

PRODUCT INFORMATION

Tag C-Flag Tag **Target** CACB4

Synonyms CAB4, CACNLB4, EA5, EIG9, EJM, EJM4, EJM6 Human CACB4 full length protein-synthetic **Description**

nanodisc **Delivery** 6~8weeks **Uniprot ID** O00305 **Expression Host HEK293**

Ion Channels: Other **Protein Families**

Protein Pathways N/A

Formulation &

Reconstitution

Storage & Shipping

Background

The human full length CACB4 protein has a MW of **Molecular Weight**

58.2kDa Lyophilized from nanodisc solubilization buffer (20

mM Tris-HCl, 150 mM NaCl, pH 8.0). Normally 5% – 8% trehalose is added as protectants before lyophilization. Please see Certificate of Analysis

for specific instructions. Do not use solvents with a pH below 6.5 or those containing high concentrations of divalent metal ions (greater than 5 mM) in subsequent experiments. Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not

intended for use within a month, aliquot and store at -80°C (Avoid repeated freezing and thawing). Lyophilized proteins are shipped at ambient

témperature.

This gene encodes a member of the beta subunit family of voltage-dependent calcium channel complex proteins. Calcium channels mediate the

influx of calcium ions into the cell upon

membrane polarization and consist of a complex of alpha-1, alpha-2/delta, beta, and gamma subunits in a 1:1:1:1 ratio. Various versions of each of these subunits exist, either expressed from similar genes or the result of alternative splicing. The protein encoded by this locus plays

an important role in calcium channel function by modulating G protein inhibition, increasing peak calcium current, controlling the alpha-1 subunit membrane targeting and shifting the voltage dependence of activation and inactivation. Certain mutations in this gene have been associated with idiopathic generalized epilepsy (IGE), juvenile myoclonic epilepsy (JME), and episodic ataxia, type 5. [provided by RefSeq, Aug

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2016]

Usage Research use only Conjugate Unconjugated

