

## Product datasheet for **RC402273**

### G protein alpha S (GNAS) (NM\_000516) Human Mutant ORF Clone

#### Product data:

Product Type:	Mutant ORF Clones
Product Name:	G protein alpha S (GNAS) (NM_000516) Human Mutant ORF Clone
Mutation Description:	R160C
Affected Codon#:	160
Affected NT#:	478
Nucleotide Mutation:	GNAS Mutant (R160C), Myc-DDK-tagged ORF clone of Homo sapiens GNAS complex locus (GNAS), transcript variant 1 as transfection-ready DNA
Effect:	Albrih herediry oseodysrophy
Symbol:	GNAS
Synonyms:	AHO; C20orf45; GNAS1; GPSA; GSA; GSP; NESP; PITA3; POH; SCG6; SgVI
E. coli Selection:	Kanamycin (25 ug/mL)
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
Tag:	Myc-DDK
ACCN:	NM_000516
ORF Size:	1182 bp
Restriction Sites:	Sgfl-NotI



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ORF Nucleotide  
Sequence:

>RC402273 representing NM\_000516  
Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
GCCCGGATCGCC

ATGGGCTGCCTCGGGAACAGTAAGACCGAGGACCAGCGCAACGAGGAGAAGGCGCAGCGTGAGGCCAACA  
AAAAGATCGAGAAGCAGCTGCAGAAGGACAAGCAGGTCTACCGGCCACGCACCGCCTGCTGCTGCTGGG  
TGCTGGAGAATCTGGTAAAAGCACCAATTGTGAAGCAGATGAGGATCCTGCATGTTAATGGTTTTAATGGA  
GAGGGCGGCGAAGAGGACCCGCAGGCTGCAAGGAGCAACAGCGATGGTGAGAAGGCAACCAAAAGTGCAGG  
ACATCAAAAACAACTGAAAGAGGGCATTGAAACCATTGTGGCCGCCATGAGCAACCTGGTGCCCCCGT  
GGAGCTGGCAACCCCGAGAACCAGTTCAAGTGGACTACATCCTGAGTGTGATGAACGTGCCTGACTTT  
GACTTCCTCCCGAATTCATGAGCATGCCAAGGCTCTGTGGGAGGATGAAGGAGTGTGTGCCTGCTACG  
AACGCTCAACGAGTACCAGCTGATTGACTGTGCCAGTACTTCTGGACAAGATCGACGTGATCAAGCA  
GGCTGACTATGTGCCGAGCGATCAGGACCTGCTTCGCTGCCGTGCTCCTGACTTCTGGAATCTTTGAGACC  
AAGTTCAGGTGGACAAAGTCAACTTCCACATGTTTGACGTGGGTGGCCAGCGCATGAACGCCGCAAGT  
GGATCCAGTGTCTCAACGATGTGACTGCCATCATCTTCGTGGTGGCCAGCAGCAGCTACAACATGGTCAT  
CCGGGAGGACAACCAGACCAACCGCCTGCAGGAGGCTCTGAACCTCTCAAGAGCATCTGGAACAACAGA  
TGGCTGCGCACCATCTCTGTGATCCTGTTCTCAACAAGCAAGATCTGCTCGCTGAGAAAGTCTTTGCTG  
GGAAATCGAAGATTGAGGACTACTTCCAGAATTTGCTCGCTACACTACTCCTGAGGATGCTACTCCCGA  
GCCCGGAGAGGACCCACGCGTGACCCGGGCCAAGTACTTATTCGAGATGAGTTTCTGAGGATCAGCACT  
GCCAGTGGAGATGGGCGTCACTACTGCTACCCTCATTTACCTGCGCTGTGGACTGAGAACATCCGCC  
GTGTGTTCAACGACTGCCGTGACATCATTACGGCATGCACCTTCGTCACTACGAGCTGCTC

AGCGGACCGACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCC  
TGGATTACAAGGATGACGACGA TAAGGTTTAA

## Protein Sequence:

>RC402273 representing NM\_000516  
Red=Cloning site Green=Tags(s)

MGCLGNSKTEDQRNEEKAQREANKKIEKQLQKDKQVYRATHRLLLLGAGESGKSTIVKQMRILHVNGFNG  
EGGEEDPQAARSNSDGEKATKVQDIKNNLKEAIETIVAAMSNLVPPVELANPENQFRVDYILSVMNVPDF  
DFPPEFYEHAKALWEDEGVCACYERSNEYQLIDCAQYFLDKIDVIKQADYVPSDQLLRVLTSGIFET  
KFQVDKVNFMFDVGGQRDERRKIQCNDVTAIIFVVASSSYNMVIREDNQTNRLQEALNLFKSIWNNR  
WLRTISVILFLNKQDLLAEKVLGKSKIEDYFPEFARYTTPEDATPEPGEPRVTRAKYFIRDEFRLIST  
ASGDGRHYCYPHFTCAVDTENIRRVFNDCRDI IQRMHLRQYELL

SGPTRRRLEQKLI SEEDLAANDILDYKDDDDKV

## Restriction Sites:

Sgfl-NotI



<b>Protein Pathways:</b>	Calcium signaling pathway, Dilated cardiomyopathy, Gap junction, GnRH signaling pathway, Long-term depression, Melanogenesis, Taste transduction, Vascular smooth muscle contraction, Vibrio cholerae infection
<b>MW:</b>	43.3 kDa
<b>Gene Summary:</b>	<p>This locus has a highly complex imprinted expression pattern. It gives rise to maternally, paternally, and biallelically expressed transcripts that are derived from four alternative promoters and 5' exons. Some transcripts contain a differentially methylated region (DMR) at their 5' exons, and this DMR is commonly found in imprinted genes and correlates with transcript expression. An antisense transcript is produced from an overlapping locus on the opposite strand. One of the transcripts produced from this locus, and the antisense transcript, are paternally expressed noncoding RNAs, and may regulate imprinting in this region. In addition, one of the transcripts contains a second overlapping ORF, which encodes a structurally unrelated protein - Alex. Alternative splicing of downstream exons is also observed, which results in different forms of the stimulatory G-protein alpha subunit, a key element of the classical signal transduction pathway linking receptor-ligand interactions with the activation of adenylyl cyclase and a variety of cellular responses. Multiple transcript variants encoding different isoforms have been found for this gene. Mutations in this gene result in pseudohypoparathyroidism type 1a, pseudohypoparathyroidism type 1b, Albright hereditary osteodystrophy, pseudopseudohypoparathyroidism, McCune-Albright syndrome, progressive osseous heteroplasia, polyostotic fibrous dysplasia of bone, and some pituitary tumors. [provided by RefSeq, Aug 2012]</p>