abcam

Product datasheet

Anti-ADAMTS13 antibody ab28274

★★★★★ 1 Abreviews 4 References 1 Image

Overview

Product name Anti-ADAMTS13 antibody

Description Rabbit polyclonal to ADAMTS13

Host species Rabbit

Specificity ab28274 recognises the metalloproteinase domain of ADAMTS13.

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide corresponding to Human ADAMTS13.

(Peptide available as ab41249)

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

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: 7.40

Constituent: PBS

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

1

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab28274 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. Detects a band of approximately 190 kDa (predicted molecular weight: 154 kDa). 1/1000 when using colorimetric substrates such as BCIP/NBT - 1/5000, when using chemiluminescent substrates. Glycosylation and the abundance of cysteine residues gives ADAMTS 13 an apparent molecular weight of 190 kDa on reduced SDS PAGE gels. Several bands at 110-190 kDa are observed on Western blots, possibly

Function		

Cleaves the vWF multimers in plasma into smaller forms.

Tissue specificity

Target

Plasma. Expressed primarily in liver.

Involvement in disease

Defects in ADAMTS13 are the cause of thrombotic thrombocytopenic purpura congenital (TTP) [MIM:274150]; also known as Upshaw-Schulman syndrome (USS). A hematologic disease characterized by hemolytic anemia with fragmentation of erythrocytes, thrombocytopenia, diffuse and non-focal neurologic findings, decreased renal function and fever.

Sequence similarities

Contains 2 CUB domains.
Contains 1 disintegrin domain.
Contains 1 peptidase M12B domain.
Contains 8 TSP type-1 domains.

Domain

The pro-domain is not required for folding or secretion and does not perform the common function of maintening enzyme latency.

The spacer domain is necessary to recognize and cleave vWF. The C-terminal TSP type-1 and

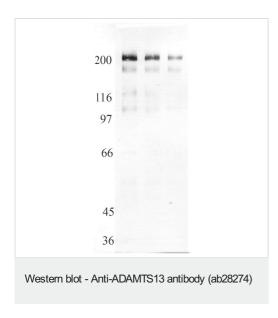
CUB domains may modulate this interaction.

Post-translational modifications

May contain a C-mannosylation site and O-fucosylation sites in the TSP type-1 domains. The precursor is processed by a furin endopeptidase which cleaves off the pro-domain.

Cellular localization Secreted.

Images



All lanes: Anti-ADAMTS13 antibody (ab28274) at 1 µg/ml

Lane 1: Recombinant Human ATS-13 at 0.08 μg

Lane 2 : Recombinant Human ATS-13 at 0.04 μg

Lane 3: Recombinant Human ATS-13 at 0.02 μg

Predicted band size: 154 kDa

Glycosylation and the abundance of cysteine residues gives ADAMTS-13 an apparent molecular weight of about 190 kDa on reduced SDS PAGE gels.

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

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- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

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