

Product datasheet for **RC401071**

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:	S282X
Affected Codon#:	282
Affected NT#:	845
Nucleotide Mutation:	CDKN1C Mutant (S282X), 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:	843 bp
Restriction Sites:	Sgfl-RsrII



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ORF Nucleotide Sequence:

>RC401071 representing NM_000076
 Red=Cloning site Blue=ORF Green=Tags(s)
 TTTTGTAAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCCGCGATCGCC

ATGTCGACGCGTCCCTCCGACGACATCCACGATGGAGCGTCTTGTCGCCCGTGGGACCTTCCCAGTAC
 TAGTGCGCACCAAGCGCCTGCCGACGCTCTCGGGCCGGTGGACCACGAGGAGCTGAGCCGCGAGCTGCA
 GGCCCGCTGGCCGAGCTGAACGCCGAGGACCAGAACCCTGGGATTACGACTTCCAGCAGGACATGCCG
 CTGCGGGGCCCTGGACGCTGCAGTGGACCGAAGTGGACAGCGACTCGGTGCCCGCGTTCTACCGCGAGA
 CGGTGCAGGTGGGGCGCTGCCGCTGCTGCTGGCGCCGCGCCGTCGCGGTGCGGTGGCTGTAGCC
 GCCCTCGAGCCGCCGCTGAGTCCCTCGACGGCTCGAGGAGGCGCCGAGCAGCTGCCTAGTGTCCC
 GTCCCGGCCCGCGTCCACCCGCCCCAGTCCCGGTCTGGCTCCAGCCCGGCCCGGCTCCGGCTC
 CGGTGCGGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGT
 TCCGGTCCGGTCCGGTCCAGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGTCCGGT
 GCCCGGCCCGGCCCGGACGCGCGCCTCAAGAGAGCGCCGAGCAGGGCGCAACAGGGGCAGCGCG
 GCCAGGAGCTCTCGCTGACCAGCTGACTCGGGATTTCGGGACGTCCCGCGGCCGGCACCGCGGCCG
 CAGCGCAACGCGCGCGATCAAGAAGCTGTCCGGCCTCTGATCTCCGATTTCTCGCAAGCGCAAG
 AGA

AGCGGACCGACGCGTACGCGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC
 TGGATTACAAGGATGACGACGA TAAGGTTTAA

Protein Sequence:

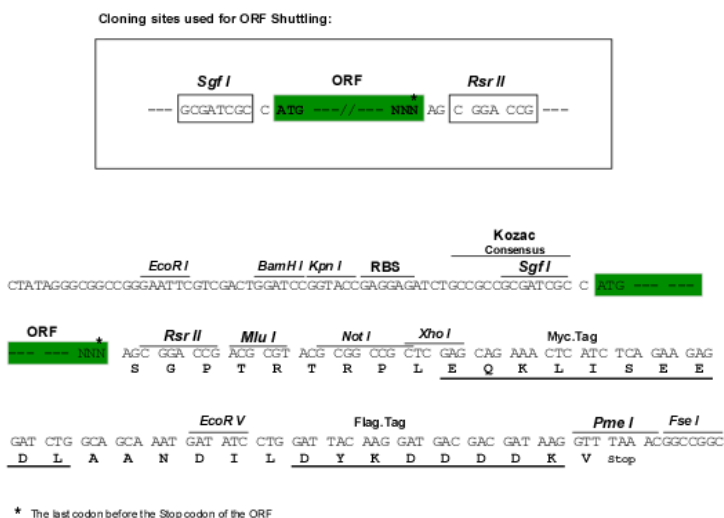
>RC401071 representing NM_000076
 Red=Cloning site Green=Tags(s)
 MSDASLRSTMERLVARGTFPVLVRTSACRSLFGPVDHEELSRELQARLAELNAEDQNRWDYDFQQDMP
 LRGPGRLLQWTEVDSVPAFYRETVQVGRCLLLAPRPVAVAVAVSPPLEPAAESLDGLEEAPEQLPSVP
 VPAPASTPPPVPVLAPAPAPAPVAAPVAAPVAVAVLAPAPAPAPAPAPAPVAAPAPAPAPAPAPAP
 APAPAPDAAPQESAEQGANQQRGQEPLADQLHSGISGRPAAGTAAASANGAAIKKLSGPLISDFFAKRK
 R

SGPTRRRLLEQKLISEEDLAANDILDYKDDDDKV

Restriction Sites:

Sgfl-RsrII

Cloning Scheme:



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 custsupport@origene.com 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. [More info](#)

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:

[NP_000067](#)

RefSeq Size:

843 bp

RefSeq ORF:

951 bp

Locus ID:

1028

Cytogenetics:

11p15.4

Domains:

CDI

Protein Families:

Druggable Genome

Protein Pathways:

Cell cycle

MW: 30.9 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 syndrome, 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]