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Datasheet for ABIN881389 anti-ADAMTSL2 antibody (AA 522-580) (Alexa Fluor 555)



Overview

| Quantity: | 100 µL | |
|----------------------|--------------------------------------------------------------------------------------------------------------------------------|--|
| Target: | ADAMTSL2 | |
| Binding Specificity: | AA 522-580 | |
| Reactivity: | Human | |
| Host: | Rabbit | |
| Clonality: | Polyclonal | |
| Conjugate: | This ADAMTSL2 antibody is conjugated to Alexa Fluor 555 | |
| Application: | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Western Blotting (WB) | |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human ADAMTSL2 |
|-----------------------|--------------------------------------------------------------|
| lsotype: | lgG |
| Cross-Reactivity: | Human |
| Predicted Reactivity: | Mouse,Rat |
| Purification: | Purified by Protein A. |
| Target Details | |
| Target: | ADAMTSL2 |
| Alternative Name: | ADAMTSL2 (ADAMTSL2 Products) |

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| Target Details | | |
|---------------------|-----------------------------------------------------------------------------------------------|--|
| Background: | Synonyms: ADAMTS like 2, ADAMTS like protein 2, ADAMTS-like protein 2, ADAMTSL 2, | |
| | ADAMTSL-2, ADAMTSL2, ATL2_HUMAN. | |
| | Background: ADAMTS (A Disintegrin And Metalloproteinase Domain with Thrombospondin type | |
| | 1 Modules) is a family of zinc-dependent proteases that are implicated in a variety of normal | |
| | and pathological conditions, including arthritis and cancer. ADAMTS protein family members | |
| | contain an amino-terminal propeptide domain, a metalloproteinase domain, a disintegrin-like | |
| | domain and a carboxy-terminus that contains a varying number of Thrombospondin type 1 | |
| | (TSP-1) motifs. ADAMTS-L2 (ADAMTS-like protein 2) is a 951 amino acid secreted protein that | |
| | is highly expressed in lung, kidney and liver. Mutations in the gene encoding ADAMTS are the | |
| | cause of geleophysic dysplasia, an autosomal recessive disorder characterized by cardiac | |
| | vavular anomalies, short stature, thick skin and brachydactyly. In individuals affected with | |
| | geleophysic dysplasia, there is a significant increase in total active TGF-beta 1 and nuclear | |
| | locations of p-SAMD2 in fibroblasts. Interestingly, ADAMTS-L2 interacts with LTBP-1, a | |
| | glycoprotein that is part of the platelet-derived TGF-beta 1 complex. | |
| Gene ID: | 9719 | |
| Application Details | | |
| Application Notes: | IF(IHC-P) 1:50-200 | |
| | IF(IHC-F) 1:50-200 | |
| | IF(ICC) 1:50-200 | |
| Restrictions: | For Research Use only | |
| Handling | | |
| Format: | Liquid | |
| Concentration: | 1 µg/µL | |
| Buffer: | Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and | |
| | 50 % Glycerol. | |
| Preservative: | ProClin | |
| Precaution of Use: | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be | |
| | handled by trained staff only. | |
| Storage: | -20 °C | |
| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. | |
| | | |

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Expiry Date:

12 months

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