

OriGene Technologies, Inc.

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Product datasheet for PH313692

SAR1B (NM_001033503) Human Mass Spec Standard

Product data:

Description: SAR1B MS Standard C13 and N15-labeled recombinant protein (NP_001028675) Species: Lluman	
Charles Human	
Species: Human	
Expression Host: HEK293	
Expression cDNA Clone RC213692 or AA Sequence:	
Predicted MW: 22.4 kDa	
Protein Sequence:>RC213692 protein sequenceRed=Cloning site Green=Tags(s)	
MSFIFDWIYSGFSSVLQFLGLYKKTGKLVFLGLDNAGKTTLLHMLKDDRLGQHVPTLHPTSEELTIAGMT FTTFDLGGHVQARRVWKNYLPAINGIVFLVDCADHERLLESKEELDSLMTDETIANVPILILGNKIDRPE AISEERLREMFGLYGQTTGKGSISLKELNARPLEVFMCSVLKRQGYGEGFRWMAQYID	
TRTRPLEQKLISEEDLAANDILDYKDDDDKV	
Tag:C-Myc/DDK	
Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining	
Concentration: >0.05 μg/μL as determined by microplate BCA method	
Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine	
Buffer:25 mM Tris-HCl, 100 mM glycine, pH 7.3	
Storage:Store at -80°C. Avoid repeated freeze-thaw cycles.	
Stability: Stable for 3 months from receipt of products under proper storage and handling	conditions.
RefSeq: NP 001028675	
RefSeq Size: 6651	
RefSeq ORF:594	
Synonyms: ANDD; CMRD; GTBPB; SARA2	
Locus ID: 51128	
UniProt ID: <u>Q9Y6B6</u>	



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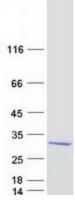
Cytogenetics:

5q31.1

Summary:

The protein encoded by this gene is a small GTPase that acts as a homodimer. The encoded protein is activated by the guanine nucleotide exchange factor PREB and is involved in protein transport from the endoplasmic reticulum to the Golgi. This protein is part of the COPII coat complex. Defects in this gene are a cause of chylomicron retention disease (CMRD), also known as Anderson disease (ANDD). Two transcript variants encoding the same protein have been found for this gene. [provided by RefSeq, Mar 2010]

Product images:



Coomassie blue staining of purified SAR1B protein (Cat# [TP313692]). The protein was produced from HEK293T cells transfected with SAR1B cDNA clone (Cat# [RC213692]) using MegaTran 2.0 (Cat# [TT210002]).

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