

## Product datasheet for **RC209228L3V**

### eNOS (NOS3) (NM\_000603) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

Product Type:	Lentiviral Particles
Product Name:	eNOS (NOS3) (NM_000603) Human Tagged ORF Clone Lentiviral Particle
Symbol:	eNOS
Synonyms:	ECNOS; eNOS
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-Myc-DDK-P2A-Puro (PS100092)
Tag:	Myc-DDK
ACCN:	NM_000603
ORF Size:	3609 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC209228).
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. <a href="#">More info</a>
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:	<a href="#">NM_000603.3</a>
RefSeq Size:	4345 bp
RefSeq ORF:	3612 bp
Locus ID:	4846
UniProt ID:	<a href="#">P29474</a>
Cytogenetics:	7q36.1
Domains:	flavodoxin, NAD_binding_1, FAD_binding_1, NO_synthase
Protein Families:	Druggable Genome



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**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, VEGF signaling pathway

**MW:** 133.3 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. Nitric oxide is synthesized from L-arginine by nitric oxide synthases. Variations in this gene are associated with susceptibility to coronary spasm. Alternative splicing and the use of alternative promoters results in multiple transcript variants. [provided by RefSeq, Oct 2016]