

## Product datasheet for **SC330223**

### CD16b (FCGR3B) (NM\_001244753) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	CD16b (FCGR3B) (NM_001244753) Human Untagged Clone
Tag:	Tag Free
Symbol:	CD16b
Synonyms:	CD16; CD16A; CD16b; FCG3; FCGR3; FCGR3A; FCR-10; FCRIII; FCRIIIb
Mammalian Cell Selection:	Neomycin
Vector:	<u>PCMV6-Neo</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_001244753, the custom clone sequence may differ by one or more nucleotides

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ATGGGTGGAGGGACTGGGAAAGGCTGTTACTCCCTCCTGTCTAGTCGGCTTGGTCCCTTTAGGGCTCC
GGATATCTTTGGTGACTTGCCACTCCAGTGTGGCATCATGTGGCAGCTGCTCCTCCCAACTGCTCTGCT
ACTTCTAGTTTCAGCTGGCATGCGGACTGAAGATCTCCCAAAGGCTGTGGTGTTCCTGGAGCCTCAATGG
TACAGCGTGCTTGAGAAGGACAGTGTGACTCTGAAGTCCAGGGAGCCTACTCCCCTGAGGACAATTCCA
CACAGTGGTTTCACAATGAGAACCTCATCTCAAGCCAGGCCTCGAGCTACTTCATTGACGCTGCCACAGT
CAACGACAGTGGAGAGTACAGGTGCCAGACAAACCTCTCCACCCTCAGTGACCCGGTGCAGCTAGAAGTC
CATATCGGCTGGCTGTTGCTCCAGGCCCTCGGTGGGTGTTCAAGGAGGAAGACCCTATTCACCTGAGGT
GTCACAGCTGGAAGAAGACTGCTCTGCATAAAGTACATATTTACAGAATGGCAAAGACAGGAAGTATTT
TCATCATAAATTGACTTCCACATTCCAAAAGCCACACTCAAAGATAGCGGCTCCTACTTCTGCAGGGGG
CTTGTTGGGAGTAAAAATGTGTCTTCAGAGACTGTGAACATCACCATCAAGGTTTGGCAGTGTCAA
CCATCTCATCTCTCCACCTGGGTACCAAGTCTCTTCTGCTTGGTGATGGTACTCCTTTTTGCAGT
GGACACAGGACTATTTCTCTGTGAAGACAAACATTTGA
```

Restriction Sites:	Sgfl-MluI
ACCN:	NM_001244753



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**OTI Disclaimer:** Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

**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\\_001244753.1](#), [NP\\_001231682.1](#)

**RefSeq Size:** 2394 bp

**RefSeq ORF:** 810 bp

**Locus ID:** 2215

**UniProt ID:** [O75015](#)

**Cytogenetics:** 1q23.3

**Protein Families:** ES Cell Differentiation/IPS, Secreted Protein, Transmembrane

**Protein Pathways:** Natural killer cell mediated cytotoxicity, Systemic lupus erythematosus

**Gene Summary:**

The protein encoded by this gene is a low affinity receptor for the Fc region of gamma immunoglobulins (IgG). The encoded protein acts as a monomer and can bind either monomeric or aggregated IgG. This gene may function to capture immune complexes in the peripheral circulation. Several transcript variants encoding different isoforms have been found for this gene. A highly-similar gene encoding a related protein is also found on chromosome 1. [provided by RefSeq, Aug 2012]

Transcript Variant: This variant (1) encodes the longest isoform (2). Variants 1 and 2 encode the same isoform. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments. CCDS Note: The coding region has been updated to shorten the N-terminus to one that is more supported by conservation.