

Product datasheet for TP316645

Factor I (CFI) (NM_000204) Human Recombinant Protein

Product data:

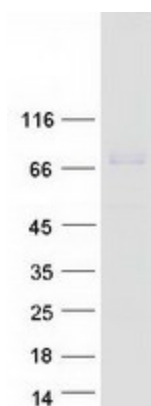
Product Type:	Recombinant Proteins
Description:	Recombinant protein of human complement factor I (CFI), 20 µg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC216645 representing NM_000204 Red =Cloning site Green =Tags(s)
	<p>MKLLHVFLFLCFHLRFCKVITYTSQEDLVEKKCLAKKYTHLSCDKVFCQPWQRCIEGTCVCKLPYQCPKN GTAVCATNRRSFPTYCQQKSLECLHPGKFLNNGTCTAEGKFSVSLKHGNTDSEGIVEVKLVDQDKTMFI CKSSWSMREANVACLDLGFQQGADTQRRFKLSDLSINSTECLHVHCRGLETSLAECTFTKRRTMGYQDFA DVVICYTQKADSPMDDFFQCVNGKYISQMKACDGINDCGDQSDCLCKACQGKGFHCKSGVCIPSQYQCNG EVDICITGEDEVGCAGFASVAQEETEILTADMDAERRRIKSLLPKLSGKVNRMHIRRKRIVGGKRAQLGD LPWQVAIKDASGITCGGIYIGGCWILTAAHCLRASKTHRYQIWTTVVDWIHPDLKRIVIEYVDRIIFHEN YNAGTYQNDIALIEMKKDGNKKDCELPRSIPACVPWSPYLFQPNDT CIVSGWGREKDNERVFLQWGEVK LISNCSKFGNRFYEKEMECAGTYDGSIDACKGDSGGPLVCM DANNVTVVWGVVSWG ENCGKPEFPGVYV KVANYFDWISYHVGRPFISQYNV</p> <p>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</p>
Tag:	C-Myc/DDK
Predicted MW:	63.4 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.



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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_000195</u>
Locus ID:	3426
UniProt ID:	<u>P05156, A8K3L0</u>
RefSeq Size:	1963
Cytogenetics:	4q25
RefSeq ORF:	1749
Synonyms:	AHUS3; ARMD13; C3b-INA; C3BINA; FI; IF; KAF
Summary:	This gene encodes a serine proteinase that is essential for regulating the complement cascade. The encoded preproprotein is cleaved to produce both heavy and light chains, which are linked by disulfide bonds to form a heterodimeric glycoprotein. This heterodimer can cleave and inactivate the complement components C4b and C3b, and it prevents the assembly of the C3 and C5 convertase enzymes. Defects in this gene cause complement factor I deficiency, an autosomal recessive disease associated with a susceptibility to pyogenic infections. Mutations in this gene have been associated with a predisposition to atypical hemolytic uremic syndrome, a disease characterized by acute renal failure, microangiopathic hemolytic anemia and thrombocytopenia. Primary glomerulonephritis with immune deposits and age-related macular degeneration are other conditions associated with mutations of this gene. [provided by RefSeq, Dec 2015]
Protein Families:	Druggable Genome, Protease, Secreted Protein
Protein Pathways:	Complement and coagulation cascades

Product images:



Coomassie blue staining of purified CFI protein (Cat# TP316645). The protein was produced from HEK293T cells transfected with CFI cDNA clone (Cat# [RC216645]) using MegaTran 2.0 (Cat# [TT210002]).