abcam

Product datasheet

Recombinant Human Ephrin B1 protein (Fc Chimera His Tag) ab235877

1 Image

Description

Product name Recombinant Human Ephrin B1 protein (Fc Chimera His Tag)

Purity > 90 % SDS-PAGE.

Affinity purified

Endotoxin level < 1.000 Eu/µg

Expression system Baculovirus infected insect cells

Accession P98172

Protein length Protein fragment

Animal free No

Nature Recombinant

Species Human

Sequence ADPLAKNLEPVSWSSLNPKFLSGKGLVIYPKIGDKLDIICP

RAEAGRPYE

YYKLYLVRPEQAAACSTVLDPNVLVTCNRPEQEIRFTIKFQ

EFSPNYMGL

 ${\tt EFKKHHDYYITSTSNGSLEGLENREGGVCRTRTMKIIMKVG}$

QDPNAVTPE

QLTTSRPSKEADNTVKMATQAPGSRGSLGDSDGKHETV

NQEEKSGPGASG

GSSGDPDGFFNSKLEPKSCDKTHTCPPCPAPELLGGPS

VFLFPPKPKDTL

MISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK

PREEQYNSTYR

VVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG

QPREPQVYTL

PPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENN

YKTTPPVLDSD

GSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKS

LSLSPGKHHHH HH

Predicted molecular weight 50 kDa including tags

Amino acids 28 to 237

Tags His tag C-Terminus

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Additional sequence information Fused to hlgG-His-tag at C-terminus. NP 004420. Extracellular domain.

Specifications

Our Abpromise guarantee covers the use of ab235877 in the following tested applications.

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

Applications SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

pH: 7.40

Constituents: PBS, 10% Glycerol (glycerin, glycerine)

General Info

Function Binds to the receptor tyrosine kinases EPHB1 and EPHA1. Binds to, and induce the collapse of,

commissural axons/growth cones in vitro. May play a role in constraining the orientation of

longitudinally projecting axons.

Tissue specificity Heart, placenta, lung, liver, skeletal muscle, kidney, pancreas.

Involvement in disease Defects in EFNB1 are a cause of craniofrontonasal syndrome (CFNS) [MIM:304110]; also known

as craniofrontonasal dysplasia (CFND). CFNS is an X-linked inherited syndrome characterized by hypertelorism, coronal synostosis with brachycephaly, downslanting palpebral fissures, clefting of the nasal tip, joint anomalies, longitudinally grooved fingernails and other digital anomalies.

Sequence similarities Belongs to the ephrin family.

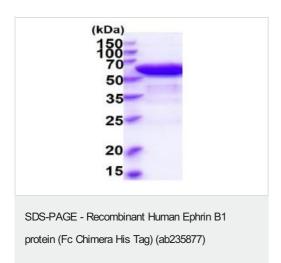
Post-translational

modifications

Inducible phosphorylation of tyrosine residues in the cytoplasmic domain.

Cellular localization Membrane.

Images



15% SDS-PADE analysis of 3 µg ab235877.

50-70 kDa (Under reducing conditions)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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