSLLRTPFLCGLLWAFCAPGARAEEPAASFSQPGSMGLDKNTVHDQEHIMEHLEGVINKPEAEMSP Q | |
| | ELQLHYFKMHDYDGNNLLDGLELSTAITHVHKEEGSEQAPLMSEDELINIIDGVLRDDDKNNDGYIDYAE FAKSLQ | |
| | TRTRPLEQKLISEEDLAANDILDYKDDDDKV | |
| Tag: | C-Myc/DDK | |
| Predicted MW: | 13.5 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: | <u>NP 644808</u> | |
| Locus ID: | 90411 | |
| UniProt ID: | <u>Q8NI22</u> | |



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| GRIGENE MCFD2 (NM_139279) Human Recombinant Protein – TP302230L | |
|--|--|
| RefSeq Size: | 4196 |
| Cytogenetics: | 2p21 |
| RefSeq ORF: | 438 |
| Synonyms: | F5F8D; F5F8D2; LMAN1IP; SDNSF |
| Summary: | This gene encodes a soluble luminal protein with two calmodulin-like EF-hand motifs at its C- terminus. This protein forms a complex with LMAN1 (lectin mannose binding protein 1; also known as ERGIC-53) that facilitates the transport of coagulation factors V (FV) and VIII (FVIII) from the endoplasmic reticulum to the Golgi apparatus via an endoplasmic reticulum Golgi intermediate compartment (ERGIC). Mutations in this gene cause combined deficiency of FV and FVIII (F5F8D); a rare autosomal recessive bleeding disorder characterized by mild to moderate bleeding and coordinate reduction in plasma FV and FVIII levels. This protein has also been shown to maintain stem cell potential in adult central nervous system and is a marker for testicular germ cell tumors. The 3' UTR of this gene contains a transposon-like human repeat element named 'THE 1'. A processed RNA pseudogene of this gene is on chromosome 6p22.1. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Apr 2016] |

Product images:

| 116 - | - |
|--------------|-----|
| 66 - | - |
| 45 - | - 1 |
| 35 - | - 1 |
| 25 - | - |
| 18 - 14 - | = |

Coomassie blue staining of purified MCFD2 protein (Cat# [TP302230]). The protein was produced from HEK293T cells transfected with MCFD2 cDNA clone (Cat# [RC202230]) using MegaTran 2.0 (Cat# [TT210002]).

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