

## Product datasheet for RC222143L3V

## OriGene Technologies, Inc.

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# P2RX7 (NM\_002562) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** P2RX7 (NM\_002562) Human Tagged ORF Clone Lentiviral Particle

Symbol: P2RX7
Synonyms: P2X7

Mammalian Cell Puromycin

Selection:

Vector:

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

 Tag:
 Myc-DDK

 ACCN:
 NM\_002562

 ORF Size:
 1785 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC222143).

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 002562.4

 RefSeq Size:
 3155 bp

 RefSeq ORF:
 1788 bp

 Locus ID:
 5027

 UniProt ID:
 Q99572

 Cytogenetics:
 12q24.31

**Domains:** P2X\_receptor

**Protein Families:** Druggable Genome, Ion Channels: ATP Receptors, Transmembrane





## P2RX7 (NM\_002562) Human Tagged ORF Clone Lentiviral Particle - RC222143L3V

**Protein Pathways:** Calcium signaling pathway, Neuroactive ligand-receptor interaction

MW: 68.4 kDa

**Gene Summary:** The product of this gene belongs to the family of purinoceptors for ATP. This receptor

functions as a ligand-gated ion channel and is responsible for ATP-dependent lysis of macrophages through the formation of membrane pores permeable to large molecules. Activation of this nuclear receptor by ATP in the cytoplasm may be a mechanism by which cellular activity can be coupled to changes in gene expression. Multiple alternatively spliced variants have been identified, most of which fit nonsense-mediated decay (NMD) criteria.

[provided by RefSeq, Jul 2010]