

Product datasheet for AR09416PU-N

HRAS (1-186, His-tag) Human Protein

Product data:

OriGene Technologies, Inc.

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Product Type:	Recombinant Proteins	
Description:	HRAS (1-186, His-tag) human recombinant protein, 0.1 mg	
Species:	Human	
Expression Host:	E. coli	
Expression cDNA Clone or AA Sequence:	MTEYKLVVVG AGGVGKSALT IQLIQNHFVD EYDPTIEDSY RKQVVIDGET CLLDILDTAG QEEYSAMRDQ YMRTGEGFLC VFAINNTKSF EDIHQYREQI KRVKDSDDVP MVLVGNKCDL AARTVESRQA QDLARSYGIP YIETSAKTRQ GVEDAFYTLV REIRQHKLRK LNPPDESGPG CMSCKC <u>LEHH HHHH</u>	
Tag:	His-tag	
Concentration:	lot specific	
Purity:	>90% by SDS - PAGE	
Buffer:	Presentation State: Purified State: Liquid purified protein Buffer System: Tris-HCl buffer (pH 8.0) containing 20% glycerol, 0.1 M NaCl	
Preparation:	Liquid purified protein	
Protein Description:	Recombinant human HRAS protein, fused to His-tag, was expressed in E.coli and purified by using conventional chromatography techniques.	
Storage:	Store undiluted at 2-8°C for up to two weeks or (in aliquots) at -20°C or -70°C for longer. Avoid repeated freezing and thawing.	
Stability:	Shelf life: one year from despatch.	
RefSeq:	<u>NP 001123914</u>	
Locus ID:	3265	
UniProt ID:	<u>P01112</u>	
Cytogenetics:	11p15.5	
Synonyms:	GTPase HRas, HRAS1, p21ras, H-Ras-1, c-H-ras, Ha-Ras, H-Ras	



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🖢 ORÏGENE 🛛 HRAS (1-186, His-tag) Human Protein – AR09416PU-N

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]

Protein Families: Druggable Genome

Protein Pathways: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

Product images:

(kDa) 70 57	-
40	-
28	-
18	-
13.5	-
8.5	

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