Anti-Antithrombin III / ATIII / SERPINC1 Antibody [15F1]

EM1901-11



Product Type: Mouse monoclonal IgG1, primary antibodies

Species reactivity: Human, Mouse, Rat, Zebrafish

Applications: WB, IHC-P, FC

Molecular Wt: Predicted band size: 53 kDa

Clone number: 15F1

Description: The protein encoded by this gene, antithrombin III, is a plasma protease inhibitor and a

member of the serpin superfamily. This protein inhibits thrombin as well as other activated serine proteases of the coagulation system, and it regulates the blood coagulation cascade. The protein includes two functional domains: the heparin binding-domain at the N-terminus of the mature protein, and the reactive site domain at the C-terminus. The inhibitory activity is enhanced by the presence of heparin. Numerous mutations have been identified for this gene, many of which are known to cause antithrombin-III deficiency which constitutes a strong risk factor for thrombosis. A reduction in the serum level of this protein is associated

with severe cases of Coronavirus Disease 19 (COVID-19).

Immunogen: Recombinant protein within Human SERPINC1 aa 150-370 / 464.

Positive control: Mouse liver tissue lysate, HL-60 cell lysates, HepG2 cell lysate, SiHa cell lysate, U937 cell

lysate, zebrafish tissue lysates, rat testis tissue, human lung carcinoma tissue, human liver

tissue, mouse kidney tissue, HCT116.

Subcellular location: Extracellular space or secreted.

Database links: SwissProt: P01008 Human | P32261 Mouse | Q5M7T5 Rat

Recommended Dilutions:

WB 1:500-1:2,000 ICC/ IHC-P 1:50-1:200 FC 1:50-1:100

Storage Buffer: 1*PBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide.

Storage Instruction: Shipped at 4℃. Store at +4℃ short term (1-2 weeks). It is recommended to aliquot into

single-use upon delivery. Store at -20 ℃ long term.

Purity: Protein A affinity purified.

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Images

kDa M. M. Berninghi 250-150-100-75-55-45-35-25-14-HSP90 **Fig1:** Western blot analysis of Antithrombin III / ATIII / SERPINC1 on different lysates with Mouse anti-Antithrombin III / ATIII / SERPINC1 antibody (EM1901-11) at 1/1,000 dilution.

Lane 1: Mouse liver tissue lysate

Lane 2: Mouse brain tissue lysate (low expression)

Lysates/proteins at 20 µg/Lane.

Predicted band size: 53 kDa Observed band size: 53 kDa

Exposure time: 23 seconds; ECL: K1801;

4-20% SDS-PAGE gel.

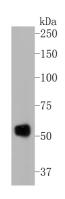


Fig2: Western blot analysis of Antithrombin III / ATIII / SERPINC1 on HL-60 cell lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (EM1901-11, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Mouse IgG - HRP Secondary Antibody (HA1006) at 1:5,000 dilution was used for 1 hour at room temperature.

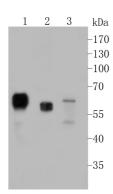


Fig3: Western blot analysis of Antithrombin III / ATIII / SERPINC1 on different lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (EM1901-11, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Mouse IgG - HRP Secondary Antibody (HA1006) at 1:5,000 dilution was used for 1 hour at room temperature.

Positive control:

Lane 1: HepG2 cell lysate Lane 2: SiHa cell lysate Lane 3: U937 cell lysate

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Orders:0086-571-88062880

Technical:0086-571-89986345

Service mail:support@huabio.cn

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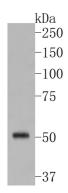


Fig4: Western blot analysis of Antithrombin III / ATIII / SERPINC1 on zebrafish tissue lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (EM1901-11, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Mouse IgG - HRP Secondary Antibody (HA1006) at 1:5,000 dilution was used for 1 hour at room temperature.

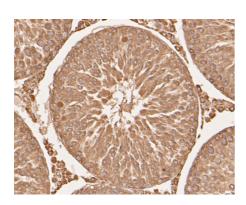


Fig5: Immunohistochemical analysis of paraffin-embedded rat testis tissue using anti-Antithrombin III / ATIII / SERPINC1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (EM1901-11, 1/50) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

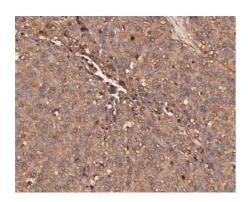


Fig6: Immunohistochemical analysis of paraffin-embedded human lung carcinoma tissue using anti-Antithrombin III / ATIII / SERPINC1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (EM1901-11, 1/50) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

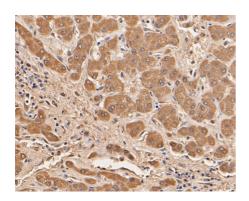
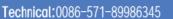


Fig7: Immunohistochemical analysis of paraffin-embedded human liver tissue using anti-Antithrombin III / ATIII / SERPINC1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (EM1901-11, 1/200) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

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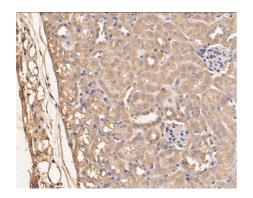


Fig8: Immunohistochemical analysis of paraffin-embedded mouse kidney tissue using anti-Antithrombin III / ATIII / SERPINC1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (EM1901-11, 1/50) for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

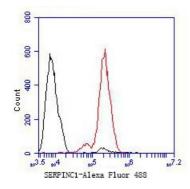


Fig9: Flow cytometric analysis of Antithrombin III / ATIII / SERPINC1 was done on HCT116 cells. The cells were fixed, permeabilized and stained with the primary antibody (EM1901-11, 1/50) (red). After incubation of the primary antibody at room temperature for an hour, the cells were stained with a Alexa Fluor 488-conjugated Goat anti-Mouse IgG Secondary antibody at 1/1000 dilution for 30 minutes.Unlabelled sample was used as a control (cells without incubation with primary antibody; black).

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- 1. Sungurlu S et al. Role of Antithrombin III and Tissue Factor Pathway in the Pathogenesis of Sepsis. Crit Care Clin. 2020 Apr
- 2. Farrell DH et al. Antithrombin III Levels and Outcomes Among Patients With Trauma. JAMA Netw Open. 2024 Aug

