

Product datasheet for RC221263L3V

OriGene Technologies, Inc.

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RANK (TNFRSF11A) (NM 003839) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: RANK (TNFRSF11A) (NM 003839) Human Tagged ORF Clone Lentiviral Particle

Symbol: RANK

Synonyms: CD265; FEO; LOH18CR1; ODFR; OFE; OPTB7; OSTS; PDB2; RANK; TRANCE-R; TRANCER

Mammalian Cell

Selection:

ACCN:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

NM 003839

Tag: Myc-DDK

ORF Size: 1848 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC221263).

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.

RefSeg: NM 003839.2

 RefSeq Size:
 3133 bp

 RefSeq ORF:
 1851 bp

 Locus ID:
 8792

 UniProt ID:
 Q9Y6Q6

Cytogenetics: 18q21.33

Domains: TNFR

Protein Families: Druggable Genome, Transmembrane





RANK (TNFRSF11A) (NM_003839) Human Tagged ORF Clone Lentiviral Particle - RC221263L3V

Protein Pathways: Cytokine-cytokine receptor interaction

MW: 66.03 kDa

Gene Summary: The protein encoded by this gene is a member of the TNF-receptor superfamily. This

receptors can interact with various TRAF family proteins, through which this receptor induces the activation of NF-kappa B and MAPK8/JNK. This receptor and its ligand are important regulators of the interaction between T cells and dendritic cells. This receptor is also an essential mediator for osteoclast and lymph node development. Mutations at this locus have been associated with familial expansile osteolysis, autosomal recessive osteopetrosis, and Paget disease of bone. Alternatively spliced transcript variants have been described for this

locus. [provided by RefSeq, Aug 2012]