

## Product datasheet for RC213091L3V

## OriGene Technologies, Inc.

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## TrkA (NTRK1) (NM\_001012331) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: TrkA (NTRK1) (NM\_001012331) Human Tagged ORF Clone Lentiviral Particle

Symbol: TrkA

Synonyms: MTC; p140-TrkA; TRK; Trk-A; TRK1; TRKA

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK

**ACCN:** NM\_001012331

ORF Size: 2370 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC213091).

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

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:** NM 001012331.1, NP 001012331.1

RefSeq Size: 2647 bp
RefSeq ORF: 2373 bp
Locus ID: 4914
UniProt ID: P04629

Cytogenetics: 1q23.1

**Protein Families:** Druggable Genome, Protein Kinase, Transmembrane





## TrkA (NTRK1) (NM\_001012331) Human Tagged ORF Clone Lentiviral Particle - RC213091L3V

Protein Pathways: Apoptosis, Endocytosis, MAPK signaling pathway, Neurotrophin signaling pathway, Pathways

in cancer, Thyroid cancer

MW: 86.7 kDa

**Gene Summary:** This gene encodes a member of the neurotrophic tyrosine kinase receptor (NTKR) family. This

kinase is a membrane-bound receptor that, upon neurotrophin binding, phosphorylates itself and members of the MAPK pathway. The presence of this kinase leads to cell differentiation and may play a role in specifying sensory neuron subtypes. Mutations in this gene have been associated with congenital insensitivity to pain, anhidrosis, self-mutilating behavior, cognitive disability and cancer. Alternate transcriptional splice variants of this gene have been found,

but only three have been characterized to date. [provided by RefSeq, Jul 2008]