

Product datasheet for RC220860

HFE (NM_139011) Human Tagged ORF Clone

Product data:

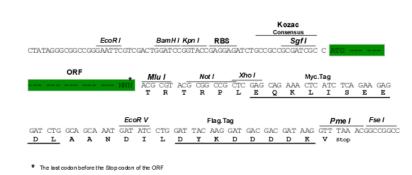
Tag:

Product Type: **Expression Plasmids Product Name:** HFE (NM_139011) Human Tagged ORF Clone Myc-DDK Symbol: HFE HFE1; HH; HLA-H; MVCD7; TFQTL2 Synonyms: **Mammalian Cell** Neomycin Selection: Vector: pCMV6-Entry (PS100001) E. coli Selection: Kanamycin (25 ug/mL) **Restriction Sites:** Sgfl-Mlul Cloning sites used for ORF Shuttling: **Cloning Scheme:**

NM 139011

228 bp





ACCN: **ORF Size:**



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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

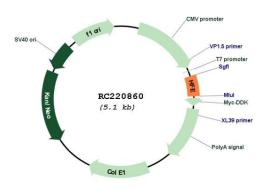
GRIGENE HFE (NM_139011) Human Tagged ORF Clone – RC220860	
OTI Disclaimer:	The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), 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. <u>More info</u>
OTI Annotation:	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
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:	 Centrifuge at 5,000xg for 5min. Carefully open the tube and add 100ul of sterile water to dissolve the DNA. Close the tube and incubate for 10 minutes at room temperature. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom. 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 139011.3</u>
RefSeq Size:	1406 bp
RefSeq ORF:	231 bp
Locus ID:	3077
UniProt ID:	<u>Q30201</u>
Cytogenetics:	6p22.2
Protein Families:	Druggable Genome, Transmembrane
MW:	8.2 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]

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Product images:



Circular map for RC220860

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