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Product datasheet

Recombinant Human Dynamin 2 protein ab132947

1 Image

Description

Product name Recombinant Human Dynamin 2 protein

Expression system Wheat germ
Accession P50570

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MGNRGMEELIPLVNKLQDAFSSIGQSCHLDLPQIAVVGGQ

SAGKSSVLEN

FVGRDFLPRGSGIVTRRPLILQLIFSKTEHAEFLHCKSKKFT

DFDEVRQE

IEAETDRVTGTNKGISPVPINLRVYSPHVLNLTLIDLPGITKV

PVGDQPP

DIEYQIKDMILQFISRESSLILAVTPANMDLANSDALKLAKE

VDPQGLRT

IGVITKLDLMDEGTDARDVLENKLLPLRRGYIGVVNRSQKDI

EGKKDIRA

ALAAERKFFLSHPAYRHMADRMGTPHLQKTLNQQLTNHIR

ESLPALRSKL

QSQLLSLEKEVEEYKNFRPDDPTRKTKALLQMVQQFGVD

FEKRIEGSGDQ

VDTLELSGGARINRIFHERFPFELVKMEFDEKDLRREISYAI

KNIHGVRT

GLFTPDLAFEAIVKKQVVKLKEPCLKCVDLVIQELINTVRQ

CTSKLSSYP

RLREETERIVTTYIREREGRTKDQILLLIDIEQSYINTNHEDFIG

FANAQ

QRSTQLNKKRAIPNQGEILVIRRGWLTINNISLMKGGSKEYW

FVLTAESL

SWYKDEEEKEKKYMLPLDNLKIRDVEKGFMSNKHVFAIFN

TEQRNVYKDL

RQIELACDSQEDVDSWKASFLRAGVYPEKDQAENEDGA

QENTFSMDPQLE

 ${\sf RQVETIRNLVDSYVAIINKSIRDLMPKTIMHLMINNTKAFIHHE}$

LLAYLY

I

SSADQSSLMEESADQAQRRDDMLRMYHALKEALNIIGDIS

TSTVSTPVPP

PVDDTWLQSASSHSPTPQRRPVSSIHPPGRPPAVRGPTP

GPPLIPVPVGA

AASFSAPPIPSRPGPQSVFANSDLFPAPPQIPSRPVRIPP

GIPPGVPSRR PPAAPSRPTIIRPAEPSLLD

Predicted molecular weight

125 kDa including tags

Amino acids

1 to 870

Specifications

Our Abpromise quarantee covers the use of ab132947 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications

ELISA

Western blot SDS-PAGE

Form

Liquid

Additional notes

Preparation and Storage

Stability and Storage

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.31% Glutathione, 0.79% Tris HCI

General Info

Function

Microtubule-associated force-producing protein involved in producing microtubule bundles and able to bind and hydrolyze GTP. Most probably involved in vesicular trafficking processes, in particular endocytosis.

Tissue specificity

Ubiquitously expressed.

Involvement in disease

Defects in DNM2 are a cause of centronuclear myopathy autosomal dominant (ADCNM) [MIM:160150]; also known as autosomal dominant myotubular myopathy. Centronuclear myopathies (CNMs) are congenital muscle disorders characterized by progressive muscular weakness and wasting involving mainly limb girdle, trunk, and neck muscles. It may also affect distal muscles. Weakness may be present during childhood or adolescence or may not become evident until the third decade of life. Ptosis is a frequent clinical feature. CNMs comprise a wide spectrum of phenotypes, ranging from severe neonatal to mild late-onset familial forms. The most prominent histopathologic features include high frequency of centrally located nuclei in muscle fibers not secondary to regeneration, radial arrangement of sarcoplasmic strands around the central nuclei, and predominance and hypotrophy of type 1 fibers.

Defects in DNM2 are the cause of Charcot-Marie-Tooth disease dominant intermediate type B (CMTDIB) [MIM:606482]. Charcot-Marie-Tooth disease (CMT) is a clinically and genetically heterogeneous disorder of the peripheral nervous system, characterized by progressive

weakness and atrophy, initially of the peroneal muscles and later of the distal muscles of the arms. CMTDIB is a form of Charcot-Marie-Tooth disease characterized by clinical and pathologic features intermediate between demyelinating and axonal peripheral neuropathies, and motor

median nerve conduction velocities ranging from 25 to 45 m/sec.

Sequence similarities Belongs to the dynamin family.

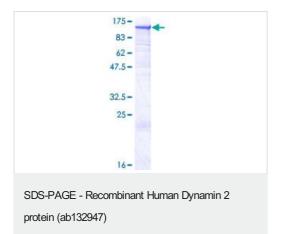
Contains 1 GED domain. Contains 1 PH domain.

Cytoplasm. Cytoplasm > cytoskeleton. Cell junction > synapse > postsynaptic cell membrane >

postsynaptic density. Cell junction > synapse. Microtubule-associated. Also found in the

postsynaptic density of neuronal cells.

Images



12.5% SDS-PAGE analysis of ab132947 stained with Coomassie Blue

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