

## Product datasheet for **RC402233**

### 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:	W239C
Affected Codon#:	239
Affected NT#:	717
Nucleotide Mutation:	STK11 Mutant (W239C), 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:**

>RC402233 representing NM\_000455  
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGGAGGTGGTGGACCCGACGAGCTGGGCATGTTACGGAGGGCGAGCTGATGTGGTGGGATGGACA  
 CGTTCATCCACCGCATCGACTCCACCGAGGTACTCTACCAGCCGCGCCGAAGCGGGCCAAGCTCATCGG  
 CAAGTACCTGATGGGGGACCTGCTGGGGGAAGGCTCTTACGGCAAGGTGAAGGAGGTGCTGGACTCGGAG  
 ACGCTGTGCAGGAGGGCCGCAAGATCCTCAAGAAGAAGAAGTTGCGAAGGATCCCCAACGGGGAGGCCA  
 ACGTGAAGAAGGAAATCAACTACTGAGGAGGTTACGGCACAAAAATGTCATCCAGCTGGTGGATGTGTT  
 ATACAACGAAGAGAAGCAGAAAATGTATATGGTGTGGAGTACTGCGTGTGTGGCATGCAGGAAATGCTG  
 GACAGCGTGCCGGAAGCGTTCCAGTGTGCCAGGCCACGGGTACTTCTGTCAGCTGATTGACGGCC  
 TGGAGTACCTGCATAGCCAGGGCATTGTGCACAAGGACATCAAGCCGGGGAACCTGCTGCTCACCACCGG  
 TGGCACCTCAAATCTCGACCTGGGCGTGGCCGAGGCACTGCACCCGTTTCGGCGGGACGACACCTGC  
 CGGACCAGCCAGGGCTCCCCGGCTTCCAGCCGCCGAGATTGCCAACGGCTGGACACCTTCTCCGGCT  
 TCAAGGTGGACATCTGCTCGGCTGGGGTCAACCTCTACAACATCACACGGGTCTGTACCCCTTCGAAGG  
 GGACAACATCTACAAGTTGTTTGAACATCGGGAAGGGGAGCTACGCCATCCCGGGCGACTGTGGCCCC  
 CCGCTCTCTGACTGTGAAAGGGATGCTTGAGTACGAACCGCCAAGAGGTTCTCCATCCGGCAGATCC  
 GGCAGCACAGCTGGTCCGGAAGAAACATCCTCCGGCTGAAGCACCAGTGCCATCCCACCGAGCCGAGA  
 CACCAAGGACCGGTGGCGCAGCATGACTGTGGTGCCGACTTGGAGGACCTGCACGGCGGGACGAGGAC  
 GAGGACCTCTTCGACATCGAGGATGACATCATCTACACTCAGGACTTACGGTGCCTGGACAGGTCACG  
 AAGAGGAGGCCAGTCACAATGGACAGCGCCGGGCTCCCCAAGGCGTGTGTATGAACGGCACAGAGGC  
 GGCGCAGCTGAGCACAAATCCAGGGCGAGGGCCGGGCCCCCAACCTGCCCGCAAGGCTGCTCCGCC  
 AGCAGCAAGATCCGCCGCTGTCGGCTGCAAGCAGCAG

AG**CGGACCG**ACGCGTACGCGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC  
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

**Protein Sequence:**

>RC402233 representing NM\_000455  
 Red=Cloning site Green=Tags(s)

MEVVDPPQLGMFTEGELMSVGMDFIHRIDSTEVYQPRRKRAKLIGKYLMDLLGEGSYGKVKVLDSE  
 TLCRRAVKILKKKLRIPNGEANVKKEIQLLRRLRHKNVQLVDVLYNEEKQKMYMMEYCVCGMQEML  
 DSVPEKRFVPCQAHGYFCQLIDGLEYLHSQIVHKDIKPNLLTTGGTLKISDLGVAEALHPFAADTTC  
 RTSQGSAPAFQPPEIANGLDTFSGFKVDICSAGVTLYNITGLYPFEGDNIYKLFENIGKGSYAIPGDCGP  
 PLSDLLKGMLEYEPAKRFSIRQIRQHSWFRKKHPPAEAPVPIPPSPDTKDRWRSMTVVPYLEDLHGADED  
 EDLFDIEDDIIYTQDFVPGQVPEEEASHNGQRRGLPKAVCMNGTEAAQLSTKSRAEGRAPNPARKACSA  
 SSKIRRLSACKQQ

SGP**TRTRRLEQKLI**SEEDLAANDILDYKDDDDKV

**Restriction Sites:**

Sgfl-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]