

Product datasheet for RC210390L1V

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

alpha 1 Glycine Receptor (GLRA1) (NM_000171) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: alpha 1 Glycine Receptor (GLRA1) (NM_000171) Human Tagged ORF Clone Lentiviral Particle

Symbol: alpha 1 Glycine Receptor

Synonyms: HKPX1; STHE

Mammalian Cell

Selection:

None

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

 Tag:
 Myc-DDK

 ACCN:
 NM_000171

 ORF Size:
 1347 bp

ORF Nucleotide

Sequence:

The ORF insert of this clone is exactly the same as(RC210390).

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.

RefSeq: <u>NM 000171.1</u>

 RefSeq Size:
 1811 bp

 RefSeq ORF:
 1350 bp

 Locus ID:
 2741

 UniProt ID:
 P23415

 Cytogenetics:
 5q33.1

Protein Families: Druggable Genome, Ion Channels: Cys-loop Receptors, Transmembrane

Protein Pathways: Neuroactive ligand-receptor interaction





alpha 1 Glycine Receptor (GLRA1) (NM_000171) Human Tagged ORF Clone Lentiviral Particle – RC210390L1V

MW: 51.7 kDa

Gene Summary:

The protein encoded by this gene is a subunit of a pentameric inhibitory glycine receptor, which mediates postsynaptic inhibition in the central nervous system. Defects in this gene are a cause of startle disease (STHE), also known as hereditary hyperekplexia or congenital stiff-person syndrome. Multiple transcript variants encoding different isoforms have been found. [provided by RefSeq, Dec 2015]