

## OriGene Technologies, Inc.

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## Product datasheet for RC212989L3V

## CNGB3 (NM\_019098) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

Product Type:	Lentiviral Particles
Product Name:	CNGB3 (NM_019098) Human Tagged ORF Clone Lentiviral Particle
Symbol:	CNGB3
Synonyms:	ACHM1
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-Myc-DDK-P2A-Puro (PS100092)
Tag:	Myc-DDK
ACCN:	NM_019098
ORF Size:	2427 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC212989).
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 019098.2</u>
RefSeq Size:	4369 bp
RefSeq ORF:	2430 bp
Locus ID:	54714
UniProt ID:	<u>Q9NQW8</u>
Cytogenetics:	8q21.3
Protein Families:	Druggable Genome, Ion Channels: Cyclic nucleotide gated, Transmembrane
MW:	92.2 kDa



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Gene Summary:This gene encodes the beta subunit of a cyclic nucleotide-gated ion channel. The encoded<br/>beta subunit appears to play a role in modulation of channel function in cone<br/>photoreceptors. This heterotetrameric channel is necessary for sensory transduction, and<br/>mutations in this gene have been associated with achromatopsia 3, progressive cone<br/>dystrophy, and juvenile macular degeneration, also known as Stargardt Disease. [provided by<br/>RefSeq, Feb 2010]

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