

## Product datasheet for RC213645L4V

## OriGene Technologies, Inc.

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## RPS6KA3 (NM\_004586) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: RPS6KA3 (NM 004586) Human Tagged ORF Clone Lentiviral Particle

Symbol: RPS6KA3

Synonyms: CLS; HU-3; ISPK-1; MAPKAPK1B; MRX19; p90-RSK2; pp90RSK2; RSK; RSK2; S6K-alpha3

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_004586 **ORF Size:** 2220 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC213645).

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 004586.2

 RefSeq Size:
 7723 bp

 RefSeq ORF:
 2223 bp

 Locus ID:
 6197

 UniProt ID:
 P51812

 Cytogenetics:
 Xp22.12

**Domains:** pkinase, S\_TK\_X, TyrKc, S\_TKc

**Protein Families:** Druggable Genome, Protein Kinase





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**Protein Pathways:** Long-term potentiation, MAPK signaling pathway, mTOR signaling pathway, Neurotrophin

signaling pathway, Oocyte meiosis, Progesterone-mediated oocyte maturation

**MW:** 83.7 kDa

**Gene Summary:** This gene encodes a member of the RSK (ribosomal S6 kinase) family of serine/threonine

kinases. This kinase contains 2 non-identical kinase catalytic domains and phosphorylates various substrates, including members of the mitogen-activated kinase (MAPK) signalling pathway. The activity of this protein has been implicated in controlling cell growth and differentiation. Mutations in this gene have been associated with Coffin-Lowry syndrome

(CLS). [provided by RefSeq, Jul 2008]