

## **Product datasheet for SC208272**

## OriGene Technologies, Inc.

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## Repulsive Guidance Molecule C (HFE2) (NM\_145277) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: Repulsive Guidance Molecule C (HFE2) (NM\_145277) Human 3' UTR Clone

**Symbol:** Repulsive Guidance Molecule C

Synonyms: HFE2; HFE2A; JH; RGMC

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_145277

**Insert Size:** 637 bp

Insert Sequence: >SC208272 3'UTR clone of NM\_145277

The sequence shown below is from the reference sequence of NM\_145277. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

 ${\sf TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC}$ 

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).





## Repulsive Guidance Molecule C (HFE2) (NM\_145277) Human 3' UTR Clone - SC208272

**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:** NM 145277.5

Summary: The product of this gene is involved in iron metabolism. It may be a component of the

signaling pathway which activates hepcidin or it may act as a modulator of hepcidin

expression. It could also represent the cellular receptor for hepcidin. Two uORFs in the 5' UTR negatively regulate the expression and activity of the encoded protein. Alternatively spliced transcript variants encoding different isoforms have been identified for this gene. Defects in this gene are the cause of hemochromatosis type 2A, also called juvenile hemochromatosis (JH). JH is an early-onset autosomal recessive disorder due to severe iron overload resulting in

hypogonadotrophic hypogonadism, hepatic fibrosis or cirrhosis and cardiomyopathy,

occurring typically before age of 30. [provided by RefSeq, Oct 2015]

**Locus ID:** 148738

MW: 24.5