

Product datasheet for SC212048

FMN2 (NM_020066) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: FMN2 (NM_020066) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: FMN2

ACCN: NM_020066

Insert Size: 1063 bp

Insert Sequence: >SC212048 3'UTR clone of NM_020066

The sequence shown below is from the reference sequence of NM_020066. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TAATAAAGAATAAAGATACTTGCAAAA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul



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FMN2 (NM_020066) Human 3' UTR Clone - SC212048

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 020066.5</u>

Summary: This gene is a member of the formin homology protein family. The encoded protein is

thought to have essential roles in organization of the actin cytoskeleton and in cell polarity. This protein mediates the formation of an actin mesh that positions the spindle during oogenesis and also regulates the formation of actin filaments in the nucleus. This protein also forms a perinuclear actin/focal-adhesion system that regulates the shape and position of the nucleus during cell migration. Mutations in this gene have been associated with infertility and also with an autosomal recessive form of intellectual disability (MRT47). Alternatively

spliced transcript variants have been identified. [provided by RefSeq, Jul 2017]

Locus ID: 56776

MW: 40.9