

Product datasheet for RC223780L1V

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FADS2 (NM_004265) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: FADS2 (NM_004265) Human Tagged ORF Clone Lentiviral Particle

Symbol: FADS2

Synonyms: D6D; DES6; FADSD6; LLCDL2; SLL0262; TU13

Mammalian Cell

Selection:

None

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

Tag: Myc-DDK
ACCN: NM 004265

ORF Size: 1332 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC223780).

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 004265.2</u>

 RefSeq Size:
 3149 bp

 RefSeq ORF:
 1335 bp

 Locus ID:
 9415

 UniProt ID:
 095864

 Cytogenetics:
 11q12.2

Domains: heme_1, FA_desaturase

Protein Families: Transmembrane





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Protein Pathways: alpha-Linolenic acid metabolism, Biosynthesis of unsaturated fatty acids, PPAR signaling

pathway

MW: 52.1 kDa

Gene Summary: The protein encoded by this gene is a member of the fatty acid desaturase (FADS) gene

family. Desaturase enzymes regulate unsaturation of fatty acids through the introduction of double bonds between defined carbons of the fatty acyl chain. FADS family members are considered fusion products composed of an N-terminal cytochrome b5-like domain and a C-terminal multiple membrane-spanning desaturase portion, both of which are characterized by conserved histidine motifs. This gene is clustered with family members at 11q12-q13.1; this cluster is thought to have arisen evolutionarily from gene duplication based on its similar exon/intron organization. Alternative splicing results in multiple transcript variants encoding

different isoforms. [provided by RefSeq, Jul 2013]