

Product datasheet for RC214037L4V

OriGene Technologies, Inc.

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FUSIP1 (SRSF10) (NM_054016) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: FUSIP1 (SRSF10) (NM 054016) Human Tagged ORF Clone Lentiviral Particle

Symbol: FUSIP1

Synonyms: FUSIP1; FUSIP2; NSSR; PPP1R149; SFRS13; SFRS13A; SRp38; SRrp40; TASR; TASR1; TASR2

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_054016

ORF Size: 786 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC214037).

Sequence:
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. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeg: NM 054016.2

 RefSeq Size:
 7792 bp

 RefSeq ORF:
 789 bp

 Locus ID:
 10772

 UniProt ID:
 075494

 Cytogenetics:
 1p36.11

 Domains:
 RRM

Protein Families: Transcription Factors





Protein Pathways: Spliceosome

MW: 31.3 kDa

Gene Summary: This gene product is a member of the serine-arginine (SR) family of proteins, which are

involved in constitutive and regulated RNA splicing. Members of this family are characterized by N-terminal RNP1 and RNP2 motifs, which are required for binding to RNA, and multiple C-terminal SR/RS repeats, which are important in mediating association with other cellular proteins. This protein interacts with the oncoprotein TLS, and abrogates the influence of TLS on adenovirus E1A pre-mRNA splicing. This gene has pseudogenes on chromosomes 4, 9, 14, 18, and 20. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul

2014]