

## Product datasheet for RC201153L4V

## 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

## FVT1 (KDSR) (NM\_002035) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** FVT1 (KDSR) (NM\_002035) Human Tagged ORF Clone Lentiviral Particle

Symbol: FVT1

Synonyms: DHSR; EKVP4; FVT1; SDR35C1

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_002035

ORF Size: 996 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC201153).

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.

**RefSeq:** <u>NM 002035.1</u>

 RefSeq Size:
 5198 bp

 RefSeq ORF:
 999 bp

 Locus ID:
 2531

 UniProt ID:
 Q06136

 Cytogenetics:
 18q21.33

**Domains:** adh\_short

**Protein Families:** Druggable Genome, Transmembrane





## FVT1 (KDSR) (NM\_002035) Human Tagged ORF Clone Lentiviral Particle - RC201153L4V

**Protein Pathways:** Metabolic pathways, Sphingolipid metabolism

MW: 36.2 kDa

**Gene Summary:** The protein encoded by this gene catalyzes the reduction of 3-ketodihydrosphingosine to

dihydrosphingosine. The putative active site residues of the encoded protein are found on the cytosolic side of the endoplasmic reticulum membrane. A chromosomal rearrangement involving this gene is a cause of follicular lymphoma, also known as type II chronic lymphatic leukemia. The mutation of a conserved residue in the bovine ortholog causes spinal muscular

atrophy. [provided by RefSeq, Jul 2008]