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Product datasheet for RC401066

p57 Kip2 (CDKN1C) (NM_000076) Human Mutant ORF Clone

Product data:

Product Type:	Mutant ORF Clones
Product Name:	p57 Kip2 (CDKN1C) (NM_000076) Human Mutant ORF Clone
Mutation Description:	L50P
Affected Codon#:	50
Affected NT#:	149
Nucleotide Mutation:	CDKN1C Mutant (L50P), Myc-DDK-tagged ORF clone of Homo sapiens cyclin-dependent kinase inhibitor 1C (p57, Kip2) (CDKN1C), transcript variant 1 as transfection-ready DNA
Effect:	Beckwith-Wiedemann syndrome
Symbol:	CDKN1C
Synonyms:	BWCR; BWS; KIP2; p57; p57Kip2; WBS
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_000076
ORF Size:	948 bp
Restriction Sites:	Sgfl-RsrII

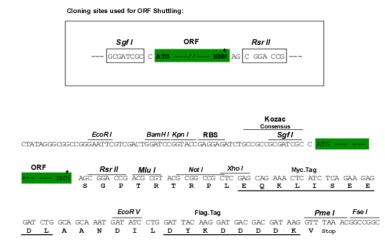


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	p57 Kip2 (CDKN1C) (NM_000076) Human Mutant ORF Clone – RC401066
ORF Nucleotide	>RC401066 representing NM_000076
Sequence:	<pre>Red=Cloning site Blue=ORF Green=Tags(s)</pre>
	TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC GCC <mark>GCGATCGC</mark> C
	ATGTCCGACGCGTCCCTCCGCAGCACATCCACGATGGAGCGTCTTGTCGCCCGTGGGACCTTCCCAGTAC TAGTGCGCACCAGCGCCTGCCGCAGCCTCTTCGGGCCGGTGGACCACGAGGAGCTGAGCCGCAGGCTGCA GGCCCGCCGGCCGAGCTGAACGCCGAGGACCAGAACCGCTGGGATTACGACTTCCAGCAGGACATGCCG CTGCGGGGCCCTGGACGCCTGCAGTGGACCGAAGTGGACAGCGACTCGGTGCCGCGTTCTACCGCGAGA CGGTGCAGGTGGGGGCGCTGCCGCTGCTGCTGGCGCCGCGGCCCGTCGCGGTGGCGGTGCCAGCC GCCCCTCGAGCCGGCCGCTGAGTCCCTCGACGGCCCCGGGCCCGGAGCAGCTGCCTAGTGTCCCG GTCCCGGCCCCGGCGTCCACCCCGCCCCAGTCCCGGTCCTGGCCCCGGCCCCGGCCCCGGCTC CGGTCGCGGCCCCGGCCTCCAGTCGCGGTCCCGGGTCCTGGCCCCGGCCCCGGCCCCGGCTC GGCCCGGCCCCGGCCCCGGCCCCAGTCCCGGTCCCGGCCCCGGCCCCGGCCCCGGCCCCGGC TCCGGCTCCGGCCCCGGACGCGCGCCCAAGAGGCGCGAGCAGCGCGGCACCAGGGCAGCGC GCCCGGCCCCGGCCCCGGACCGCGCCCCAGCCCCGGCCCCGGCCCCGGCCCCGGCCCCGGCCCCG GCCCGGCCCCGGCCCCGGACCGCGCCCCAGCCCGGACCAGGGCGCGGCACCAGGGCAGCGC GCCAGGAGCCTCTCGCTGACCAGCTGCAGCTGCCGGCCTCTGATCTCCGATTTCTTCGCCAAGCGCCGC CAGCGCCAACGGCGCGGCG
	TGGATTACAAGGATGACGACGA TAAG GTTTAA
Protein Sequence	<pre>>RC401066 representing NM_000076 Red=Cloning site Green=Tags(s)</pre>
	MSDASLRSTSTMERLVARGTFPVLVRTSACRSLFGPVDHEELSRELQARPAELNAEDQNRWDYDFQQDMP LRGPGRLQWTEVDSDSVPAFYRETVQVGRCRLLLAPRPVAVAVAVSPPLEPAAESLDGLEEAPEQLPSVP VPAPASTPPPVPVLAPAPAPAPAPVAAPVAAPVAVAVLAPAPAPAPA
	SGPTRTRRLEQKLISEEDLAANDILDYKDDDDKV
Restriction Sites:	Sgfl-RsrII

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Cloning Scheme:



* The last codon before the Stop codon of the ORF

OTI Disclaimer: Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at <u>custsupport@origene.com</u> or by calling 301.340.3188 option 3 for pricing and delivery.

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. <u>More info</u>

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

RefSeq:	<u>NP 000067</u>
RefSeq Size:	948 bp
RefSeq ORF:	951 bp
Locus ID:	1028
Cytogenetics:	11p15.4
Domains:	CDI
Protein Families:	Druggable Genome
Protein Pathways:	Cell cycle

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	p57 Kip2 (CDKN1C) (NM_000076) Human Mutant ORF Clone – RC401066
MW:	34.8 kDa
Gene Summary:	This gene is imprinted, with preferential expression of the maternal allele. The encoded protein is a tight-binding, strong inhibitor of several G1 cyclin/Cdk complexes and a negative regulator of cell proliferation. Mutations in this gene are implicated in sporadic cancers and Beckwith-Wiedemann syndorome, suggesting that this gene is a tumor suppressor candidate. Three transcript variants encoding two different isoforms have been found for this gene. [provided by RefSeq, Oct 2010]

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