

Product datasheet for RC209557L2V

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

SLC27A4 (NM_005094) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SLC27A4 (NM_005094) Human Tagged ORF Clone Lentiviral Particle

Symbol: SLC27A4

Synonyms: ACSVL4; FATP4; IPS

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_005094 **ORF Size:** 1929 bp

ORF Nucleotide

OTI Disclaimer:

- - - -

Sequence:

The ORF insert of this clone is exactly the same as(RC209557).

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 005094.2

 RefSeq Size:
 2991 bp

 RefSeq ORF:
 1932 bp

 Locus ID:
 10999

 UniProt ID:
 Q6P1M0

 Cytogenetics:
 9q34.11

Domains: AMP-binding

Protein Families: Transmembrane





SLC27A4 (NM_005094) Human Tagged ORF Clone Lentiviral Particle - RC209557L2V

Protein Pathways: PPAR signaling pathway

MW: 71.9 kDa

Gene Summary: This gene encodes a member of a family of fatty acid transport proteins, which are involved

in translocation of long-chain fatty acids cross the plasma membrane. This protein is expressed at high levels on the apical side of mature enterocytes in the small intestine, and appears to be the principal fatty acid transporter in enterocytes. Clinical studies suggest this gene as a candidate gene for the insulin resistance syndrome. Mutations in this gene have

been associated with ichthyosis prematurity syndrome. [provided by RefSeq, Apr 2010]