

Product datasheet for **RC402231**

LKB1 (STK11) (NM_000455) Human Mutant ORF Clone

Product data:

Product Type:	Mutant ORF Clones
Product Name:	LKB1 (STK11) (NM_000455) Human Mutant ORF Clone
Mutation Description:	S232P
Affected Codon#:	232
Affected NT#:	694
Nucleotide Mutation:	STK11 Mutant (S232P), Myc-DDK-tagged ORF clone of Homo sapiens serine/threonine kinase 11 (STK11) as transfection-ready DNA
Effect:	Peutz-Jeghers syndrome
Symbol:	STK11
Synonyms:	hLKB1; LKB1; PJS
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_000455
ORF Size:	1299 bp
Restriction Sites:	Sgfi-Mlul



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ORF Nucleotide Sequence:

>RC402231 representing NM_000455
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCC**CGGATCGCC**

ATGGAGGTGGTGGACCCGACGAGCTGGGCATGTTACGGAGGGCGAGCTGATGTGGTGGGTATGGACA
 CGTTCATCCACCGCATCGACTCCACCGAGGTCACTACAGCCGCGCCGCAAGCGGGCCAAGCTCATCGG
 CAAGTACCTGATGGGGGACCTGCTGGGGGAAGGCTCTTACGGCAAGGTGAAGGAGGTGCTGGACTCGGAG
 ACGCTGTGCAGGAGGGCCGCAAGATCCTCAAGAAGAAGAAGTTGCGAAGGATCCCCAACGGGGAGGCCA
 ACGTGAAGAAGGAAATCAACTACTGAGGAGGTTACGGCACAATAATGTCATCCAGCTGGTGGATGTGTT
 ATACAACGAAGAGAAGCAGAAAATGTATATGGTATGGAGTACTGCGTGTGTGGCATGCAGGAAATGCTG
 GACAGCGTGCCGGAAGCGTTCCAGTGTGCCAGGCCACGGGTACTTCTGTCAGCTGATTGACGGCC
 TGGAGTACCTGCATAGCCAGGGCATTGTGCACAAGGACATCAAGCCGGGGAACCTGCTGCTCACCACCGG
 TGGCACCTCAAATCTCGACCTGGGCGTGGCCGAGGCACTGCACCCGTTTCGGCGGGACGACACCTGC
 CGGACCAGCCAGGGCTCCCCGGCTTCCAGCCGCCGAGATTGCCAACGGCTGGACACCTTCCCCGGCT
 TCAAGGTGGACATCTGGTCCGGTGGGGTACCCTCTACAACATCACACGGGTCTGTACCCCTTCGAAGG
 GGACAACATCTACAAGTTGTTTGAACATCGGGAAGGGGAGCTACGCCATCCCGGGCGACTGTGGCCCC
 CCGCTCTCTGACTGTGAAAGGGATGCTTGTAGTACGAACCGGCAAGAGGTTCTCCATCCGGCAGATCC
 GGCAGCACAGCTGGTCCGGAAGAAACATCCTCCGGCTGAAGCACCAGTGCCATCCCACCGAGCCGAGA
 CACCAAGGACCGGTGGCGCAGCATGACTGTGGTGGCGTACTTGGAGGACCTGCACGGCGGGACGAGGAC
 GAGGACCTCTTCGACATCGAGGATGACATCATCTACACTCAGGACTTACGGTGGCCGGACAGGTCACG
 AAGAGGAGGCCAGTCACAATGGACAGCGCCGGGCTCCCCAAGGCCGTGTGTATGAACGGCACAGAGGC
 GGCGCAGCTGAGCACCAATCCAGGGCGAGGGCCGGGCCCCCAACCCCTGCCCGCAAGGCCTGCTCCGCC
 AGCAGCAAGATCCGCCGGCTGTGCGCTGCAAGCAGCAG

AG**CGGACCG**ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

Protein Sequence:

>RC402231 representing NM_000455
 Red=Cloning site Green=Tags(s)

MEVVDPPQLGMFTEGELMSVGMDFIHRIDSTEVIYQPRRKRAKLIGKYLMDLLGEGSYGKVKVLDSE
 TLCRRAVKILKKKLRIPNGEANVKKEIQLLRRLRHKVNIQLVDVLYNEEKQKMYMMEYCVCGMQEML
 DSVPEKRFVPCQAHGYFCQLIDGLEYLHSQIVHKDIKPNLLTTGGTLKISDLGVAEALHPFAADTTC
 RTSQGSAPAFQPPEIANGLDTFPGFKVDIWSAGVTLYNITGLYPFEGDNIYKLFENIGKGSYAIPGDCGP
 PLSDLLKGMLEYEPAKRFSIRQIRQHSWFRKKHPPAEAPVPIPPSPDTKDRWRSMTVVPYLEDLHGADED
 EDLFDIEDDIIYTQDFVPGQVPEEEASHNGQRRGLPKAVCMNGTEAAQLSTKSRAEGRAPNPARKACSA
 SSKIRRLSACKQQ

SGP**TRTRRLEQKLI**SEEDLAANDILDYKDDDDKV

Restriction Sites:

SgfI-MluI

MW: 47.6 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]