

Product datasheet for **RC402244**

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: | L285P |
| Affected Codon#: | 285 |
| Affected NT#: | 854 |
| Nucleotide Mutation: | STK11 Mutant (L285P), 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-MluI |



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ORF Nucleotide Sequence:

>RC402244 representing NM_000455
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCC**CGATCGCC**

ATGGAGGTGGTGGACCCGCAGCAGCTGGGCATGTTACGGAGGGCGAGCTGATGTCGGTGGGTATGGACA
 CGTTCATCCACCGCATCGACTCCACCGAGGTACTTACCAGCCGCGCCGAAGCGGGCCAAGCTCATCGG
 CAAGTACCTGATGGGGGACCTGCTGGGGGAAGGCTCTTACGGCAAGGTGAAGGAGGTGCTGGACTCGGAG
 ACGCTGTGCAGGAGGGCCGCAAGATCCTCAAGAAGAAGAAGTTGCGAAGGATCCCCAACGGGGAGGCCA
 ACGTGAAGAAGGAAATCAACTACTGAGGAGGTTACGGCACAAAAATGTCATCCAGCTGGTGGATGTGTT
 ATACAACGAAGAGAAGCAGAAAATGTATATGGTATGGAGTACTGCGTGTGTGGCATGCAGGAAATGCTG
 GACAGCGTGCCGGAAGCGTTCCAGTGTGCCAGGCCACGGGTACTTCTGTCAGCTGATTGACGGCC
 TGGAGTACCTGCATAGCCAGGGCATTGTGCACAAGGACATCAAGCCGGGGAACCTGCTGCTCACCACCGG
 TGGCACCTCAAATCTCCGACCTGGGCGTGGCCGAGGCACTGCACCCGTTTCGGCGGGACGACACCTGC
 CGGACCAGCCAGGGCTCCCCGGCTTCCAGCCGCCGAGATTGCCAACGGCTGGACACCTTCTCCGGCT
 TCAAGGTGGACATCTGGTCCGGTGGGGTACCCTCTACAACATCACACGGGTCTGTACCCCTTCGAAGG
 GGACAACATCTACAAGTTGTTTGAACATCGGGAAGGGGAGCTACGCCATCCCGGGCGACTGTGGCCCC
 CCGCTCTCTGACCCGCTGAAAGGGATGCTTGTGAGTACGAACCGCCAAGAGGTTCTCCATCCGGCAGATCC
 GGCAGCACAGCTGGTCCGGAAGAAACATCCTCCGGCTGAAGCACCAGTGCCATCCCACCGAGCCGAGA
 CACCAAGGACCGGTGGCGCAGCATGACTGTGGTGGCGTACTTGGAGGACCTGCACGGCGGGACGAGGAC
 GAGGACCTCTTCGACATCGAGGATGACATCATCTACACTCAGGACTTACGGTGGCCGGACAGGTCACG
 AAGAGGAGGCCAGTCACAATGGACAGCGCCGGGCTCCCCAAGGCGTGTGTATGAACGGCACAGAGGC
 GGCGCAGCTGAGCACAAATCCAGGGCGAGGGCCGGGCCCCCAACCCCTGCCCGCAAGGCTGCTCCGCC
 AGCAGCAAGATCCGCCGGCTGTGCGCTGCAAGCAGCAG

AG**CGGACCG**ACGCGTACGCGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

Protein Sequence:

>RC402244 representing NM_000455
 Red=Cloning site Green=Tags(s)

MEVVDPPQLGMFTEGELMSVGMDFIHRIDSTEVIYQPRRKRAKLIGKYLMDLLGEGSYGKVKVLDSE
 TLCRRAVKILKKKLRIPNGEANVKKEIQLLRRLRHKVNIQLVDVLYNEEKQKMYMMEYCVCGMQEML
 DSVPEKRFVPCQAHGYFCQLIDGLEYLHSQIVHKDIKPNLLTTGGTLKISDLGVAEALHPFAADDT
 RTSQGSAPAFQPEIANGLDTFSGFKVDIWSAGVTLYNITGLYPFEGDNIYKLFENIGKGSYAIPGDCGP
 PLSPLKGMLEYEPAKRFSIRQIRQHSWFRKKHPPAEAPVPIPPSPDTKDRWRSMTVVPYLEDLHGADED
 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]