

## Product datasheet for RC201357L3V

## OriGene Technologies, Inc.

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## NALP2 (NLRP2) (NM 017852) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** NALP2 (NLRP2) (NM\_017852) Human Tagged ORF Clone Lentiviral Particle

Symbol:

CLR19.9; NALP2; NBS1; PAN1; PYPAF2 Synonyms:

**Mammalian Cell** 

Selection:

Puromycin

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

Myc-DDK Tag: ACCN: NM 017852

**ORF Size:** 3186 bp

**ORF Nucleotide** 

120.3 kDa

Sequence: OTI Disclaimer:

MW:

The ORF insert of this clone is exactly the same as(RC201357).

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: NM 017852.1

RefSeq Size: 3531 bp RefSeq ORF: 3189 bp Locus ID: 55655 **UniProt ID:** Q9NX02 Cytogenetics: 19q13.42 **Domains:** LRR, LRR RI





## **Gene Summary:**

This gene is a member of the nucleotide-binding and leucine-rich repeat receptor (NLR) family, and is predicted to contain an N-terminal pyrin effector domain (PYD), a centrally-located nucleotide-binding and oligomerization domain (NACHT) and C-terminal leucine-rich repeats (LRR). Members of this gene family are thought to be important regulators of immune responses. This gene product interacts with components of the IkB kinase (IKK) complex, and can regulate both caspase-1 and NF-kB (nuclear factor kappa-light-chain-enhancer of activated B cells) activity. The pyrin domain is necessary and sufficient for suppression of NF-kB activity. An allelic variant (rs147585490) has been found that is incapable of blocking the transcriptional activity of NF-kB. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Dec 2016]