

Product datasheet for RC402766

p21 Ras (HRAS) (NM_005343) Human Mutant ORF Clone

Product data:

Product Type:	Mutant ORF Clones
Product Name:	p21 Ras (HRAS) (NM_005343) Human Mutant ORF Clone
Mutation Description:	G12S
Affected Codon#:	12
Affected NT#:	34
Nucleotide Mutation:	HRAS Mutant (G12S), Myc-DDK-tagged ORF clone of Homo sapiens v-Ha-ras Harvey rat sarcoma viral oncogene homolog (HRAS), transcript variant 1 as transfection-ready DNA
Effect:	Cosello syndrome
Symbol:	HRAS
Synonyms:	C-BAS/HAS; C-H-RAS; C-HA-RAS1; CTLO; H-RASIDX; HAMSV; HRAS1; p21ras; RASH1
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_005343
ORF Size:	567 bp
Restriction Sites:	SgfI-Mlul

OriGene Technologies, Inc.

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	p21 Ras (HRAS) (NM_005343) Human Mutant ORF Clone – RC402766									
ORF Nucleotide Sequence:	<pre>>RC402766 representing NM_005343 Red=Cloning site Blue=ORF Green=Tags(s) TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC GCCGCGATCGCC</pre>									
	ATGACGGAATATAAGCTGGTGGTGGTGGGGGCGCCAGCGGTGTGGGCAAGAGTGCGCTGACCATCCAGCTGA TCCAGAACCATTTTGTGGACGAATACGACCCCACTATAGAGGATTCCTACCGGAAGCAGGTGGTCATTGA TGGGGAGACGTGCCTGTTGGACATCCTGGATACCGCCGGCCAGGAGGAGTACAGCGCCATGCGGGACCAG TACATGCGCACCGGGGAGGGCTTCCTGTGTGTGTTTGCCATCAACAACACCAAGTCTTTTGAGGACATCC ACCAGTACAGGGAGCAGATCAAACGGGTGAAGGACTCGGATGACGTGCCCATGGTGGTGGGGGAACAA GTGTGACCTGGCTGCACGCACTGTGGAATCTCGGCAGGCTCAGGACCTCGCCCGAAGCTACGGCATCCCC TACATCGAGACCTCGGCCAGGACCGGCAGGGAGTGGAGGATGCCTTCTACACGGTGGGGGAACCA GGCAGCACAAGCTGCGGCAAGACCCGGCAGGGAGTGGAGGATGCCTTCTACACGTTGGTGCGTGAGATCC GGCAGCACAAGCTGCGGAAGCTGAACCCTCCTGATGAGAGTGGCCCCGGCTGCATGAGCTGCAAGTGTGT GCTCTC									
	AGCGGACCG ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC TGGATTACAAGGATGACGACGA TAAG GTTTAA									
Protein Sequence:	<pre>>RC402766 representing NM_005343 Red=Cloning site Green=Tags(s)</pre>									
	MTEYKLVVVGASGVGKSALTIQLIQNHFVDEYDPTIEDSYRKQVVIDGETCLLDILDTAGQEEYSAMRDQ YMRTGEGFLCVFAINNTKSFEDIHQYREQIKRVKDSDDVPMVLVGNKCDLAARTVESRQAQDLARSYGIP YIETSAKTRQGVEDAFYTLVREIRQHKLRKLNPPDESGPGCMSCKCVLS									
	SGP TRTRRLEQKLISEEDLAANDILDYKDDDDKV									
Restriction Sites:	Sgfl-Mlul									
Cloning Scheme:	Cloning sites used for ORF Shuttling: Sgf I ORF Mlu I GCGATCGC C ATG NINN ACG CGT									
	Kozac Consensus EcoRI BamHI Kpn I RBS Sgf1 CTATAGGGGGGGGGGGGAATTOGTOGGACTGGGTCGGGTACGCC ATG									
	ORF <u>Miu i Noti Xhoi</u> Myc.Tag AGU GET ACG GGG GGG CCC CTC GAG CAA CTC ATC TCA GAA GAG T R T R P L E Q K L I S E E									

					Eco	RV				Flag.T	ag					F	me l	Fse I
GAT	CTG	GCA	GCA	AAT	GAT	ATC	CTG	GAT	TAC	AAG	GAT	GAC	GAC	GAT	AAG	GTT	TAA	ACGGCCGGCC
D	L	A	А	N	D	I	L	D	Y	к	D	D	D	D	к	v	Stop	

* The last codon before the Stop codon of the ORF

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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 005334</u>								
RefSeq Size:	567 bp								
RefSeq ORF:	570 bp								
Locus ID:	3265								
Cytogenetics:	11p15.5								
Protein Families:	Druggable Genome								
Protein Pathway	S: Acute myeloid leukemia, Axon guidance, B cell receptor signaling pathway, Bladder cancer, Chemokine signaling pathway, Chronic myeloid leukemia, Endocytosis, Endometrial cancer, ErbB signaling pathway, Fc epsilon RI signaling pathway, Focal adhesion, Gap junction, Glioma, GnRH signaling pathway, Insulin signaling pathway, Long-term depression, Long-term potentiation, MAPK signaling pathway, Melanogenesis, Melanoma, Natural killer cell mediated cytotoxicity, Neurotrophin signaling pathway, Non-small cell lung cancer, Pathways in cancer, Prostate cancer, Regulation of actin cytoskeleton, Renal cell carcinoma, T cell receptor signaling pathway, Thyroid cancer, Tight junction, VEGF signaling pathway								
MW:	20.8 kDa								

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CRIGENE p21 Ras (HRAS) (NM_005343) Human Mutant ORF Clone – RC402766

Gene Summary: This gene belongs to the Ras oncogene family, whose members are related to the transforming genes of mammalian sarcoma retroviruses. The products encoded by these genes function in signal transduction pathways. These proteins can bind GTP and GDP, and they have intrinsic GTPase activity. This protein undergoes a continuous cycle of de- and re-palmitoylation, which regulates its rapid exchange between the plasma membrane and the Golgi apparatus. Mutations in this gene cause Costello syndrome, a disease characterized by increased growth at the prenatal stage, growth deficiency at the postnatal stage, predisposition to tumor formation, cognitive disability, skin and musculoskeletal abnormalities, distinctive facial appearance and cardiovascular abnormalities. Defects in this gene are implicated in a variety of cancers, including bladder cancer, follicular thyroid cancer, and oral squamous cell carcinoma. Multiple transcript variants, which encode different isoforms, have been identified for this gene. [provided by RefSeq, Jul 2008]

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