

Product datasheet

Recombinant Human Acid Phosphatase 2 protein (denatured) ab202151

[1 Image](#)

Description

Product name	Recombinant Human Acid Phosphatase 2 protein (denatured)	
Purity	> 85 % SDS-PAGE.	
Expression system	Escherichia coli	
Accession	<u>P11117</u>	
Protein length	Protein fragment	
Animal free	No	
Nature	Recombinant	
Species	Human	
Sequence	MGSSHHHHHHSSGLVPRGSHMGSRLRFVTLRYRHGDR SPVKTYPKDPYQ EEEWPQGFQQLTKEGMLQHWELGQALRQRYHGFLNTSY HRQEVYVRSTDF DRTLMSAEANLAGLFPPNGMQRFNPNISWQPIPVHTVPIT EDRLKFPPLG PCPRYEQQLNETRQTPEYQNESSRNAQFLDMVANETGLT DLTLETVWNVY DTLFCEQTHGLRLPPWASPQTMQRLSRLKDFSFRFLFGIY QQA EKARLQG GVLLAQIRKNLTLMATTSQLPKLLVYSAHD TTLVALQMALD VYNGEQAPY ASCHFELYQEDSGNFSVEMYFRNESDKAPWPLSLPGCP HRCPLQDFLRL TEPVVPKDWQQECQLASGPADTE	
Predicted molecular weight	43 kDa including tags	
Amino acids	31 to 380	
Tags	His tag N-Terminus	
Additional sequence information	Lumenal domain (NP_001601).	
Description	Recombinant Human Acid Phosphatase 2 protein	

Specifications

Our **Abpromise guarantee** covers the use of **ab202151** 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: 8.00

Constituents: 10% Glycerol (glycerin, glycerine), 0.32% Tris HCl

General Info

Involvement in disease Defects in ACP2 are a cause of acid phosphatase deficiency (ACPHD) [MIM:200950]. The clinical features are intermittent vomiting, hypotonia, lethargy, opisthotonos, terminal bleeding, and death in early infancy. Lysosomal acid phosphatase is deficient in cultured fibroblasts and multiple tissues.

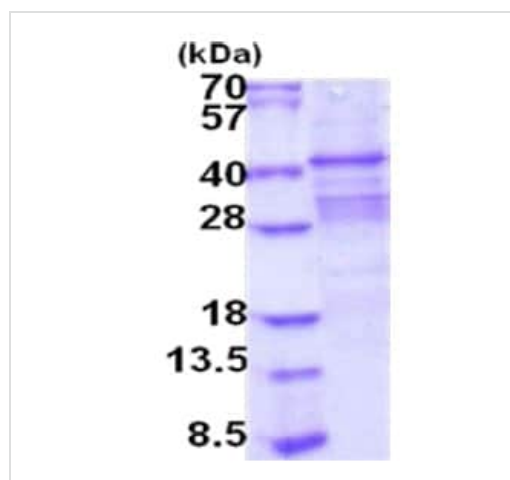
Sequence similarities Belongs to the histidine acid phosphatase family.

Post-translational modifications The membrane-bound form is converted to the soluble form by sequential proteolytic processing. First, the C-terminal cytoplasmic tail is removed. Cleavage by a lysosomal protease releases the soluble form in the lysosome lumen.

N-glycosylated. The intermediates formed during enzymatic deglycosylation suggest that all eight predicted N-glycosylation sites are used.

Cellular localization Lysosome membrane. Lysosome lumen. The soluble form arises by proteolytic processing of the membrane-bound form.

Images



15% SDS-PAGE analysis of ab202151 (3 µg).

SDS-PAGE - Recombinant Human Acid

Phosphatase 2 protein (denatured) (ab202151)

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

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