

Product datasheet for **SC318048**

COG8 (NM_032382) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	COG8 (NM_032382) Human Untagged Clone
Tag:	Tag Free
Symbol:	COG8
Synonyms:	CDG2H; DOR1
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)



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Fully Sequenced ORF: >NCBI ORF sequence for NM_032382, the custom clone sequence may differ by one or more nucleotides
 ATGGCGACCGCGGCGACTATCCCATCGGTAGCCACGGCCACAGCAGCGGCTCTCGGCGAG
 GTGGAGGATGAAGGGCTCCTGGCGTCGCTGTTCCGGGACCGCTTCCCCGAGGCCAGTGG
 CGCGAGCGGCCGATGTGGGCGCTACCTCCGGGAGTTGAGCGGCTCGGGGCTGGAGCGG
 CTGCGCGCGAGCCGAGCGCCTGGCGGAGGAGCGGGCGCAGCTGCTGCAGCAGACGCGC
 GACTTGGCCTTCGCTAACTACAAGACCTTCATCCGCGGCGCCGAGTGCACCGAGCGCATC
 CACCGCCTGTTTGGCGACGTGGAGGCGTCGCTCGGCCGCTGCTCGACCGTTTGCCGAGC
 TTCCAGCAGAGCTGCAGGAACCTTGTGAAGGAAGCCGAGGAGATCAGCTCCAACCGCCGG
 ATGAATAGCCTGACCCTAAACCGGCACACAGAAATTTGGAAATACTGGAGATTCCTCAG
 CTCATGGACACCTGTGTCCGGAACAGTTATTATGAAGAGGCCCTGGAGCTTGACAGCTAC
 GTACGCCGACTGGAGAGGAAATACTCTTCCATCCCTGTCATCCAGGGCATCGTGAACGAA
 GTGCGCCAGTCCATGCAGCTGATGCTGAGCCAGCTGATCCAGCAACTGAGGACCAACATC
 CAGCTTCTCGCTGCCTCCGTGTCATTGGCTACCTGCGGCGCATGGACGTCTTCACTGAG
 GCTGAGTTGAGGGTGAAGTTTCTCAGGCCGAGATGCTTGGCTCCGGTCCATCCTGACT
 GCCATTCTAATGATGATCCCTATTTCCATATTACAAAAACCATCGAGGCCCTCCCGTGT
 CATCTCTTTGATATCATACCCAGTACCGTGCCATCTTCTCAGACGAGGCCCACTGCTG
 CCCCCTGCCATGGGTGAGCACACTGTGAATGAGAGTGCCATCTTCCATGGCTGGGTGCTA
 CAGAAGGTCTCACAATTCCTGCAGGTGCTGGAGACCGACCTTTACCGGGGCATAGGCGGC
 CACCTGGACTCTCTGCTGGGCGCGTGCATGTACTTTGGGCTGTCTTACGCCGGTGGGA
 GCTGATTTCCGGGGTCAAGTTGGCTCCTGTTTTCCAGCGGGTGGCCATCAGCACTTCCAG
 AAAGCAATTCAGGAAACAGTGGAGAAATCCAGGAAGAAATGAACTCCTACATGCTCATC
 TCGGCTCCAGCCATCCTGGGCACAGTAACATGCCTGCTGCTGTGCCAGCCACCCAGCCG
 GGGACGCTGCAGCCACCCATGGTCTCCTAGATTTCCACCCCTCGCCTGCTTTCTCAAC
 AATATTCTGGTTGCCTTCAATGATCTGCGCCTCTGCTGCCCTGTGGCCCTGGCGCAGGAT
 GTGACTGGGGCCTTGAAGATGCCCTTGCCAAGGTAATAAAAATAATCCTGGCCTTCCAT
 CGCGCTGAAGAGGCTGCCTTACGAGCGGGGAGCAAGAGCTTTTGTCCAGTTCTGCACT
 GTCTTCTGGAAGACCTTGTCCGTATTTAAATCGCTGTCTCCAAGTCTTTTTCCACCA
 GCTCAGATAGCACAGACTTTAGGCATTCCTCCCACTCAGCTCTCCAAGTACGGTAACCTA
 GGGCATGTGAACATCGGCGCCATTGAGGAGCCCTCGCCTTTATCCTGCCAAAGAGAGAG
 ACGCTTTTACCCTGGATGACCAGGCGCTGGGGCCGAGCTCACAGCTCCAGCACCAGAG
 CCTCCCGCCGAGGAGCCACGCTGGAGCCCGCGGGCCAGCCTGCCCGGAGGGAGGGCGA
 GCGGAGACGCAGGCCAACCGCCAGCGTGGGGCCC

Restriction Sites: Please inquire

ACCN: NM_032382

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method:

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: [NM_032382.3](#), [NP_115758.2](#)

RefSeq Size: 2482 bp

RefSeq ORF: 1839 bp

Locus ID: 84342

UniProt ID: [Q96MW5](#)

Cytogenetics: 16q22.1

Domains: Dor1

Protein Families: Druggable Genome

Gene 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]