

Product datasheet for SC323046

GFI1 (NM 001127216) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: GFI1 (NM_001127216) Human Untagged Clone

Tag: Tag Free

Symbol: GFI1

Synonyms: GFI-1; GFI1A; SCN2; ZNF163

Mammalian Cell

Selection:

Neomycin

Vector: pCMV6-Entry (PS100001)

E. coli Selection: Kanamycin (25 ug/mL)

Fully Sequenced ORF: >SC323046 representing NM_001127216.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GCTCGTTTAGTGAACCGTCAGAATTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ATGCCGCGCTCATTTCTCGTCAAAAGCAAGAAGGCTCACAGCTACCACCAGCCGCGCTCCCCAGGACCA GACTATTCCCTCCGTTTAGAGAATGTACCGGCGCCTAGCCGAGCAGCACACTTCAAATGCAGGCGGG GCATCCCCAGACAGCTGCGAAGGCAGCGTCTGCGAACGGAGCTCGGAGTTTGAGGACTTCTGGAGGCCC CCGTCACCCTCCGCGTCTCCAGCCTCGGAGAAGTCAATGTGCCCATCGCTGGACGAAGCCCAGCCCTTC CCCCTGCCTTTCAAACCGTACTCATGGAGCGGCCTGGCGGGTTCTGACCTGCGGCACCTGGTGCAGAGC TACCGACCGTGTGGGGCCCTGGAGCGTGGCGTGGCCTGGGCCTCTTCTGCGAACCCGCCCCGGAGCCT GGCCACCCGGCCGCGCTGTACGGCCCGAAGCGGGCTGCCGGCGGCGCGGGGGCCCGGGGGCGCCAGGGAGC TGCAGCGCAGGGGCCGGTGCCACCGCTGGCCTGGCCTAGGGCTCTACGGCGACTTCGGGTCTGCGGCA GCCGGGCTGTATGAGAGGCCCACGGCAGCGGCGGGCTTGCTGTACCCCGAGCGTGGCCACGGGCTGCAC GCAGACAAGGGCGCTGGCGTCAAGGTGGAGTCGGAGCTGCTGTGCACCCGCCTGCTGCTGGGCGGCGGC TCCTACAAGTGCATCAAGTGCAGCAAGGTGTTCTCCACGCCGCACGGGCTCGAGGTGCACGTGCGCAGG TCCCACAGCGGTACCAGACCCTTTGCCTGCGAGATGTGCGGCAAGACCTTCGGGCACGCGGTGAGCCTG GAGCAGCACAAAGCCGTGCACTCGCAGGAACGGAGCTTTGACTGTAAGATCTGTGGGAAGAGCTTCAAG AGGTCATCCACACTGTCCACACACCTGCTTATCCACTCAGACACTCGGCCCTACCCCTGTCAGTACTGT GGCAAGAGGTTCCACCAGAAGTCAGACATGAAGAAACACACTTTCATCCACACTGGTGAGAAGCCTCAC AAGTGCCAGGTGTGCGGCAAGGCATTCAGCCAGAGCTCCAACCTCATCACCCACAGCCGCAAACACACA GGCTTCAAGCCCTTCGGCTGCGACCTCTGTGGGAAGGGTTTCCAGAGGAAGGTGGACCTCCGAAGGCAC CGGGAGACGCAGCATGGGCTCAAATGA

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com **Restriction Sites:** Sgfl-Mlul

ACCN: NM_001127216

Insert Size: 1269 bp

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: <u>NM 001127216.1</u>

 RefSeq Size:
 2709 bp

 RefSeq ORF:
 1269 bp

 Locus ID:
 2672

 UniProt ID:
 Q99684

Protein Families: Druggable Genome, Transcription Factors

1p22.1

MW: 45.3 kDa

Cytogenetics:





Gene Summary:

This gene encodes a nuclear zinc finger protein that functions as a transcriptional repressor. This protein plays a role in diverse developmental contexts, including hematopoiesis and oncogenesis. It functions as part of a complex along with other cofactors to control histone modifications that lead to silencing of the target gene promoters. Mutations in this gene cause autosomal dominant severe congenital neutropenia, and also dominant nonimmune chronic idiopathic neutropenia of adults, which are heterogeneous hematopoietic disorders that cause predispositions to leukemias and infections. Multiple alternatively spliced variants, encoding the same protein, have been identified for this gene. [provided by RefSeq, Jul 2008] Transcript Variant: This variant (3) differs in the 5' UTR compared to variant 1. Variants 1, 2 and 3 encode the same protein. 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.