

Product datasheet for RC202885L2V

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

LKB1 (STK11) (NM_000455) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: LKB1 (STK11) (NM_000455) Human Tagged ORF Clone Lentiviral Particle

Symbol: LKB1

Synonyms: hLKB1; LKB1; PJS

Mammalian Cell None

Selection:

Vector:

pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_000455 **ORF Size:** 1299 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC202885).

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 000455.4

 RefSeq Size:
 3286 bp

 RefSeq ORF:
 1302 bp

 Locus ID:
 6794

 UniProt ID:
 Q15831

 Cytogenetics:
 19p13.3

Domains: pkinase, TyrKc, S_TKc

Protein Families: Druggable Genome, Protein Kinase





LKB1 (STK11) (NM_000455) Human Tagged ORF Clone Lentiviral Particle - RC202885L2V

Protein Pathways: Adipocytokine signaling pathway, mTOR signaling pathway

MW: 48.5 kDa

Gene Summary: This gene, which encodes a member of the serine/threonine kinase family, regulates cell

polarity and functions as a tumor suppressor. Mutations in this gene have been associated with Peutz-Jeghers syndrome, an autosomal dominant disorder characterized by the growth of polyps in the gastrointestinal tract, pigmented macules on the skin and mouth, and other neoplasms. Alternate transcriptional splice variants of this gene have been observed but have

not been thoroughly characterized. [provided by RefSeq, Jul 2008]