

## **Product datasheet for SC211621**

## CFHR5 (NM 030787) Human 3' UTR Clone

## **Product data:**

**Product Type:** 3' UTR Clones

**Product Name:** CFHR5 (NM\_030787) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: CFHR5

**Synonyms:** CFHL5; CFHR5D; FHR-5; FHR5

**ACCN:** NM\_030787

**Insert Size:** 1025 bp

Insert Sequence: >SC211621 3'UTR clone of NM\_030787

The sequence shown below is from the reference sequence of NM\_030787. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TACCTATCCAATACTAAATACCCCTTAAAGTATTAAATGCACTATCTGCTGTAAAAGAA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul



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## CFHR5 (NM\_030787) Human 3' UTR Clone - SC211621

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.

**RefSeq:** <u>NM 030787.4</u>

Summary: This gene is a member of a small complement factor H (CFH) gene cluster on chromosome 1.

Each member of this gene family contains multiple short consensus repeats (SCRs) typical of regulators of complement activation. The protein encoded by this gene has nine SCRs with the first two repeats having heparin binding properties, a region within repeats 5-7 having heparin binding and C reactive protein binding properties, and the C-terminal repeats being similar to a complement component 3 b (C3b) binding domain. This protein co-localizes with C3, binds C3b in a dose-dependent manner, and is recruited to tissues damaged by C-reactive protein. Allelic variations in this gene have been associated, but not causally linked, with two different forms of kidney disease: membranoproliferative glomerulonephritis type II

(MPGNII) and hemolytic uraemic syndrome (HUS). [provided by RefSeq, Jan 2010]

**Locus ID:** 81494

MW: 38.8