

Product datasheet for **SC207624**

IFNGR2 (NM_005534) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones
Product Name: IFNGR2 (NM_005534) Human 3' UTR Clone
Vector: pMirTarget (PS100062)
Symbol: IFNGR2
Synonyms: AF-1; IFGR2; IFNGT1; IMD28
ACCN: NM_005534
Insert Size: 584 bp
Insert Sequence: >SC207624 3'UTR clone of NM_005534
 The sequence shown below is from the reference sequence of NM_005534. The complete sequence of this clone may contain minor differences, such as SNPs.
 Blue=Stop Codon Red=Cloning site

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GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
GAGCAAGAAGATGTTCTCAAACGCTTTGACCAAAGCATGGGCCTAGCCACTGGCTCCCTGGAAGAG
ATCAAGCCATCGGAGCTGCTAGAGTTCTGTCTGGACTTCCAGAGACCAGTATCCCTTTTGTGCCTC
TAAAAGGCTGTCCCTGCAGACATGAGAGACAGCAGGTCTCATGGGGGTGACAAGCTTTTTTTTTTTT
CTTAAAGAATTTTCAAATCAAATTCAGAATGATTTTACGGAGATATCCAGGAAAATTAAGGCTTCT
CTTAAACACTAAAAGGCATGTAATTGCTTGTAGCAAAATGGATATGACACATCTCTGATACTTTTTT
CATTATTGGTTGGGCTGAGCAGTCAGAAGACCTGGTCGTCGTCTTGACTTTGGCAAATGAGCCGGAGCC
CCTTGGGCAGGTCACACAACCTGTCCCAGCGAGGGACACCGAGTGGCCCTTCATGTACATCCATGGTGT
GCTGGCTTAAAATGTAATTAATCTGTAAATATACTCCTAGTAATTTAAGATTTTGTTTTTAAACTGGA
AATAAAAGATTGTATAGTCATGTTTTTAA
ACGCGTAAGCGGCCGCGCATCTAGATTGAAAGAAATGACCGACCAAGCGACGCCCAACCTGCCATCA
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_005534.4](#)

Summary: This gene (IFNGR2) encodes the non-ligand-binding beta chain of the gamma interferon receptor. Human interferon-gamma receptor is a heterodimer of IFNGR1 and IFNGR2. Defects in IFNGR2 are a cause of mendelian susceptibility to mycobacterial disease (MSMD), also known as familial disseminated atypical mycobacterial infection. MSMD is a genetically heterogeneous disease with autosomal recessive, autosomal dominant or X-linked inheritance. [provided by RefSeq, Jul 2008]

Locus ID: 3460

MW: 22.5