

Product datasheet for RC206831L2V

OriGene Technologies, Inc.

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SEPT5 (SEPTIN5) (NM 002688) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SEPT5 (SEPTIN5) (NM_002688) Human Tagged ORF Clone Lentiviral Particle

Symbol: SEPTIN5

Synonyms: CDCREL; CDCREL-1; CDCREL1; H5; HCDCREL-1; PNUTL1; SEPT5

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_002688 **ORF Size:** 1107 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC206831).

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.

RefSeq: <u>NM 002688.4</u>

 RefSeq Size:
 2090 bp

 RefSeq ORF:
 1110 bp

 Locus ID:
 5413

 UniProt ID:
 Q99719

 Cytogenetics:
 22q11.21

Domains: GTP_CDC

Protein Families: Druggable Genome





Protein Pathways: Parkinson's disease

MW: 42.8 kDa

Gene Summary: This gene is a member of the septin gene family of nucleotide binding proteins, originally

described in yeast as cell division cycle regulatory proteins. Septins are highly conserved in yeast, Drosophila, and mouse and appear to regulate cytoskeletal organization. Disruption of septin function disturbs cytokinesis and results in large multinucleate or polyploid cells. This gene is mapped to 22q11, the region frequently deleted in DiGeorge and velocardiofacial syndromes. A translocation involving the MLL gene and this gene has also been reported in patients with acute myeloid leukemia. Alternative splicing results in multiple transcript variants. The presence of a non-consensus polyA signal (AACAAT) in this gene also results in read-through transcription into the downstream neighboring gene (GP1BB; platelet

glycoprotein lb), whereby larger, non-coding transcripts are produced. [provided by RefSeq,

Dec 2010]