

# AAT (13702): sc-73431

## BACKGROUND

Cumulative damage to lung tissue by neutrophil elastase is responsible for the development of pulmonary emphysema, an irreversible lung disease characterized by loss of lung elasticity.  $\alpha$ 1-antitrypsin (AAT), a 394 amino acid hepatic acute phase protein, predominantly inhibits neutrophil elastase. AAT is highly expressed in liver and in cultured hepatoma cells and, to a lesser extent, in macrophages. AAT is a highly polymorphic glycosylated serum protein with characteristic isoelectric-focusing patterns for most variants. The gene encoding AAT maps to a region of human chromosome 14 that includes a related serine protease inhibitor (Serpin) gene which encodes corticosteroid-binding globulin. Oxidation of the methionine 358 residue in the active center of AAT results in a dramatic decrease in inhibitory activity towards elastase. AAT also has a moderate affinity for plasmin and Thrombin. AAT deficiency is associated with a 20-30 fold increased risk of precocious pulmonary emphysema.

## REFERENCES

1. Okayama, H., et al. 1991. Characterization of the molecular basis of the  $\alpha$ 1-antitrypsin F allele. *Am. J. Hum. Genet.* 48: 1154-1158.
2. Seyama, K., et al. 1991. Siiyama (Serine 53 (TCC) to phenylalanine 53 (TTC)). A new  $\alpha$ 1-antitrypsin-deficient variant with mutation on a predicted conserved residue of the Serpin backbone. *J. Biol. Chem.* 266: 12627-12632.
3. Rosenberg, S., et al. 1994. Synthesis in yeast of a functional oxidation-resistant mutant of human  $\alpha$ -antitrypsin. *Nature* 312: 77-80.
4. Graziadei, I., et al. 2000. A novel-binding site for the native hepatic acute-phase protein  $\alpha$ -antitrypsin expressed on the human hepatoma cell line Hep G2 and intestinal cell line Caco 2. *Liver* 20: 240-246.
5. Rollini, P. and Fournier, R.E. 2000. Differential regulation of gene activity and chromatin structure within the human Serpin gene cluster at 14q32.1 in macrophage microcell hybrids. *Nucleic Acids Res.* 28: 1767-1777.
6. Hsu, P.I., et al. 2007.  $\alpha$ 1-antitrypsin precursor in gastric juice is a novel biomarker for gastric cancer and ulcer. *Clin. Cancer Res.* 13: 876-883.
7. Churg, A., et al. 2007.  $\alpha$ 1-antitrypsin suppresses TNF $\alpha$  and MMP-12 production by cigarette smoke-stimulated macrophages. *Am. J. Respir. Cell Mol. Biol.* 37: 144-151.
8. Zhang, B., et al. 2007.  $\alpha$ 1-antitrypsin protects  $\beta$  cells from apoptosis. *Diabetes* 56: 1316-1323.

## CHROMOSOMAL LOCATION

Genetic locus: SERPINA1 (human) mapping to 14q32.13.

## SOURCE

AAT (13702) is a mouse monoclonal antibody raised against  $\alpha$ 1-antitrypsin of human origin.

## PRODUCT

Each vial contains 100  $\mu$ g IgG<sub>1</sub> in 1.0 ml PBS with < 0.1% sodium azide and 0.1% gelatin.

## APPLICATIONS

AAT (13702) is recommended for detection of AAT of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)]; non cross-reactive with  $\alpha$ 1-antichymotrypsin (AACT).

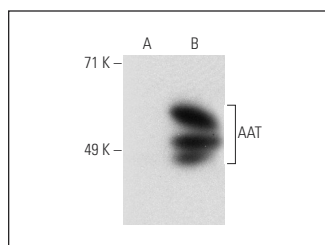
Suitable for use as control antibody for AAT siRNA (h): sc-40945, AAT shRNA Plasmid (h): sc-40945-SH and AAT shRNA (h) Lentiviral Particles: sc-40945-V.

Molecular Weight of luminal AAT: 51 kDa.

Molecular Weight of mature AAT: 55 kDa.

Positive Controls: human plasma extract: sc-364374, AAT (h): 293 Lysate: sc-112989 or Hep G2 cell lysate: sc-2227.

## DATA



AAT (13702): sc-73431. Western blot analysis of AAT expression in non-transfected: sc-110760 (A) and human AAT transfected: sc-112989 (B) 293 whole cell lysates.

## SELECT PRODUCT CITATIONS

1. Novotna, A., et al. 2011. Construction and characterization of hepatocyte nuclear factor HNF4 $\alpha$ 1 over-expressing cell line derived from human hepatoma Hep G2 cells. *Eur. J. Pharmacol.* 669: 45-50.
2. Novotna, A., et al. 2013. Construction and characterization of peroxisome proliferator-activated receptor- $\gamma$  co-activator 1 alpha (PGC-1 $\alpha$  over-expressing cell line derived from human hepatocyte carcinoma Hep G2 cells). *Biomed. Pap. Med. Fac. Univ. Palacky Olomouc Czech. Repub.* 157: 214-221.
3. Yu, Y.B., et al. 2018. Differentiation of umbilical cord mesenchymal stem cells into hepatocytes in comparison with bone marrow mesenchymal stem cells. *Mol. Med. Rep.* 18: 2009-2016.

## STORAGE

Store at 4° C, **\*\*DO NOT FREEZE\*\***. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.