

Product datasheet for RC226485L1V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Alpha Fodrin (SPTAN1) (NM 001130438) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Alpha Fodrin (SPTAN1) (NM 001130438) Human Tagged ORF Clone Lentiviral Particle

Symbol: Alpha Fodrin

Synonyms: DEE5; EIEE5; NEAS; SPTA2

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

ACCN: NM_001130438

ORF Size: 7431 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC226485).

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: NM 001130438.2, NP 001123910.1

 RefSeq Size:
 7907 bp

 RefSeq ORF:
 7434 bp

 Locus ID:
 6709

 UniProt ID:
 Q13813

 Cytogenetics:
 9q34.11

Protein Families: Druggable Genome

Protein Pathways: Tight junction





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MW: 285.5 kDa

Gene Summary:

Spectrins are a family of filamentous cytoskeletal proteins that function as essential scaffold proteins that stabilize the plasma membrane and organize intracellular organelles. Spectrins are composed of alpha and beta dimers that associate to form tetramers linked in a head-to-head arrangement. This gene encodes an alpha spectrin that is specifically expressed in nonerythrocytic cells. The encoded protein has been implicated in other cellular functions including DNA repair and cell cycle regulation. Mutations in this gene are the cause of early infantile epileptic encephalopathy-5. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Sep 2010]