

# Product datasheet for RC206010L2V

### OriGene Technologies, Inc.

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## ULK2 (NM\_014683) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

Product Type: Lentiviral Particles

Product Name: ULK2 (NM 014683) Human Tagged ORF Clone Lentiviral Particle

Symbol: ULK2

**Synonyms:** ATG1B; Unc51.2

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_014683 **ORF Size:** 3108 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC206010).

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 014683.2

 RefSeq Size:
 9165 bp

 RefSeq ORF:
 3111 bp

 Locus ID:
 9706

 UniProt ID:
 Q8IYT8

 Cytogenetics:
 17p11.2

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

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



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**Protein Pathways:** mTOR signaling pathway, Regulation of autophagy

MW: 112.7 kDa

**Gene Summary:** This gene encodes a protein that is similar to a serine/threonine kinase in C. elegans which is

involved in axonal elongation. The structure of this protein is similar to the C. elegans protein in that both proteins have an N-terminal kinase domain, a central proline/serine rich (PS) domain, and a C-terminal (C) domain. The gene is located within the Smith-Magenis syndrome region on chromosome 17. Alternatively spliced transcript variants encoding the

same protein have been identified. [provided by RefSeq, Dec 2008]