

Product datasheet for RC205511L4V

OriGene Technologies, Inc.

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LASS3 (CERS3) (NM_178842) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: LASS3 (CERS3) (NM_178842) Human Tagged ORF Clone Lentiviral Particle

Symbol: LASS3

Synonyms: ARCI9; LASS3

Mammalian Cell Puromycin

Selection:

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

Tag: mGFP

ACCN: NM_178842

ORF Size: 1149 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC205511).

OTI Disclaimer:

Sequence:

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 178842.3

 RefSeq Size:
 3894 bp

 RefSeq ORF:
 1152 bp

 Locus ID:
 204219

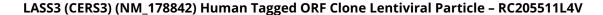
 UniProt ID:
 Q8IU89

 Cytogenetics:
 15q26.3

Protein Families: Transcription Factors, Transmembrane

MW: 46.2 kDa







Gene Summary:

This gene is a member of the ceramide synthase family of genes. The ceramide synthase enzymes regulate sphingolipid synthesis by catalyzing the formation of ceramides from sphingoid base and acyl-coA substrates. This family member is involved in the synthesis of ceramides with ultra-long-chain acyl moieties (ULC-Cers), important to the epidermis in its role in creating a protective barrier from the environment. The protein encoded by this gene has also been implicated in modification of the lipid structures required for spermatogenesis. Mutations in this gene have been associated with male fertility defects, and epidermal defects, including ichthyosis. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Aug 2015]