

## Product datasheet for RC211819L3V

## OriGene Technologies, Inc.

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## iNOS (NOS2) (NM 000625) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** iNOS (NOS2) (NM\_000625) Human Tagged ORF Clone Lentiviral Particle

Symbol:

HEP-NOS; INOS; NOS; NOS2A Synonyms:

**Mammalian Cell** 

Selection:

ACCN:

Puromycin

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

Tag: Myc-DDK NM 000625

**ORF Size:** 3459 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC211819).

Sequence:

The molecular sequence of this clone aligns with the gene accession number as a point of OTI Disclaimer: 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 000625.3

RefSeq Size: 4221 bp RefSeq ORF: 3462 bp Locus ID: 4843 **UniProt ID:** P35228

Cytogenetics: 17q11.2

**Protein Families:** Druggable Genome





## iNOS (NOS2) (NM\_000625) Human Tagged ORF Clone Lentiviral Particle - RC211819L3V

**Protein Pathways:** Alzheimer's disease, Amyotrophic lateral sclerosis (ALS), Arginine and proline metabolism,

Calcium signaling pathway, Long-term depression, Metabolic pathways, Pathways in cancer,

Small cell lung cancer

MW: 130.9 kDa

**Gene Summary:** Nitric oxide is a reactive free radical which acts as a biologic mediator in several processes,

including neurotransmission and antimicrobial and antitumoral activities. This gene encodes a nitric oxide synthase which is expressed in liver and is inducible by a combination of lipopolysaccharide and certain cytokines. Three related pseudogenes are located within the

Smith-Magenis syndrome region on chromosome 17. [provided by RefSeq, Jul 2008]