# abcam

### Product datasheet

## Native Human Antithrombin III/ATIII protein ab95077

**Description** 

Product name

Native Human Antithrombin III/ATIII protein

Purity > 95 % SDS-PAGE.

Expression system Native

Protein length Full length protein

Animal free No
Nature Native
Species Human

#### **Specifications**

Our Abpromise guarantee covers the use of ab95077 in the following tested applications.

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

**Applications** Purification

SDS-PAGE

Form Liquid

Additional notes 2ml of 1:1 slurry

1.0 mg/ml coupled

May be used repeatedly. Protocol for purifying high affinity heparin with immobilized antithrombin:

1. Equilibrate immobilized antithrombin in TBS (0.1M Tris-HCl, 0.15M NaCl, pH 7.4) or PBS

(0.05M Sodium Phosphate, 0.15M NaCl, pH 7.4).

2. Apply heparin in TBS or PBS. Binding capacity is ~0.1 mg high affinity heparin / ml resin and

must be determined by the end user.

3. Wash and elute heparin with 3M NaCl in TBS or PBS.

4. Re-equilabrate resin in TBS or PBS.

5. Add 0.02percent Sodium Azide for storage.

Previously labelled as Antithrombin III.

#### **Preparation and Storage**

**Stability and Storage** Shipped at 4°C. Store at +4°C.

pH: 7.40

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Preservative: 0.02% Sodium azide

Constituents: 1.58% Tris HCI, 0.87% Sodium chloride

#### **General Info**

**Function** Most important serine protease inhibitor in plasma that regulates the blood coagulation cascade.

AT-III inhibits thrombin as well as factors IXa, Xa and Xla. Its inhibitory activity is greatly enhanced

in the presence of heparin.

**Tissue specificity** Found in plasma.

Involvement in disease Defects in SERPINC1 are the cause of antithrombin III deficiency (AT3D) [MIM:613118]. AT3D is

an important risk factor for hereditary thrombophilia, a hemostatic disorder characterized by a tendency to recurrent thrombosis. AT3D is classified into 4 types. Type I: characterized by a 50% decrease in antigenic and functional levels. Type II: has defects affecting the thrombin-binding domain. Type III: alteration of the heparin-binding domain. Plasma AT-III antigen levels are normal

in type II and III. Type IV: consists of miscellaneous group of unclassifiable mutations.

**Sequence similarities** Belongs to the serpin family.

Post-translational

modifications

Phosphorylation sites are present in the extracelllular medium.

**Cellular localization** Secreted > extracellular space.

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