

Product datasheet for RC209655L4V

OriGene Technologies, Inc.

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ENSA (NM_207168) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ENSA (NM_207168) Human Tagged ORF Clone Lentiviral Particle

Symbol: ENSA

Synonyms: ARPP-19e

Mammalian Cell Puromycin

Selection:

Vector:

pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_207168

ORF Size: 315 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC209655).

Sequence:
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.

RefSeg: NM 207168.1, NP 997051.1

 RefSeq Size:
 771 bp

 RefSeq ORF:
 318 bp

 Locus ID:
 2029

 UniProt ID:
 043768

 Cytogenetics:
 1q21.3

Protein Families: Druggable Genome

MW: 12 kDa







Gene Summary:

The protein encoded by this gene belongs to a highly conserved cAMP-regulated phosphoprotein (ARPP) family. This protein was identified as an endogenous ligand for the sulfonylurea receptor, ABCC8/SUR1. ABCC8 is the regulatory subunit of the ATP-sensitive potassium (KATP) channel, which is located on the plasma membrane of pancreatic beta cells and plays a key role in the control of insulin release from pancreatic beta cells. This protein is thought to be an endogenous regulator of KATP channels. In vitro studies have demonstrated that this protein modulates insulin secretion through the interaction with KATP channel, and this gene has been proposed as a candidate gene for type 2 diabetes. At least eight alternatively spliced transcript variants encoding distinct isoforms have been observed. [provided by RefSeq, Jul 2008]