

# 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	<u>P98172</u>
Protein length	Protein fragment
Animal free	No
Nature	Recombinant
Species	Human
Sequence	ADPLAKNLEPVSWSSLNPKFLSGKGLVIYPKIGDKLDIICP RAEAGRPYE YYKLYLVRPEQAAACSTVLDPNVLVTCNRPEQEIRFTIKFQ EFSPNYMGL EFKKHHDYYITSTSNGLSLEGLNREGGVCRTMTKIMKVG QDPNAVTPPE QLTTSRPSKEADNTVKMATQAPGSRGSLGDSGKHETV NQEEKSGPGASG GSSGDPDGGFNSKLEPKSCDKTHTCPPCPAPELLGGPS VFLFPPKPKDTL MISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYR VVSVELTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTL PPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENN YKTTTPVLDSD GSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKS LSLSPGKHHHH HH
Predicted molecular weight	50 kDa including tags
Amino acids	28 to 237
Tags	His tag C-Terminus

**Additional sequence information** Fused to hlgG-His-tag at C- terminus. NP\_004420. Extracellular domain.

## Specifications

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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

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**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

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**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

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15% SDS-PAGE 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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