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Product datasheet for SC201447

CFHL2 (CFHR2) (NM_005666) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Product Name:	CFHL2 (CFHR2) (NM_005666) Human 3' UTR Clone
Symbol:	CFHL2
Synonyms:	CFHL2; FHR2; HFL3
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_005666
Insert Size:	572 bp
Insert Sequence:	>SC201447 3'UTR clone of NM_005666 The sequence shown below is from the reference sequence of NM_005666. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC CTGGTATATCCCAGTTGTGAAGAAAAATAGAATCAATGGCATTACTATTAGTAAAATGCACACCTTTTT CTGAATTTACTATTATATTTGTTTTCAATTTCATTTTTCAAGTACTGTTTTACTCATTTTTATTCATAA ATAAAGTTTTGTGTTGATTTGTGAAAATGCAATTACAATCTGAGATGTGTCACAATGGTGAGGACTATC TTCACCAAATCTAAGTAACAACCTAGGAATTGTCTTTTTTTTTCTTTTTAAAAAAATTGACAATAACTG TATATATTCATGGAGTACATAGTAATGTTTCCATATATATAATGTATAATGGTCAGTTAGGGTAATTAG TATATCCATTATCTCAAACATTTTTCATTTCTTTGGGTTAGGAGCATTAAATATTCTCCTTCCAGCTAT TTGGTACTTCATAGTATATTACTGGTAACTGAAGGAATTATATCTAGACGTTACCCCAGGTATCTTGAA ATGTCAATTCCTAACAGTCACAGCCTGGGAGCTCATGTTTGCCTTCTTTCAGAGCTTGTAACTATGTAT ATCCACATAAATAATCAAAA ACGCGTAAGCGGCCGCGGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG
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).

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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_005666.4
Summary:	This gene belongs to a family of complement factor H-related genes (CFHR), which are clustered together with complement factor H gene on chromosome 1, and are involved in regulation of complement. Mutations in CFHR genes have been associated with dense deposit disease and atypical haemolytic-uraemic syndrome. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Aug 2015]
Locus ID:	3080
MW:	22.9