

Product datasheet for RC223136L3V

OriGene Technologies, Inc.

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NMDAR2A (GRIN2A) (NM 000833) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: NMDAR2A (GRIN2A) (NM_000833) Human Tagged ORF Clone Lentiviral Particle

Symbol: GRIN2A

Synonyms: EPND; FESD; GluN2A; LKS; NMDAR2A; NR2A

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 000833

ORF Size: 4392 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC223136).

Sequence:
OTI Disclaimer:

Cytogenetics:

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 000833.2

 RefSeq Size:
 14450 bp

 RefSeq ORF:
 4395 bp

 Locus ID:
 2903

 UniProt ID:
 Q12879

Protein Families: Druggable Genome, Ion Channels: Glutamate Receptors, Ion Channels: Sodium,

Transmembrane

16p13.2





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Protein Pathways: Alzheimer's disease, Amyotrophic lateral sclerosis (ALS), Calcium signaling pathway, Long-

term potentiation, Neuroactive ligand-receptor interaction, Systemic lupus erythematosus

MW: 165.3 kDa

Gene Summary: This gene encodes a member of the glutamate-gated ion channel protein family. The

encoded protein is an N-methyl-D-aspartate (NMDA) receptor subunit. NMDA receptors are both ligand-gated and voltage-dependent, and are involved in long-term potentiation, an activity-dependent increase in the efficiency of synaptic transmission thought to underlie certain kinds of memory and learning. These receptors are permeable to calcium ions, and activation results in a calcium influx into post-synaptic cells, which results in the activation of several signaling cascades. Disruption of this gene is associated with focal epilepsy and speech disorder with or without cognitive disability. Alternative splicing results in multiple

transcript variants. [provided by RefSeq, May 2014]