

Product datasheet for RC205948L4V

OriGene Technologies, Inc.

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TRIM37 (NM_001005207) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: TRIM37 (NM_001005207) Human Tagged ORF Clone Lentiviral Particle

Symbol: TRIM37

Synonyms: MUL; POB1; TEF3

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

ACCN: NM_001005207

ORF Size: 2892 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC205948).

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 001005207.1</u>

 RefSeq Size:
 3588 bp

 RefSeq ORF:
 2895 bp

 Locus ID:
 4591

 UniProt ID:
 094972

 Cytogenetics:
 17q22

Protein Families: Druggable Genome

Protein Pathways: Ubiquitin mediated proteolysis







MW: 107.7 kDa

Gene Summary:

This gene encodes a member of the tripartite motif (TRIM) family, whose members are involved in diverse cellular functions such as developmental patterning and oncogenesis. The TRIM motif includes zinc-binding domains, a RING finger region, a B-box motif and a coiled-coil domain. The RING finger and B-box domains chelate zinc and might be involved in protein-protein and/or protein-nucleic acid interactions. Mutations in this gene are associated with mulibrey (muscle-liver-brain-eye) nanism, an autosomal recessive disorder that involves several tissues of mesodermal origin. TRIM37 localizes in peroxisomal membranes, and has been implicated in human peroxisomal biogenesis disorders. [provided by RefSeq, Jul 2020]