

## Product datasheet for RC216373L1V

## 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

## LRRK2 (NM\_198578) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

Product Name: LRRK2 (NM 198578) Human Tagged ORF Clone Lentiviral Particle

Symbol: LRRK2

Synonyms: AURA17; DARDARIN; PARK8; RIPK7; ROCO2

NM 198578

**Mammalian Cell** 

Selection:

ACCN:

None

**Vector:** pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

ORF Size: 7581 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC216373).

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 198578.2

 RefSeq Size:
 9234 bp

 RefSeq ORF:
 7584 bp

 Locus ID:
 120892

 UniProt ID:
 Q55007

 Cytogenetics:
 12q12

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

**Protein Pathways:** Parkinson's disease





## LRRK2 (NM\_198578) Human Tagged ORF Clone Lentiviral Particle - RC216373L1V

**MW:** 286.1 kDa

**Gene Summary:** This gene is a member of

This gene is a member of the leucine-rich repeat kinase family and encodes a protein with an ankryin repeat region, a leucine-rich repeat (LRR) domain, a kinase domain, a DFG-like motif, a RAS domain, a GTPase domain, a MLK-like domain, and a WD40 domain. The protein is present largely in the cytoplasm but also associates with the mitochondrial outer membrane. Mutations in this gene have been associated with Parkinson disease-8. [provided by RefSeq,

Jul 2008]