abcam

Product datasheet

Anti-ADAMTS10 antibody [EPR11860] ab173290

Recombinant

RabMAb

2 Images

Overview

Product name Anti-ADAMTS10 antibody [EPR11860]

Description Rabbit monoclonal [EPR11860] to ADAMTS10

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF,IHC-P or IP

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Recombinant fragment within Human ADAMTS10 aa 900-1100. The exact sequence is

proprietary.

Database link: Q9H324

Positive control Fetal heart and A549 lysates.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

Improved sensitivity and specificityLong-term security of supply

- Animal-free production

For more information **see here**.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

Properties

Form Liquid

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

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

Purity Tissue culture supernatant

1

ClonalityMonoclonalClone numberEPR11860

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab173290 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/1000 - 1/5000. Predicted molecular weight: 121 kDa.

Application notes Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

Target

Function Metalloprotease that participate in microfibrils assembly. Microfibrils are extracellular matrix

components occurring independently or along with elastin in the formation of elastic tissues.

Tissue specificity Widely expressed in adult tissues.

Involvement in disease Defects in ADAMTS10 are the cause of Weill-Marchesani syndrome 1 (WMS1) [MIM:277600].

WMS1 is a rare connective tissue disorder characterized by short stature, brachydactyly, joint stiffness, and eye abnormalities including microspherophakia, ectopia lentis, severe myopia and

glaucoma.

Sequence similaritiesContains 1 disintegrin domain.

Contains 1 peptidase M12B domain.

Contains 1 PLAC domain.
Contains 5 TSP type-1 domains.

DomainThe spacer domain and the TSP type-1 domains are important for a tight interaction with the

extracellular matrix.

Post-translational

modifications

Glycosylated. Can be O-fucosylated by POFUT2 on a serine or a threonine residue found within the consensus sequence C1-X(2)-(S/T)-C2-G of the TSP type-1 repeat domains where C1 and

C2 are the first and second cysteine residue of the repeat, respectively. Fucosylated repeats can

then be further glycosylated by the addition of a beta-1,3-glucose residue by the

glucosyltransferase, B3GALTL. Fucosylation mediates the efficient secretion of ADAMTS family

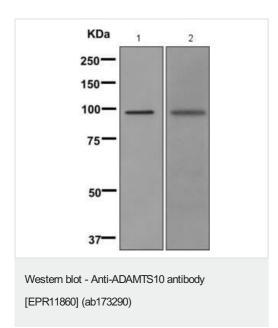
members. Also can be C-glycosylated with one or two mannose molecules on tryptophan

residues within the consensus sequence W-X-X-W of the TPRs, and N-glycosylated. These other

glycosylations can also facilitate secretion.

Cellular localization Secreted > extracellular space > extracellular matrix.

Images



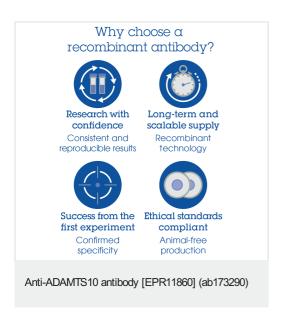
All lanes : Anti-ADAMTS10 antibody [EPR11860] (ab173290) at 1/1000 dilution

Lane 1 : Fetal heart lysate

Lane 2 : A549 cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 121 kDa



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

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