

Product datasheet for SC205685

OriGene Technologies, Inc.

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GCSF Receptor (CSF3R) (NM_156038) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: GCSF Receptor (CSF3R) (NM 156038) Human 3' UTR Clone

Symbol: GCSF Receptor

Synonyms: CD114; CD114 antigen; colony stimulating factor 3 receptor; colony stimulating factor 3

receptor (granulocyte); GCSFR; granulocyte colony stimulating factor receptor; OTTHUMP0000009703; OTTHUMP0000009704; OTTHUMP0000009705

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_156038

Insert Size: 451 bp

Insert Sequence: >SC205685 3'UTR clone of NM_156038

The sequence shown below is from the reference sequence of NM_156038. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CAGTTTTTGTTGTTTATAGACACTCTTGGGTGTA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).





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Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

RefSeq: <u>NM 156038.2</u>

Summary: The protein encoded by this gene is the receptor for colony stimulating factor 3, a cytokine

that controls the production, differentiation, and function of granulocytes. The encoded protein, which is a member of the family of cytokine receptors, may also function in some cell surface adhesion or recognition processes. Alternatively spliced transcript variants have been described. Mutations in this gene are a cause of Kostmann syndrome, also known as severe

congenital neutropenia. [provided by RefSeq, Aug 2010]

Locus ID: 1441 MW: 16.4