

Product datasheet for SC205013

OriGene Technologies, Inc.

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KIRREL 3 (KIRREL3) (NM_001161707) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: KIRREL 3 (KIRREL3) (NM 001161707) Human 3' UTR Clone

Symbol: KIRREL 3

Synonyms: KIRRE; MRD4; NEPH2; PRO4502

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_001161707

Insert Size: 402 bp

Insert Sequence: >SC205013 3'UTR clone of NM_001161707

The sequence shown below is from the reference sequence of NM_001161707. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ACAGCTGTGGGGAGTGGGTGGGGAAGCAATAAAGGAATTGCTTTGAGAAAACTTAA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

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.





KIRREL 3 (KIRREL3) (NM_001161707) Human 3' UTR Clone - SC205013

RefSeq: <u>NM 001161707.1</u>

Summary: The protein encoded by this gene is a member of the nephrin-like protein family. These

proteins are expressed in fetal and adult brain, and also in podocytes of kidney glomeruli. The cytoplasmic domains of these proteins interact with the C-terminus of podocin, also

expressed in the podocytes, cells involved in ensuring size- and charge-selective

ultrafiltration. The protein encoded by this gene is a synaptic cell adhesion molecule with multiple extracellular immunoglobulin-like domains and a cytoplasmic PDZ domain-binding motif. Mutations in this gene are associated with several neurological and cognitive disorders. Alternatively spliced transcript variants encoding different isoforms have been found for this

gene. [provided by RefSeq, Jul 2017]

Locus ID: 84623

MW: 15