

OriGene Technologies, Inc.

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Product datasheet for RC205390L2V

GABA A Receptor alpha 1 (GABRA1) (NM_000806) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type:	Lentiviral Particles
Product Name:	GABA A Receptor alpha 1 (GABRA1) (NM_000806) Human Tagged ORF Clone Lentiviral Particle
Symbol:	GABA A Receptor alpha 1
Synonyms:	DEE19; ECA4; EIEE19; EJM; EJM5
Mammalian Cell Selection:	None
Vector:	pLenti-C-mGFP (PS100071)
Tag:	mGFP
ACCN:	NM_000806
ORF Size:	1368 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC205390).
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. <u>More info</u>
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 000806.3</u>
RefSeq Size:	3678 bp
RefSeq ORF:	1371 bp
Locus ID:	2554
UniProt ID:	<u>P14867</u>
Cytogenetics:	5q34
Domains:	Neur_chan_memb, Neur_chan_LBD



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	GABA A Receptor alpha 1 (GABRA1) (NM_000806) Human Tagged ORF Clone Lentiviral Particle – RC205390L2V
Protein Families	: Druggable Genome, Ion Channels: Cys-loop Receptors, Transmembrane
Protein Pathway	/s: Neuroactive ligand-receptor interaction
MW:	51.6 kDa
Gene Summary:	This gene encodes a gamma-aminobutyric acid (GABA) receptor. GABA is the major inhibitory neurotransmitter in the mammalian brain where it acts at GABA-A receptors, which are ligand-gated chloride channels. Chloride conductance of these channels can be modulated by agents such as benzodiazepines that bind to the GABA-A receptor. GABA-A receptors are pentameric, consisting of proteins from several subunit classes: alpha, beta, gamma, delta and rho. Mutations in this gene cause juvenile myoclonic epilepsy and childhood absence epilepsy type 4. Multiple transcript variants encoding the same protein have been identified for this gene. [provided by RefSeq, Jul 2008]

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