

## **Product datasheet for SC210625**

LKB1 (STK11) (NM 000455) Human 3' UTR Clone

## **Product data:**

**Product Type:** 3' UTR Clones

Product Name: LKB1 (STK11) (NM\_000455) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: STK11

**Synonyms:** hLKB1; LKB1; PJS

**ACCN:** NM\_000455

**Insert Size:** 885 bp

Insert Sequence: >SC210625 3'UTR clone of NM\_000455

The sequence shown below is from the reference sequence of NM\_000455. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).



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## LKB1 (STK11) (NM\_000455) Human 3' UTR Clone - SC210625

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeq:** <u>NM 000455.5</u>

**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]

**Locus ID:** 6794

MW: 32.1