

Product datasheet for **SC207561**

IFNGR1 (NM_000416) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones
Product Name: IFNGR1 (NM_000416) Human 3' UTR Clone
Vector: pMirTarget (PS100062)
Symbol: IFNGR1
Synonyms: CD119; IFNGR; IMD27A; IMD27B
ACCN: NM_000416
Insert Size: 576 bp
Insert Sequence: >SC207561 3'UTR clone of NM_000416
The sequence shown below is from the reference sequence of NM_000416. The complete sequence of this clone may contain minor differences, such as SNPs.
Blue=Stop Codon **Red**=Cloning site

```
GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
CCAACAGAAGATTCCAAGAATTTTCATGAGATCAGCTAAGTTGCACCAACTTTGAAGTCTGATTTTCC
TGGACAGTTTTCTGCTTTAATTTTCATGAAAAGATTATGATCTCAGAAATTGTATCTTAGTTGGTATCAA
CCAAATGGAGTGACTTAGTGACATGAAAAGCGTAAAGAGGATGTGTGGCATTTCACCTTTGGCTTGTA
AAGTACAGACTTTTTTTTTTTTTTAAACAAAAAAGCATTGTAACCTTATGAACCTTTACATCCAGATAG
GTTACCAGTAACGGAACAGTATCCAGTACTCCTGGTTCCTAGGTGAGCAGGTGATGCCCCAGGGACCTT
TGTAGCCACTTCACTTTTTTTCTTTCTCTGCCTTGGTATAGCATATGTTTTGTAAAGTTTATGCATAC
AGTAATTTTAAGTAATTTTCAAGAAGAAATCTGCAAGCTTTTCAAATTTGGACTTAAATCTAATTCAAA
CTAATAGAATTAATGGAATATGTAATAGAAACGTGTATTTTTTATGAAACATTACAGTTAGAGATT
TTTAAATAAAGAATTTTAAACTC
ACGCGTAAGCGGCCGCGCATCTAGATTGAAGAAAATGACCGACCAAGCGACGCCAACCTGCCATCA
CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG
```

Restriction Sites: SgfI-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.



[View online »](#)

RefSeq: [NM_000416.3](#)

Summary: This gene (IFNGR1) encodes the ligand-binding chain (alpha) of the gamma interferon receptor. Human interferon-gamma receptor is a heterodimer of IFNGR1 and IFNGR2. A genetic variation in IFNGR1 is associated with susceptibility to Helicobacter pylori infection. In addition, defects in IFNGR1 are a cause of mendelian susceptibility to mycobacterial disease, also known as familial disseminated atypical mycobacterial infection. [provided by RefSeq, Jul 2008]

Locus ID: 3459

MW: 23.2