

Product datasheet

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

1 Image

Description

<b>Product name</b>	Recombinant Human Ephrin B1 protein (Fc Chimera His Tag)	
<b>Purity</b>	> 90 % SDS-PAGE.	
<b>Endotoxin level</b>	< 1.000 Eu/μg	
<b>Expression system</b>	Baculovirus infected insect cells	
<b>Accession</b>	<a href="#">P98172</a>	
<b>Protein length</b>	Protein fragment	
<b>Animal free</b>	No	
<b>Nature</b>	Recombinant	
<b>Species</b>	Human	
<b>Sequence</b>	<pre> ADPLAKNLEPVSWSLNPFLSGKGLVIYPKIGDKLDII CPRAEAGRPYE YYKLYLVRPEQAAACSTVLDPNVLVTCNRPEQEIRFTIK FQEFSPNYMGL EFKKHHDYYITSTSNGLSLEGLNREGGVCRTTRTMKIIMK VGQDPNAVTPPE QLTTSRPSKEADNTVKMATQAPGSRGSLGSDSGKHE TVNQEEKSGPGASG GSSGDPDGFNFNSKLEPKSCDKTHTCPPCPAPELLGG PSVFLFPPKPKDTL MISRTPEVTCVWVDVSHEDPEVKFNWYVDGVEVHNA KTKPREEQYNSTYR VVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTL PPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPE NNYKTTTPVLDS GSFFLYSKLTVDKSRWQQGNVFCFSVMHEALHNHYT QKSLSLSPGKHHHH HH </pre>	
<b>Predicted molecular weight</b>	50 kDa including tags	
<b>Amino acids</b>	28 to 237	
<b>Tags</b>	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: 7.4

Constituents: PBS, 10% Glycerol

## 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.

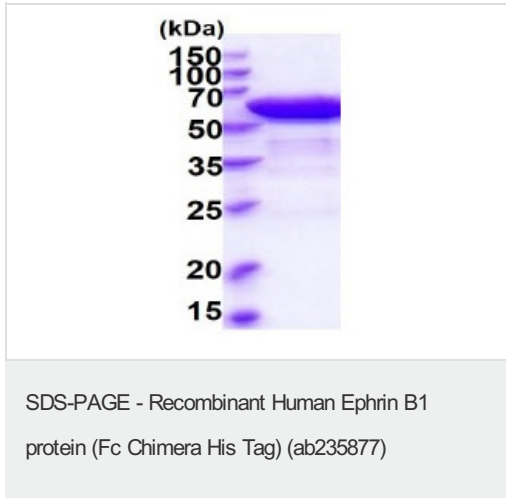
**Post-translational modifications** Inducible phosphorylation of tyrosine residues in the cytoplasmic domain.

**Cellular localization** Membrane.

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## 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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