

## Product datasheet for RC220578L3V

## 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

## CNGB1 (NM\_001297) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** CNGB1 (NM\_001297) Human Tagged ORF Clone Lentiviral Particle

Symbol: CNGB1

Synonyms: CNCG2; CNCG3L; CNCG4; CNGG1; GAR1; GARP; GARP2; RCNC2; RCNCb; RCNCbeta;

RP45

**Mammalian Cell** 

Selection:

Puromycin

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

 Tag:
 Myc-DDK

 ACCN:
 NM\_001297

 ORF Size:
 3753 bp

**ORF Nucleotide** 

Sequence:

Cytogenetics:

The ORF insert of this clone is exactly the same as(RC220578).

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 001297.4</u>

 RefSeq Size:
 5663 bp

 RefSeq ORF:
 3756 bp

 Locus ID:
 1258

 UniProt ID:
 Q14028

**Protein Families:** Druggable Genome, Ion Channels: Cyclic nucleotide gated

16q21



## CNGB1 (NM\_001297) Human Tagged ORF Clone Lentiviral Particle - RC220578L3V

**Protein Pathways:** Olfactory transduction

MW: 139.7 kDa

**Gene Summary:** In humans, the rod photoreceptor cGMP-gated cation channel helps regulate ion flow into the

rod photoreceptor outer segment in response to light-induced alteration of the levels of intracellular cGMP. This channel consists of two subunits, alpha and beta, with the protein encoded by this gene representing the beta subunit. Defects in this gene are a cause of cause of retinitis pigmentosa type 45. Three transcript variants encoding different isoforms have

been found for this gene. [provided by RefSeq, Oct 2013]