

## Product datasheet for SC204135

### GCSF Receptor (CSF3R) (NM\_000760) Human 3' UTR Clone

#### Product data:

Product Type:	3' UTR Clones
Product Name:	GCSF Receptor (CSF3R) (NM_000760) Human 3' UTR Clone
Symbol:	GCSF Receptor
Synonyms:	CD114; GCSFR; SCN7
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_000760
Insert Size:	344 bp
Insert Sequence:	<p>&gt;SC204135 3'UTR clone of NM_000760</p> <p>The sequence shown below is from the reference sequence of NM_000760. The complete sequence of this clone may contain minor differences, such as SNPs.</p> <p>Blue=Stop Codon Red=Cloning site</p>

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GGCAAGTTGGACGCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAACGATCGCC
CATGGGATGGAGGCGCTGGGAGCTTCTAGGGCTTCTGGGGTTCCTTCTTGGGCTGCCTCTTAAAG
GCCTGAGCTAGCTGGAGAAGAGGGGAGGGTCCATAAGCCCATGACTAAAACTACCCAGCCCAGGCTC
TCACCATCTCCAGTCACCAGCATCTCCCTCTCCTCCCAATCTCCATAGGCTGGGCTCCCAGGCGATCT
GCATACTTTAAGGACCAGATCATGCTCCATCCAGCCCCACCAATGGCCTTTTGTGCTTGTTCCTATA
ACTTCAGTATTGTAACTAGTTTTTGGTTTGAGTTTTTGTGTTGTTTATAGACTCTTGGGTGTA
ACGCGTAAGCGGCCGCGCATCTAGATTGGAAGAAATGACCGACCAAGCGACGCCAACCTGCCATCA
CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG
  
```

Restriction Sites:	Sgfl-MluI
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).
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_000760.4</u>


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**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:** 12.6