

Product datasheet for AR09849PU-N

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OriGene Technologies, Inc.

CIB2 (1-187, His-tag) Human Protein

Product data:

Product Type: Recombinant Proteins

Description: CIB2 (1-187, His-tag) human recombinant protein, 0.1 mg

Species: Human
Expression Host: E. coli

Expression cDNA Clone MGSSHHHHHH SSGLVPRGSH MGNKQTIFTE EQLDNYQDCT FFNKKDILKL HSRFYELAPN

or AA Sequence: LVPMDYRKSP IVHVPMSLII QMPELRENPF KERIVAAFSE DGEGNLTFND FVDMFSVLCE

SAPRELKANY AFKIYDFNTD NFICKEDLEL TLARLTKSEL DEEEVVLVCD KVIEEADLDG DGKLGFADFE

DMIAKAPDFL STFHIRI

Tag: His-tag

Predicted MW: 23.8 kDa

Concentration: lot specific

Purity: >90%

Buffer: Presentation State: Purified

State: Liquid purified protein

Buffer System: 20 mM Tris-HCl buffer (pH 8.0) containing 10% glycerol, 2 mM DTT, 100 mM

NaCl, 1 mM PMSF

Preparation: Liquid purified protein

Protein Description: Recombinant human CIB2 protein, fused to His-tag at N-terminus, 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 001258817</u>

Locus ID: 10518

 UniProt ID:
 O75838

 Cytogenetics:
 15q25.1

Synonyms: DFNB48; KIP2; USH1J





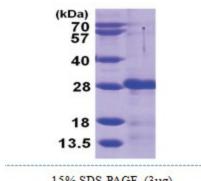
Summary:

The protein encoded by this gene is similar to that of KIP/CIB, calcineurin B, and calmodulin. The encoded protein is a calcium-binding regulatory protein that interacts with DNAdependent protein kinase catalytic subunits (DNA-PKcs), and it is involved in photoreceptor cell maintenance. Mutations in this gene cause deafness, autosomal recessive, 48 (DFNB48), and also Usher syndrome 1J (USH1J). Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2014]

Protein Families:

Druggable Genome

Product images:



15% SDS-PAGE (3ug)