

Product datasheet for TP319727L

COG8 (NM_032382) Human Recombinant Protein

Product data:

Product Type:	Recombinant Proteins
Description:	Recombinant protein of human component of oligomeric golgi complex 8 (COG8), 1 mg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC219727 protein sequence Red =Cloning site Green =Tags(s)

MATAATIPSVATATAAALGEVEDEGLLASLFRDRFPEAQWRERPDVGRYLRELSGSLERLRREPERLAE
ERAQLLQQTRDLAFANYKTFIRGAECTERIHRLFGDVEASLGRLLDRLPSFQQSCRNFVKEAEEISSNRR
MNSLTLNRHTEILEILEIPQLMDTCVRNSYEEALELAAYVRRLEKYSSIPVIQGIVNEVRQSMQLMLS
QLIQQLRTNIQLPACLRIVIGYLRRMDVFTEAELRVKFLQARDAWLRSILTAIPNDDPYFHITKTIEASRV
HLFDIITQYRAIFSDDEDPLPPAMGEHTVNESAIFHWVWLQKVSQFLQVLETDLYRGIGGHLDSSLGQCM
YFGLSFSRVGADFRGQLAPVFQRVAISTFQKAIQETVEKFQEEMNSYMLISAPAILGTSNMPAAVPATQP
GTLQPPMVLLDFPPLACFLNNILVAFNDLRLCCPVALAQDVTGALEDALAKVTKIILAFHRAEEAAFFSSG
EQELFVQFCTVFLEDLVPYLNRLQVLFPPAQIAQTLGIPPTQLSKYGNLGHVNIQAIQEPLAFILPKRE
TLFTLDDQALGPELTAPAPEPPAEEPRLEPAGPACPEGGRAETQAEPSPVGP

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

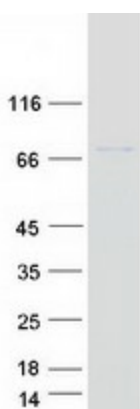
Tag:	C-Myc/DDK
Predicted MW:	68.2 kDa
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol
Preparation:	Recombinant protein was captured through anti-DDK affinity 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:	NP_115758
Locus ID:	84342
UniProt ID:	Q96MW5
RefSeq Size:	2522
Cytogenetics:	16q22.1
RefSeq ORF:	1836
Synonyms:	CDG2H; DOR1
Summary:	This gene encodes a protein that is a component of the conserved oligomeric Golgi (COG) complex, a multiprotein complex that plays a structural role in the Golgi apparatus, and is involved in intracellular membrane trafficking and glycoprotein modification. Mutations in this gene cause congenital disorder of glycosylation, type IIh, a disease that is characterized by under-glycosylated serum proteins, and whose symptoms include severe psychomotor retardation, failure to thrive, seizures, and dairy and wheat product intolerance. [provided by RefSeq, Jul 2008]
Protein Families:	Druggable Genome

Product images:



Coomassie blue staining of purified COG8 protein (Cat# [TP319727]). The protein was produced from HEK293T cells transfected with COG8 cDNA clone (Cat# [RC219727]) using MegaTran 2.0 (Cat# [TT210002]).