

## Product datasheet for RC217560L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## HFE (NM\_139007) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

Product Name: HFE (NM 139007) Human Tagged ORF Clone Lentiviral Particle

Symbol: HFE

**Synonyms:** HFE1; HH; HLA-H; MVCD7; TFQTL2

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_139007

ORF Size: 780 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC217560).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeg:** NM 139007.2

 RefSeq Size:
 1958 bp

 RefSeq ORF:
 783 bp

 Locus ID:
 3077

 UniProt ID:
 Q30201

 Cytogenetics:
 6p22.2

**Protein Families:** Druggable Genome, Transmembrane

**MW:** 29.6 kDa







## **Gene Summary:**

The protein encoded by this gene is a membrane protein that is similar to MHC class I-type proteins and associates with beta2-microglobulin (beta2M). It is thought that this protein functions to regulate iron absorption by regulating the interaction of the transferrin receptor with transferrin. The iron storage disorder, hereditary haemochromatosis, is a recessive genetic disorder that results from defects in this gene. At least nine alternatively spliced variants have been described for this gene. Additional variants have been found but their full-length nature has not been determined. [provided by RefSeq, Jul 2008]