

Product datasheet

Recombinant Human Protective protein/Cathepsin A (PPCA) ab159166

[1 Image](#)

Overview

Product name Recombinant Human Protective protein/Cathepsin A (PPCA)
Protein length Full length protein

Description

Nature Recombinant
Source Wheat germ
Amino Acid Sequence
Species Human

Sequence

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MIRAAPPPLFLLLLLLLLLLSWASRGEAAPDQDEIQRL
PGLAKQPSFRQY
SGYLKSGSGSKHLHYWFVESQKDPENSPVVLWLNGGP
GCSSLDGLLTHEGP
FLVQPDGVTLEYNPYSWNLIANVLYLESPAGVGFSYSYD
DKFYATNDTEVA
QSNFEALQDFFRLFPEYKNNKFLTGESYAGMPTLAV
LVMQDPSMNLQ
GLAVGNGLSSYEQNDNSLVYFAYYHGLLGNRLWSSLQ
THCCSQNKCNFYD
NKDLECVTNLQEVARVGNISGLNINLYAPCAGGVPSH
FRYEKDTVVVQD
LGNIFTRLPLKRMWHQALLRSGDKVRMDPPCTNTTAA
STYLNNPYVRKAL
NIPEQLPQWDMCNFLVNLQYRRLYRSMNSQYLKLLSS
QKYQILLYNGDVD
MACNFMGDEWFVDSL NQKMEVQRRPWLVKYGDSGE
QIAGFVKEFSHIAFL
TIKAGAHMVPTDKPLAAFTMFSRFLNKQPY
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Amino acids 1 to 480
Tags proprietary tag N-Terminus

Specifications

Our [Abpromise guarantee](#) covers the use of **ab159166** 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
Form	Liquid
Additional notes	Protein concentration is above or equal to 0.05 mg/ml.

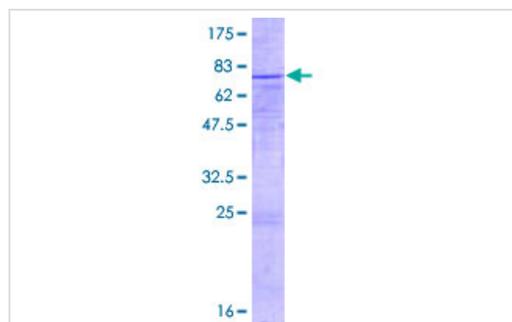
Preparation and Storage

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 HCl
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General Info

Function	Protective protein appears to be essential for both the activity of beta-galactosidase and neuraminidase, it associates with these enzymes and exerts a protective function necessary for their stability and activity. This protein is also a carboxypeptidase and can deamidate tachykinins.
Involvement in disease	Defects in CTSA are the cause of galactosialidosis (GSL) [MIM:256540]. A lysosomal storage disease associated with a combined deficiency of beta-galactosidase and neuraminidase, secondary to a defect in cathepsin A. All patients have clinical manifestations typical of a lysosomal disorder, such as coarse facies, cherry red spots, vertebral changes, foam cells in the bone marrow, and vacuolated lymphocytes. Three phenotypic subtypes are recognized. The early infantile form is associated with fetal hydrops, edema, ascites, visceromegaly, skeletal dysplasia, and early death. The late infantile type is characterized by hepatosplenomegaly, growth retardation, cardiac involvement, and a normal or mildly affected mental state. The juvenile/adult form is characterized by myoclonus, ataxia, angiokeratoma, mental retardation, neurologic deterioration, absence of visceromegaly, and long survival.
Sequence similarities	Belongs to the peptidase S10 family.
Cellular localization	Lysosome.

Images



ab159166 on a 12.5% SDS-PAGE stained with Coomassie Blue.

SDS-PAGE - Recombinant Human Protective protein/Cathepsin A (PPCA) (ab159166)

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