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Fortsættedes...
DESCRIPTION

FIELD OF THE INVENTION

[0001] The invention relates to an alpha-1-antitrypsin (AAT, also known as alpha-1 proteinase inhibitor, API, and A 1-PI) composition suitable for pharmaceutical use.

BACKGROUND OF THE INVENTION

[0002] Alpha-1-antitrypsin (AAT) is a glycopeptide inhibitor of proteases, and is found in human serum and other fluids. Protease inhibition by AAT is an essential component of the regulation of tissue proteolysis, and AAT deficiency is implicated in the pathology of several diseases. Individuals who inherit an alpha-1 antitrypsin deficiency, for example, have increased risk of suffering from severe early-onset emphysema, the result of unregulated destruction of lung tissue by human leukocyte elastase. The administration of exogenous human AAT has been shown to inhibit elastase and is associated with improved survival and reduction in the rate of decline of lung function in AAT-deficient patients (Crystal et al., Am. J. Respir. Crit. Care Med. 158:49-59 (1998); see R. Mahadeva and D. Lomas, Thorax 53:501-505 (1998) for a review.)

[0003] Because of its therapeutic utility, commercial AAT production has been the subject of considerable research. Much progress has been made in the production of recombinant AAT in E. coli (R. Bischoff et al., Biochemistry 30:3464-3472 (1991)), yeast (K. Kwon et al., J. Biotechnology 42:191-195 (1995); Bollen et al., U.S. patent 4,629,567), and plants (J. Huang et al., Biotechnol. Prog. 17:126-33 (2001)), and by secretion in the milk of transgenic mammals (G. Wright et al., Biotechnology, 9:830-834 (1991); A.L. Archibald, Proc. Natl. Acad. Sci. USA, 87:5178-5182 (1990)). However, isolation of AAT from human plasma is presently the most efficient practical method of obtaining AAT in quantity, and human plasma is the only FDA-approved source.

[0004] A number of processes for isolating and purifying AAT from human plasma fractions have been described, involving combinations of precipitation, adsorption, extraction, and chromatographic steps. In order to minimize the risk of pathogen transfer, pooled human plasma intended for production of human AAT for therapeutic use is screened for the hepatitis B surface antigen, and for antibodies to the human immunodeficiency virus. As an additional precaution against transmission of infectious agents, the purified product is ordinarily pasteurized by heating to 60 °C for 10 hours (Mitra et al., Am. J. Med. 84(sup. 6A):87-90 (1988)) and sterile filtered.

[0005] Most published processes for AAT isolation begin with one or more fractions of human plasma known as the Cohn fraction IV precipitates, e.g. Cohn fraction IV 1 or fraction IV 1-4, which are obtained from plasma as a paste after a series of ethanol precipitations and pH adjustments (E. J. Cohn et al., J. Amer. Chem. Soc., 68:459-475 (1946)).

[0006] U.S. patent 3,301,842 describes a method for isolation of AAT from Cohn fraction IV 1 wherein an acridine or quinoline derivative is added to the paste in a buffer at pH 6, the precipitate is discarded, and the pH adjusted to 7.0. Additional acridine or quinoline is added, and the precipitate is collected. This precipitate is dissolved in a pH 5.0 buffer, sodium chloride is added, and the resulting precipitate discarded. The solution, containing the AAT, is further processed by methanol precipitation. Alternatively, ammonium sulfate precipitations at pH 8 and at pH 5 are conducted with plasma, with the pH 5 supernatant being further processed as above with quinoline or acridine additives.

[0007] Glaser et al., Preparative Biochemistry, 5:333-348 (1975), disclosed a method for isolating AAT from Cohn fraction IV 1 paste. The paste is stirred in a phosphate buffer at pH 8.5 in order to reactivate the AAT, which is largely deactivated by the pH of 5.2 employed in the Cohn fractionation. After dialysis and centrifugation, the supernatant is subjected to two rounds of anion exchange chromatography at pH 6.0 to 7.6 and at pH 8.6, followed by further chromatographic processing at pH 7.6 and at pH 8.0, to produce AAT in about a 30% overall yield.

[0008] M. H. Coan et al., in U.S. patents 4,379,087 and 4,439,356 (see also M. H. Coan et al., Vox Sang., 48:333-342 (1985)); M. H. Coan, Amer. J. Med., 84(sup 6A):32-36 (1988); and R. H. Hein et al., Eur. Respir. J., 3(sup 9):16s-20s (1990)), disclosed a procedure wherein Cohn fraction IV 1 paste is dissolved in a pH 6.5 to 8.5 buffer, polyethylene glycol is added, and the pH is lowered to the range of 4.6 to 5.7 to precipitate unwanted proteins. After centrifugation, AAT is isolated from the supernatant by anion exchange chromatography. Further processing provides a 45% yield of AAT with a purity of about 60%. Methods employing polyethylene glycol as a precipitant are also described in U.S. 4,697,003, US 4,856,254, and Japanese patent JP 08099999,
described below; and also by Hao et al., Proc. Intl. Workshop on Technology for Protein Separation and Improvement of Blood Plasma Fractionation, Sept. 7-8, 1977, Reston, VA.

[0009] Dubin et al., Preparative Biochemistry. 20:63-70 (1990), disclosed a two step chromatographic purification, in which AAT was first eluted from Blue SEPHAROSE™ and then purified by gel filtration chromatography.


[0011] Lebing and Chen, in U.S. Patent 5,610,285, disclosed a purification process which employs an initial anion exchange chromatography, followed by cation exchange chromatography at low pH and low ionic strength, to purify human AAT from plasma and plasma fractions. The cation chromatography takes advantage of the fact that active AAT does not bind to the ion exchange column under these conditions while contaminating proteins, including denatured AAT and albumin, are retained.


[0013] Shearer et al., in European patent application EP 0 224 811 and in the corresponding U.S. patent 4,656,254, disclosed an improved method for extracting AAT from Cohn fraction IV paste, in which the improvement consisted of treating the paste with a larger volume of buffer, at a higher pH, than had been customary in the prior art. The combination of higher volume and higher pH increased the amount of AAT extracted from the paste. Unwanted proteins were precipitated by addition of polyethylene glycol, followed by centrifugation. An alternative procedure is disclosed, which is essentially the procedure of Coan et al., wherein after addition of polyethylene glycol, the pH is adjusted to the range of 4.6 to 5.7, and the acidified mixture held for from one to sixty minutes to further precipitate unwanted proteins. The AAT is precipitated by addition of additional polyethylene glycol, and further purified by anion exchange chromatography.

[0014] Arrighi et al., in European application EP 0717049, disclosed a process wherein fraction IV paste is stirred in a pH 8.2 buffer at 40 °C for one hour, followed by precipitation of unwanted proteins with ammonium sulfate. The AAT is isolated from the supernatant by hydrophobic interaction chromatography at pH 7.

[0015] Kress et al., in Preparative Biochemistry 3:541-552 (1973), dialyzed the precipitate from an 80% ammonium sulfate treatment of human plasma, then chromatographed it on DEAE-cellulose. The product was dialyzed again and gel filtered on SEPHADEX™ G-100. AAT-containing fractions were then chromatographed on DE-52 cellulose to give AAT.

[0016] Japanese patent 59-128335 discloses the precipitation of unwanted proteins from a plasma fraction by addition of polyethylene glycol at a pH between 5 and 7, followed by anion exchange chromatography.

[0017] Bollen et al., in U.S. patent 4,629,567, disclose the isolation of AAT from a culture of yeast carrying recombinant plasmids expressing AAT. The process begins with polyethylene glycol precipitation at pH 8.5 to remove contaminating proteins, followed by anion exchange chromatography at pH 6.5 and subsequent chromatographic steps.

[0018] Dove and Mitra, in U.S. patent 4,684,723, disclose a variant of the method of Coan et al. (US 4,379,087 and US 4,439,385) in which AAT is purified by a process comprising the steps of (a) holding a solution containing AAT at a pH of 6.5 to 8.5 for up to 24 hours, (b) adding polyethylene glycol and an inorganic salt, so as to obtain a two-phase mixture, and (c) isolating the aqueous salt phase, which contains purified AAT.

[0019] Taniguchi et al., in PCT application WO 95/35306, disclose a similar process, involving precipitation with polyethylene glycol in the presence of zinc chloride, followed by anion-exchange chromatography and chromatography on a metal chelate resin.

[0020] Van Wietendaele et al., in U.S. patent 4,857,317, also disclose a process for isolating AAT from the crude extract of an engineered yeast culture, which comprises addition of polyethylene glycol at pH 6.1, centrifugation to remove precipitated proteins, addition of calcium chloride, storage for 24 hours at pH 7.0, and centrifugation to further remove contaminants. AAT is then isolated from the supernatant by subsequent chromatographic steps.

[0021] Coan, in U.S. patent 4,697,003, discloses a method for isolating AAT from various Cohn plasma fractions which comprises the removal of ethanol and salts from an AAT-containing fraction, followed by anion-exchange chromatography on DEAE cellulose
or a similar material under conditions such that the AAT is retained on the column while undesired proteins are eluted. Coan also describes "pasteurization" at about 60 °C or more for about 10 hours, which is stated to be sufficient to render hepatitis viruses non-infective.

[0022] Coan discloses addition of carbohydrate as a stabilizing agent, either alone or with sodium citrate, in order to stabilize the AAT at the pasteurization temperature. Suitable carbohydrates are said to be mono-, di-, and trisaccharides, and sugar alcohols such as sorbitol and mannitol. AAT is prone to both polymerization and to the adoption of inactive conformations upon heating; the presence of stabilizers reduces but does not eliminate thermal inactivation (D. Lomas et al., Eur. Resp. J. 10:672-675 (1997)). Size-exclusion HPLC analysis has shown that 10% of monomeric AAT is polymerized or aggregated when pasteurization is carried out according to the Coan process (M. H. Coan et al., Vox Sang., 48:333-342 (1985)).

[0023] T. Burnouf et al., Vox Sang., 52:291-297 (1987), disclosed substantially the same procedure for isolating AAT from Kistler-Nitschmann supernatant A. DEAE chromatography of Cohn Fractions II-III and size exclusion chromatography produced an AAT which was 80-90% pure (by SDS-PAGE) with a 36-fold increase in purity. Recovery was 65-70%.

[0024] Thierry, in European patent application EP 0282363, also discloses a method of obtaining AAT from a Kistler-Nitschmann plasma fraction. Briefly, plasma is precipitated with 10% ethanol at pH 7.4, and the supernatant precipitated again with 19% ethanol at pH 5.85. The supernatant from the second precipitation is applied to a DEAE anion-exchange column, and eluted at pH 5.2 to provide AAT of about 90% purity.

[0025] Strancar et al., in PCT patent application WO 95/24428, disclose a very similar method, employing a particular class of functionalized anion-exchange media. Desalted Cohn fraction IV1 is applied to the column, and the proteins are eluted with low salt buffer at a pH "close to the pKa of acetic acid." (The pKa of acetic acid is 4.74.) AAT is then eluted with 50 to 300 mM NaCl at pH 7.4 to 9.2.

[0026] Japanese patent JP 08099999 discloses a method of obtaining AAT from Cohn fraction IV or IV1, which involves reduction of salt concentration to below about 0.02 M, adjusting the pH to 4.5 to 5.5, and contacting the solution with a cation exchanger to adsorb contaminating proteins.


[0029] Cohn fraction IV1 precipitate which comprises the steps of (a) dissolving the paste in a pH 8.5 buffer, (b) filtering, (c) adding a dithiol such as DTT, and (d) precipitation of denatured proteins with ammonium sulfate. Glaser states that the destabilized (DTT-reduced) proteins may be precipitated by "suitable techniques such as salting, heating, change in pH, addition of solvents and the like."

[0030] Glaser et al. describe one variation in which treatment with DTT is carried out in the presence of 2.5% AEROSIL™ fumed silica, prior to precipitation with 50% saturated ammonium sulfate. Recovery of AAT was as good as it was in the absence of the silica, and the purification factor was improved by about 70%. In both references, the authors relegate the silica to a secondary role, that of an additive that improves the results of the ammonium sulfate precipitation. The effectiveness of silica alone, without ammonium sulfate precipitation, is not recognized or described. If the concentration of the protein solution appreciably exceeds about 50 mg protein/ml, AAT is reportedly lost by occlusion in the precipitate.

[0031] Ralston and Drohan, in US Patent 6,093,804, disclose a method involving the removal of lipoproteins from an initial protein suspension via a "lipid removal agent," followed by removal of "inactive AAT" via elution from an anion-exchange medium with a citrate buffer. The lipid removal agent is stated to be MICRO CEL™ E, a synthetic hydrous calcium silicate. In the presence of a non-citrate buffer, the anion-exchange medium binds active AAT while allowing "inactive AAT" to pass through. A citrate buffer is specified for subsequent elution of the AAT from the anion exchange medium, and also for later elution from a cation-exchange medium. Ralston and Drohan do not describe the use of a disulfide-reducing agent. The process is stated to provide AAT with a product purity of >90%; and manufacturing scale yields of >70%.

[0032] W. Stephan, in Vox Sanguinis 20:442-457 (1971), describes the use of fumed silica to adsorb lipoproteins from human blood serum solutions. The effect of silica adsorption on the concentrations of several plasma proteins, including AAT, was
evaluated, and there was no significant loss of AAT.

[0033] Mattes et al., in Vox Sanguinis 81:29-36 (2001), and in PCT application WO 98/16821 and published US patent application 2002/0082214, disclose a method for isolating AAT from Cohn fraction IV which involves ethanol precipitation, anion exchange chromatography, and adsorption chromatography on hydroxyapatite. The latter step is reported to remove inactive AAT, providing a product with very high specificity activity.

[0034] While AAT is an effective treatment for emphysema due to alpha-1-antitrypsin deficiency, treatment is very costly (currently about $25,000 per year), due to the limited supply and a complex manufacturing process. There remains a need for more efficient and cost-effective methods for isolating human AAT from plasma and other complex protein mixtures containing AAT. In particular, ammonium sulfate precipitation followed by dialysis is a time-consuming process, that generates substantial amounts of waste water, and there is a need for scalable processes that do not require extensive dialysis while providing high yields of high activity, high purity AAT. Thermal pasteurization of AAT effectively reduces viral contamination, but it leads to the formation of inactive AAT aggregates and polymers. Thus, there is also a need for highly pure AAT with reduced viral contamination but without significant amounts of inactive AAT aggregates and polymers. The present invention addresses these needs.

BRIEF DESCRIPTION OF THE INVENTION

[0035] The invention provides an alpha-1-antitrypsin (AAT) composition, comprising (a) 6% or less contaminating proteins based on total protein weight as determined by SDS-PAGE; and (b) less than 0.2% IgA; wherein the specific activity of the alpha-1-antitrypsin is at least 0.99 mg functional AAT per milligram of protein, when using as an extinction coefficient
\[ E_{\text{ext}, 280nm}^{\alpha} = 5.3 \]
and wherein less than 8% of the composition is of a higher molecular weight than monomeric AAT.

[0036] The present invention in particular embodiments provides a preparation of AAT characterized by the following properties:

1. (a) the alpha-1-antitrypsin contains less than 6%, preferably less than 2%, and most preferably less than 1% contaminating proteins by SDS-PAGE, and contains
2. (b) less than 0.1% Albumin;
3. (c) less than 0.8%, and preferably less than or equal to 0.2% α1-acid glycoprotein;
4. (d) less than 0.1% α2-macroglobulin;
5. (e) less than 0.1% apolipoprotein A1;
6. (f) less than 0.5%, and preferably less than or equal to 0.1% antithrombin III;
7. (g) less than 0.1 % ceruloplasmin;
8. (h) less than 0.5%, and preferably less than 0.1% haptoglobin;
9. (i) less than 0.2%, and preferably less than 0.1 % IgA;
10. (j) less than 0.1% IgG;
11. (k) less than 0.1%; transferrin;
12. (l) the specific activity of the alpha-1-antitrypsin is at least 0.99 mg functional AAT /mg, when using as an extinction coefficient
\[ E_{\text{ext}, 280nm}^{\alpha} = 5.3 \]
13. (m) less than 8%, and preferably less than 5%, of the product is of a higher molecular weight than monomeric AAT;
14. (n) the apparent ratio of active to antigenic AAT is greater than 1.08, preferably greater than 1.16, and most preferably greater than 1.23, when measured by endpoint nephelometry;
15. (o) enveloped viruses are reduced by at least 11 log_{10} units, and non-enveloped viruses by at least 6 log_{10} units, when measured in spiking studies using human and model viruses representing a wide range of physico-chemical properties; and
16. (p) the product is stable for at least 2 years when stored lyophilized at up to 25 °C.

[0037] The apparent ratio of active to antigenic AAT in the product of the present invention is greater than unity because the purity and/or activity of the product of the present invention is greater than that of the reference standard, which is a prior art composition. Antigenic levels, as determined by endpoint nephelometry, are measured against the current protein standard (product No. OQIM15, supplied by Dade-Behring, Deerfield, Illinois), which is calibrated directly against the internationally-recognized Certified Reference Material 470 (Reference Preparation for Proteins in Human Serum; see J.T. Whicher et al., Clin.
Chem. 40:934-938 (1994)), using reagents and AAT antibody (Dade-Behring product No. OSAZ15), as supplied for the Dade-Behring Nephelometer 100.

[0038] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. The term "AAT" refers to human AAT generally, whether heterogeneous or homogeneous, and whether isolated from human serum or from a recombinant organism. The term is intended to embrace pharmacologically effective naturally-occurring variants (see for example, Brantly et al., Am. J. Med. 84(sup.6A):13-31 (1988)), as well as pharmacologically effective non-natural forms of human AAT, including but not limited to those having non-human glycosylation patterns, N-terminal methionine, or altered amino acids.

BRIEF DESCRIPTION OF THE FIGURES

[0039] Figure 1 is a flow chart showing an overall AAT purification process.

Figure 2 is an SDS-PAGE gel showing the proteins present in the products produced by the process at various stages. Lane 1, molecular weight markers; Lane 2, Plasma (Cryo-Poor); Lane 3, Fraction IV1,4 Extract; Lane 4, DTT/Aerosil-Treated Extract Filtrate; Lane 5, IEC Eluate; Lane 6, HIC Effluent; Lane 7, final container.

DETAILED DESCRIPTION OF THE INVENTION

[0040] The method exemplified below employs a particular Cohn fraction IV paste as a starting material, but the use of similar plasma fractions is contemplated. Alternative starting materials include but are not limited to other AAT-containing Cohn fractions (see U.S. patent 4,687,003), a precipitate from Kistler-Nitschmann supernatants A or A+1 (P.Kistler, H.S. Nitschmann, Vox Sang., 7:414-424 (1962)), and ammonium sulfate precipitates from plasma as described by Schultze et al. in U.S. patent 3,301,842. The use of protein precipitates derived from cultures of AAT-producing recombinant cells or organisms, or precipitates derived from the milk or serum of transgenic mammals, is also contemplated.

[0041] There are many methods known in the art for selectively precipitating proteins from solution, such as by the addition of salts, alcohols, and polyethylene glycol, often in combination with cooling and various pH adjustments. It is anticipated that the method will be applicable to most AAT-containing protein precipitates containing recoverable AAT activity, regardless of how they are initially prepared. The term "crude AAT-containing protein precipitate" is used herein to refer to any AAT-containing protein precipitate prepared by one or more of these known methods, whether from serum, milk, cell culture, or other original source.

[0042] In a method described below, the crude AAT-containing protein precipitate is suspended in a Tris buffer, and treated with dithiothreitol (DTT, a preferred disulfide-reducing agent) and fumed silica (a preferred protein-adsorbing material) in order to remove contaminating proteins and lipids. Where the precipitate is Cohn fraction IV, the two major protein contaminants thus removed are albumin and transferrin. DTT and other dithiols, as well as phosphines, are known in the art to reduce intrachain and inter-chain disulfide bonds. Cleavage of structurally important disulfide bonds causes partial unfolding and destabilization of those contaminating proteins that have disulfide bonds. AAT itself is not destabilized by DTT treatment because it has no intrachain disulfide bonds.

[0043] Fumed silica is known to bind preferentially to hydrophobic proteins. It is theorized that in the method described below, the destabilized contaminating proteins bind to a protein-adsorbing material such as fumed silica because the partial unfolding caused by disulfide bond cleavage exposes the proteins' inner core of hydrophobic residues.

[0044] In the method, described below, the protein-adsorbing material, together with the adsorbed contaminating proteins, lipids, and other insoluble material, is removed from the suspension by filtration so as to obtain a clarified AAT-containing protein solution. Filtration is preferably carried out with the assistance of a filtering aid such as Celite™ diatomaceous earth, and preferably the suspension is recirculated through the filter until a clarity of <10 nephelometer turbidity units (NTU)/ml is achieved. The filtrate is further processed by chromatographic techniques to afford highly pure and highly active AAT. Other methods of separation known in the art, for example centrifugation, could also be employed in place of filtration. The practitioner will select the method appropriate to the scale of operations and the nature of the protein-adsorbing material.
After removal of insoluble materials, the AAT-containing solution may be further processed by any of the methods known in the art for protein purification, particularly the methods already known to be suitable for purification of AAT. In the method described below, the filtrate is first subjected to ion exchange chromatography ("IEC") with salt gradient elution. The chromatography column contains an anion exchange resin which consists of a porous resin support matrix to which positively charged groups are covalently attached. These positively charged groups reversibly bind anions, including proteins with anionic groups such as AAT.

AAT, and other proteins which have a net negative charge at the pH of the eluting buffer, bind to the IEC column. Contaminating proteins having little or no negative charge pass through the anion exchange resin column without binding and exit with the column effluent. Those contaminating proteins that do bind to the column are then separated from the AAT by gradient elution. The salt concentration is gradually increased as the column is eluted in order to release sequentially the various proteins that are bound to the resin.

In the method, described below, the AAT-containing eluate from the IEC column is subjected to hydrophobic interaction chromatography ("HIC"). This type of chromatography employs a support matrix to which moieties are covalently attached. In an aqueous environment, these hydrophobic moieties bind reversibly to hydrophobic molecules, such as the contaminating proteins remaining in the IEC eluate. AAT is relatively non-hydrophobic, therefore the majority of the AAT flows through the column during the elution of the column with buffer, while the more hydrophobic contaminating proteins remain bound to the column. The column effluent thus contains the purified AAT. In practice, AAT has been found to have a slight affinity for certain HIC column media, and in such cases further elution with several volumes of wash buffer may be desirable in order to recover substantially all of the AAT in the originally-applied sample.

After such additional purification steps as are required to reach the desired level of purity and activity, the AAT solution is then concentrated and sterilized. In the method described below, the AAT is at a pharmaceutically acceptable level of purity and activity after the hydrophobic interaction chromatography, and no additional steps are necessary. In the method described below, concentration is accomplished by ultrafiltration followed by dialysis filtration (dialfiltration). In these techniques, solvent and dissolved salts and small molecules are passed through a filtering membrane, leaving behind a more concentrated protein solution. Remaining salts and small molecules in the protein solution are then exchanged with a different buffer by continuous addition of several volumes of the new buffer to the product, while maintaining a constant product volume by continuously passing solution through the same membrane.

The AAT is then provided with a pharmaceutically acceptable buffer, and lyophilized by methods known in the art, preferably by methods known to be suitable for preparing AAT therapeutic formulations.

Proteins isolated from mammalian sources may contain pathogenic viral contaminants, and it is desirable to reduce or eliminate such contamination in pharmaceutical compositions. Methods of viral reduction are known to those of skill in the relevant arts. The methods include, but are not limited to, pasteurization, irradiation, solvent/detergent treatment, disinfection, filtration, and treatment with supercritical fluids. Solvent/detergent treatment can be carried out, for example, by contacting a protein solution with a polyoxyethylene sorbitan ester and tributyl phosphate (see US patent 4,820,805; see also WO 95/03506 for application of the method to an AAT composition.) Disinfection of a protein solution can be carried out by exposing the solution to a soluble pathogen inactivating agent, for example as disclosed in US patents 6,106,773, 6,369,048 and 6,436,344, or by contact with an insoluble pathogen inactivating matrix, for example as disclosed in US patent 6,098,218 and references therein. Filtration may be through 15-70 nm ultrafilters (e.g., VirAGard™ filters, A/G Technology Corp.; Planova™ filters, Asahi Kasei Corp.; Viresolve™ filters, Millipore Corp.; DV and Omega™ filters, Pall Corp.) Radiation may be with ultraviolet or gamma radiation; see for example US patent 6,187,572 and references therein. Inactivation of viruses by treatment with supercritical fluids is described in US patent 6465168. Pasteurization of a protein solution may be accomplished by heating within the limits dictated by the thermal stability of the protein to be treated. In the case of AAT, pasteurization is usually accomplished by heating to about 60-70 °C. In the method described below, viral reduction of the AAT concentrate is carried out by pasteurization and ultrafiltration.

Stabilizing additives may be added to protect the AAT from thermal degradation during the pasteurization step, as disclosed for example in US Patent 4,876,241. Sucrose and potassium acetate are preferably added as stabilizers, and the stabilized AAT solution is then pasteurized at about 60 °C to reduce viral contamination. The amount of sucrose is preferably at least 40%, more preferably at least 50%, and most preferably about 60% by weight. Use of less than 40% sucrose has been found to result in undesirable levels of aggregation of the AAT. The amount of potassium acetate is preferably at least 4%, more preferably at least 5%, and most preferably about 6% by weight.

After viral reduction, the AAT solution may optionally be diluted and ultrafiltered, then reconcentrated and sterilized, e.g.
by filtration. The sterilized AAT-containing concentrate may then be lyophilized to form a therapeutic product. A suitable composition for preparing a lyophilized AAT powder is shown in Table 1.

**Table 1**

<table>
<thead>
<tr>
<th>Component</th>
<th>Function</th>
<th>Concentration 1.0 g/vial</th>
</tr>
</thead>
<tbody>
<tr>
<td>AAT</td>
<td>Active Ingredient</td>
<td>50 mg/mLb</td>
</tr>
<tr>
<td>Sodium Phosphatec</td>
<td>Buffer, Tonicity</td>
<td>20 mM</td>
</tr>
<tr>
<td>Sodium Chloride USP</td>
<td>Tonicity</td>
<td>40 mM</td>
</tr>
<tr>
<td>Mannitol USP</td>
<td>Stabilizing Agent</td>
<td>3%</td>
</tr>
<tr>
<td>Sodium Hydroxide</td>
<td>To adjust pH</td>
<td>as needed</td>
</tr>
<tr>
<td>Hydrochloric Acid ACS</td>
<td>To adjust pH</td>
<td>as needed</td>
</tr>
<tr>
<td>Water for Injection USFd</td>
<td>Diluent/Vehicle</td>
<td>20 ml/vial</td>
</tr>
</tbody>
</table>

a The final product is ≥96% AAT as determined by SDS-PAGE and ≥93% monomer by HPLC.
b Functional AAT activity per ml.
c Added as Monobasic Sodium Phosphate Monohydrate or Dibasic Sodium Phosphate.
d Added as Sterile Water for Injection USP.

THE FINAL FORMULATION WILL DEPEND ON THE VIRAL INACTIVATION STEP(S) SELECTED AND THE INTENDED MODE OF ADMINISTRATION. Depending on whether the AAT is to be administered by injection, as an aerosol, or topically, the AAT may be stored as a lyophilized powder, a liquid, or a suspension. The composition shown in Table 1 is suitable for injection, and may be lyophilized and stored in glass vials for later reconstitution with sterile water. The composition of a suitable dry powder formulation for inhalation is shown in Table 2. Such a formulation is suitable for inhalation administration as described in US Patent 5,780,014, either with a metered dose inhaler, or with a pulmonary delivery device such as is disclosed in US Patent 6,138,668.

**Table 2**

<table>
<thead>
<tr>
<th>Component</th>
<th>Function</th>
<th>Nominal Content (per unit dose)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AAT</td>
<td>Active Ingredient</td>
<td>7,440 mg*</td>
</tr>
<tr>
<td>Sodium Citrate</td>
<td>Buffer</td>
<td>0.059 mg</td>
</tr>
<tr>
<td>Citric Acid</td>
<td>Buffer</td>
<td>0.001 mg</td>
</tr>
</tbody>
</table>

* corresponds to 6 mg functional AAT, and a delivered dose of approximately 3.6 mg functional AAT.

ASSAYS FOR DETERMINING THE QUANTITY AND QUALITY OF AAT ARE KNOWN IN THE ART AND MAY BE EMPLOYED FOR EVALUATING THE EFFICIENCY OF THE METHOD. An example of an immunocassay involving a monoclonal antibody specific for AAT, used for measuring or detecting AAT in biological fluids, is disclosed in U.S. Patent 5,114,863. An example of the use of rate nephelometry is disclosed in L. Gaidulis et al., Clin. Chem. 29:1838 (1983). AAT functional activity may be assayed by measuring its elastase inhibitory capacity using a chromogenic substrate for elastase, as described in U.S. Patent 4,697,003. AAT may also be assayed by measuring its trypsin inhibitory capacity in a similar manner. In the method described below, AAT is assayed by endpoint nephelometry, as described elsewhere in this specification.

THE QUANTITY OF PROTEINS MAY BE DETERMINED BY METHODS KNOWN IN THE ART, FOR EXAMPLE THE BRADFORD ASSAY, OR BY ABSORBANCE AT 280 nm USING AS AN EXTINCTION COEFFICIENT

\[ \varepsilon_{280}^{\text{mg/mL}} = 5.3 \]  
(R. Pannell, D. Johnson, and J. Travis, Biochemistry 13:5439-5445 (1974)). SDS-PAGE with staining and densitometry may be used to assess purity of the sample and detect the presence of contaminating proteins. A reducing agent such as dithiothreitol is preferably used with SDS-PAGE to cleave any disulfide-linked polymers, thereby facilitating the comparison of total AAT to total non-AAT protein. Size-exclusion HPLC may also be used to assess purity of the sample and detect the presence of both contaminating proteins and aggregate or polymeric forms of AAT. Analysis of four lots prepared by the method described below showed AAT protein purity by SDS-PAGE (reduced) of at least 98%, an AAT monomer content of at least 95%, and specific activity averaging 1.06 mg functional AAT/mg protein (Table 3).

**Table 3**
<table>
<thead>
<tr>
<th>Lot</th>
<th>%AAT Purity by SDS-PAGE (reduced)</th>
<th>%Monomeric AAT by HPLC</th>
<th>Specific Activity (mg functional AAT/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>98</td>
<td>95</td>
<td>1.10</td>
</tr>
<tr>
<td>B</td>
<td>99</td>
<td>95</td>
<td>1.09</td>
</tr>
<tr>
<td>C</td>
<td>98</td>
<td>95</td>
<td>1.05</td>
</tr>
<tr>
<td>D</td>
<td>98</td>
<td>96</td>
<td>1.04</td>
</tr>
</tbody>
</table>

[0055] Conditions for the method are as follows:

1. Preparation of Cohn Fraction IV1-4.

[0056] Human plasma is cooled to -2 to 2 °C and adjusted to a pH of 6.9 to 7.5. Cold ethanol is added to a concentration of 6 to 10%, and the temperature is lowered to -4 to 0 °C. The precipitate that forms ("Fraction Γ") is removed by centrifugation or filtration.

[0057] The filtrate or supernatant from the above procedure is adjusted to pH 6.7 to 7.1, and cold ethanol is added to a concentration of 18 to 22%. The temperature is lowered to -7 to -3 °C, and the mixture is again subjected to centrifugation or filtration. The precipitate that forms ("Fraction II + III") is set aside for other purposes.

[0058] The filtrate or supernatant from the above procedure is adjusted to pH 4.9 to 5.3 and the ethanol concentration is adjusted to 16 to 20 %. The temperature is adjusted to -7 to -3 °C. After the suspension settles, it is adjusted to pH 5.7 to 6.1 and the ethanol concentration is adjusted to 40 to 44%. The precipitate that forms ("Fraction IV1,4") is removed by centrifugation or filtration, and stored until needed in the form of a paste. Fraction IV1,4 contains AAT as well as contaminating proteins and lipids.

2. Purification with DTT and Silica.

[0059] The Fraction IV1,4 paste is suspended in a suspension buffer (e.g., 100 mM Tris, 20 mM NaCl, pH between about 7.5 and about 9.5, preferably between about 8 and about 9) and stirred for a minimum of one hour at low temperature. The amount of buffer used ranges from 6 to 10 kg of buffer per kg of the plasma-containing fraction.

[0060] The Tris buffer suspension is then treated with dithiothreitol (DTT) and fumed silica. DTT is added to the Tris buffer suspension at a concentration in the range of about 10-50 mM. The solution is stirred for at least 30 minutes, preferably 2-4 hours, at low temperature, and preferably at a pH of about 8.9. Fumed silica is added at a concentration of approximately 100-300 g fumed silica per kg Fraction IV precipitate. The suspension is stirred for at least 30 minutes, preferably 1-4 hours, at low temperature, at a pH of about 8-9. A filter aid such as Celite™ is added at the rate of five parts filter aid one part silica, by weight, and the mixture is stirred for approximately 15 minutes. The soluble AAT product is separated from the precipitated fumed silica and contaminating proteins using a filter press, yielding the AAT final filtrate. Preferably, the suspension is recirculated through the filter press until the desired level of clarity is obtained. The AAT final filtrate is then treated further as follows.

3. Ion Exchange Chromatography.

[0061] The AAT final filtrate is applied directly onto a chromatography column containing an anion exchange resin equilibrated with an IEC equilibration buffer. Contaminants are removed from the column by washing the column with an IEC wash buffer, and AAT is subsequently eluted using an IEC elution buffer.


[0062] The eluate from the IEC column is prepared for HC by adding ammonium sulfate to a final concentration of about 1 M.
The solution is then filtered and applied to a hydrophobic interaction chromatography column which is equilibrated in a HIC wash buffer. Initial elution with a wash buffer provides an AAT-containing effluent, and elution with additional wash buffer removes any AAT retained on the column. The combined effluent and washes are concentrated by ultrafiltration, and dialyzed into a phosphate buffer. The final AAT concentration is preferably no greater than 7 % protein.

5. Pasteurization

[0063] The AAT concentrate is stabilized for pasteurization by the addition of sucrose and potassium acetate, and pasteurized at about 60 °C for 10-11 hours. The pasteurized solution is held at 2-8 °C pending further processing.

6. Nanofiltration

[0064] The pasteurized AAT solution is diluted with a final formulation buffer. The diluted, pasteurized AAT solution is then filtered through two new YM-100 (Amicon) spiral-wound ultrafiltration cartridges. This nanofiltration step serves as a second primary viral reduction step. Viruses are retained by the membrane, which has a nominal 100,000 Dalton molecular weight cut-off, while AAT, which has an approximate molecular weight of 50 kD, passes through. The AAT is collected in the permeate of the second filter and in filter postwashes. The final filtrate is collected in a bulk receiver and held at 2-8 °C.

7. Sterile filtration and lyophilization

[0065] The AAT-containing final filtrate is concentrated and dialyzed into final formulation buffer at a temperature of no more than 15 °C to form a final bulk solution. This solution is clarified and sterilized by passage through a series of sterile, bacterial-retentive filters. The sterile bulk solution is filled into sterilized glass final containers. The filled containers are freeze-dried and then sealed under vacuum.

[0066] The product is ≥96% pure AAT as determined by both SDS-PAGE and immunological assays such as ELISA or nephelometry, and is ≥93% monomer by size exclusion HPLC. The recovery based on the functionally active AAT content of the Cohn fraction IV paste is 70%.

EXAMPLES

[0067] Fraction IV1-4 Precipitate (667 kg) was isolated via the Cohn plasma fractionation process from 9026 liters of human plasma. The material was divided into nine batches of approximately 75 kg each. Each batch was suspended in Tris Buffer, using 6 to 10 parts buffer (w/w) relative to the presscake. The suspensions were stirred for at least 15 minutes, the temperature was adjusted to 2° - 8°C, and the pH of each suspension was adjusted to 8.80-8.95 with 1 N sodium hydroxide or 1 N hydrochloric acid as necessary. The suspensions were stirred for 15 to 105 minutes (average 45 min), and monitored for protein (Bradford assay) and potency. Specific activity of each batch ranged from 0.027 to 0.045, and averaged 0.037 mg functional AAT per mg protein. Approximately 12% of the total protein was albumin, and approximately 22% was transferrin.

[0068] Dithiothreitol (DTT) was added to a final concentration of 0.01 to 0.05 M DTT (average 0.03 M) based upon the amount of Tris Buffer in each batch. After a pre-mix period of at least 15 minutes, the temperature was adjusted to 2°-8 °C and the pH re-adjusted to 8.80-8.95, and the solutions were stirred for 2 to 8 hours (average 3 hours). If necessary, the pH was again adjusted to 8.80-8.95.

[0069] Aerosil™ 380 (Degussa AG, Frankfurt-Main) was added at the rate of 13.4 to 18.6 g per liter plasma input (average 16.7 g). The suspensions were stirred for 1 to 4 hours (average 1 hour) at 2-8 °C.

[0070] Celite™ 545 was added to each suspension at the rate of 5 parts Celite to 1 part Aerosil, and the suspensions were stirred at 2-8 °C. Each suspension was then recirculated through a plate and frame filter press, holding 25 x 25 inch Cuno™ A2605-10CP filter pads (cellulose pads with inorganic filter aids; nominal cutoff 1 micron). When the turbidity was ≤ 10 NTU by nephelometry (minimum of 15 min.), re-circulation was discontinued and the filtrate was collected. The filter press was post-
washed with TRIS extraction buffer at 2-8 °C. The postwashes were combined with the initial filtrate solutions, and total protein in solution was determined by the Bradford protein assay. The filtrates were held at 2-8 °C for no longer than 19 hours. Based on AAT activity, the filtrates contained a total of 1557 g of AAT, corresponding to a 59 % yield of the activity present in the original suspension of Fraction IV paste, and a purification factor of 1.5. (In view of the activity present after subsequent processing, these values appear to be low, possibly due to the presence of unidentified factors interfering with the AAT assay.) Specific activity for each of the nine batches ranged from 0.042 to 0.064, and averaged 0.058 mg functional AAT per mg protein. Albumin and transferrin were below detection limits (total protein contained less than 0.5% albumin and less than 2.5% transferrin.)

[0071] A 92-liter, 30 cm high ion exchange chromatography (IEC) column loaded with TMAE Fractogel™ (EM Industries, Hawthorne, NY) was equilibrated with IEC equilibration buffer (50 mM Tris, pH 8.3-9.3, 20-25°C). Following equilibration, conductivity of the effluent was verified to be ≤1.25 mS/cm. Each filtrate from the previous step was warmed to 20-25°C and filtered through a Cuno Zeta Plus™ 90SP cartridge (45115-12-90SP, nominal MW cutoff of 0.1 micron) before loading onto the column with control of flow rate (≤3.0 cm/min) and column pressure (≤20 psi). Total protein loaded onto the IEC column was limited to no more than 70% of the resin capacity. The column was then washed with five column volumes of IEC wash buffer (50 mM Tris, 25-70 mM NaCl gradient, pH 7.1-7.7) at 20-25 °C, with control of flow rate (≤3.0 cm/min) and column pressure (≤20 psi). The effluent was monitored by Bradford protein determination, assay of AAT activity, and UV absorbance at 280nm.

[0072] AAT was eluted with approximately three column volumes of IEC elution buffer (50 mM Tris, 75-120 mM NaCl gradient, pH 7.1-7.7) at 20-25 °C, with control of flow rate (≤3.0 cm/min) and column pressure (≤20 psi). The effluent was monitored by Bradford protein determination, assay of AAT activity, and UV absorbance at 280nm. The entire peak that eluted after application of the elution buffer was collected for further processing.

[0073] The above procedure was repeated nine times in order to process all nine batches of filtrate. Ammonium sulfate was added to the IEC eluates to a final concentration of 0.9 to 1.1 M. The resulting solutions were either used immediately, or stored at 15-25 °C for no more than seven days. Based on AAT activity, the IEC eluates contained a total of 2241 g of AAT, corresponding to an 84% yield of the activity present in the original suspension of Fraction IV paste, and a purification factor of 16.2. Specific activity for each of the nine batches ranged from 0.416 to 0.975, and averaged 0.592 mg functional AAT per mg protein.

[0074] A Cuno™ filter (Zeta Plus™ 90SP cartridge (45115-12-90SP, nominal MW cutoff of 0.1 micron) was prepared with a hot WFI flush followed by a cold WFI rinse (WFI = Water for Injection). Water was gently blown out of the filter with compressed air. Three IEC eluates, containing ammonium sulfate, were pooled and filtered through the prepared Cuno filter and subsequently combined to provide the "filtered IEC solution". The filter was post-washed with approximately 20 liters HIC wash buffer (50 mM Tris, 1 M ammonium sulfate, pH 7.1-7.7). The post-wash and the filtrate were combined and weighed. The process was repeated three times to process the nine batches of IEC eluate.

[0075] A hydrophobic interaction column (HIC) was packed with Phenyl Sepharose™ Fast Flow HS resin (Pharmacia, Piscataway, NJ) to a volume of 49 liters (32 cm bed height), and equilibrated with HIC wash buffer (50 mM Tris, 1 M ammonium sulfate, pH 7.1-7.7). This and all column loading and subsequent elutions were carried out with control of flow rate (≤ 4 cm/min), column pressure (≤ 20 psi), and solution temperatures (20-25°C).

[0076] Each of the three batches of filtered IEC solution was loaded onto an HIC column. Total protein load onto the column was limited to ≤ 39g protein per liter of resin. Optical density (OD280) of the effluent was monitored, and collection was initiated when the OD280 rose 0.04 units higher than the baseline value. The column was washed with HIC wash buffer to elute additional AAT from the column, while non-AAT contaminants remained bound to the column. Approximately ten column volumes of HIC wash buffer was applied to the column, and effluent was collected until the A280 dropped to < 0.05 units above baseline. The AAT effluent and column wash were combined and weighed. Samples were taken for Bradford protein determination, OD Protein determination, potency, and LAL (Limulus amebocyte lysate) testing. The HIC effluents were held at 15-25 °C for no more than 72 hours. Based on AAT activity, the three batches of HIC effluent contained a total of 2090 g of AAT, corresponding to a 79% yield of the activity present in the original suspension of Fraction IV paste, and a purification factor of 25.6. Specific activity for each of the three batches ranged from 0.908 to 0.986, and averaged 0.937 mg functional AAT per mg protein.

[0077] A tangential flow ultrafiltration (UF) unit containing a polyether sulfone membrane (surface area: 50 ft²) with a molecular weight cut off range of 5,000-30,000 was integrity tested to ensure a bubble point of less than 1250 ml/minute. Dialfiltration buffer (40 mM sodium phosphate, pH 7.2-7.6, 10 kg minimum) was recirculated through the unit for a minimum of five minutes. The recirculated buffer solution was sampled to verify proper pH (7.2-7.6) and LAL (< 0.25 EU/ml). A repeat of the prewash steps was
performed if pH and LAL requirements were not met. The UF unit was held for no more than 12 hours at 2-8 °C prior to HIC effluent application.

[0079] Following concentration, the HIC effluent concentrate was diafiltered, exchanging the Tris-buffered ammonium sulfate solution with a sodium phosphate buffer. Diafiltration buffer (40 mM sodium phosphate, pH 7.2-7.6) was applied at a volume ten times the weight of the HIC effluent concentrate. Inlet pressure was maintained at ≤40 psi, and outlet pressure was monitored during the concentration process. Concentration was performed until the weight of the concentrate was approximately 10 kg.

[0080] Following diafiltration, the ultrafiltration was continued until the concentrate had a mass of approximately 6 kg. Product was then gently blown out of the UF system (≤ 25 psi). The ultrafiltration unit was postwashed twice with 1.5 kg diafiltration buffer. The UF postwashes were added to the diafiltered concentrate. The total weight of concentrate was determined and the protein concentration determined (OD at 280 nm).

[0081] Based on the OD protein observed, the AAT protein concentration was determined, and adjusted if necessary to the range 2.9-6.8%. Analysis for LAL, SDS-PAGE, Bradford protein, potency, and bioburden were performed. SDS-PAGE showed ≥ 90% AAT. Based on AAT activity, the concentrates contained a total of 2096 g of AAT, a 79% yield of the activity present in the Cohn paste suspension, and a purification factor of 26.6. Specific activity for each of the three batches ranged from 0.886 to 1.04, and averaged 0.974 mg functional AAT per mg protein.

[0082] The AAT concentrate (2.9-6.8 % protein) was adjusted to 20-25 °C, and sucrose (1.75 kg per kg AAT concentrate) and potassium acetate (0.175 kg per kg AAT concentrate) were added. The final concentration of sucrose was 59.8% ± 6% (w/w), and the final concentration of potassium acetate was 5.98% ± 0.8% (w/w). After mixing, the stabilized concentrate was transferred into one-liter sealed serum bottles. The bottles were stored at 2-8 °C for no more than 10 weeks (and at 15-25 °C for no more than 48 hours) before being heat-treated (pasteurized). Pasteurization at 60 ± 1 °C was performed for 10 - 11 hours. The pasteurized AAT solution was held at 2-8 °C for no more than 10 weeks, and at 15-25 °C for no more than 72 hours, prior to further processing.

[0083] Pasteurized AAT solution was pooled under HEPA-filtered air into two batches, and diluted with diafiltration buffer (20 mM sodium phosphate, 45 mM NaCl, 3% mannitol, pH 6.6-7.4) at a ratio of 5:1 buffer:AAT solution (w/v). The diluted solutions were sampled for LAL, protein, and potency. Based on AAT activity, the pasteurized and diluted solutions contained a total of 1941 g of AAT, a 73% yield of the activity present in the Cohn paste suspension, and a purification factor of 26.6. Specific activities for the two pasteurized batches were 0.954 and 0.993, an average of 0.973 mg functional AAT per mg protein. The percent monomer of the AAT solutions was measured by size-exclusion HPLC before and after pasteurization. The monomer fractions of the AAT concentrates (pre-pasteurization) were 97.1% to 98.5%, averaging 97.7%. The monomer fractions of the two pasteurized and diluted solutions were 95.9% and 97.5%, an average of 96.7%. Only 1.0% of the monomeric form of AAT was polymerized or aggregated during the pasteurization step.

[0084] Two YM100 filter cartridges (Millipore, Bedford, Mass.) were installed in series into a YM100 UF system, with the first cartridge operated in a tangential flow mode and the second cartridge dead-ended. The UF system was recirculated with a minimum of 5 kg diafiltration buffer. Following recirculation, the diafiltration buffer was tested to verify pH (6.8 - 7.2) and LAL (< 0.25 EU/ml). The diafiltration buffer, and all subsequent processing until lyophilization, was at 2-8 °C.

[0085] Each of the pooled AAT solutions was passed through the YM100 cartridges at 2-8 °C at an inlet pressure of ≤ 45 psi. The load did not exceed 1339 grams protein, and the weight of the YM100 filtrate plus postwashes did not exceed 337 kg. The YM100 filtrates were then ultrafiltered and diafiltered, at an inlet pressure of ≤50 psi, against diafiltration buffer (1.60 - 1.90 mg/ml sodium, 10 times the YM100 concentrate weight), using an ultrafilter containing a 10,000 M.W. membrane (≥ 25 ft² surface area) that was dedicated to the post-pasteurization process.

[0086] The diafiltered solutions were sampled inline and tested for sodium. If the sodium level of the permeate was within ± 10%
of the diafiltration buffer sodium concentration, diafiltration was considered complete. If the sodium level was not within ±10% of the diafiltration buffer sodium concentration, diafiltration was repeated with additional diafiltration buffer (5 times the YM100 filtrate weight).

[0087] A final concentration was performed until approximately 6 kg of solution was obtained. Two postwashes were performed using 1.5 kg diafiltration buffer each time. Postwashes were combined with the concentrate for determination of total volume of diafiltered YM100 filtrate. Diafiltered YM100 filtrates were held for no more than 12 days at 2-8 °C before further processing. Based on AAT activity, the diafiltrate contained a total of 1680 g of AAT, a 74% yield of the activity present in the Cohn paste suspension, with a purification factor of 27.5. Specific activities for the two batches were 0.984 and 1.03, an average of 1.01 mg functional AAT per mg protein.

[0088] After addition of diafiltration buffer to obtain a final formulation target of 50 mg functional AAT/ml, the YM100 filtrate solution pH was adjusted as necessary to pH 6.8 - 7.2. Clarification was carried out with a 0.2 micron Pall SLK-7002-NRP Filter (Pall Corp., East Hills, NY). Once clarified, the non-sterile bulk AAT solutions were combined, weighed and sampled for LAL, protein, potency, and bioburden (≤ 100 CFU/ml). The non-sterile bulk AAT was held for no longer than 73.5 hours at 2-8 °C pending sterile filtration. Based on AAT activity, the non-sterile bulk AAT solution contained a total of 1822 g of AAT, a 69% yield of the activity present in the Cohn paste suspension, with a purification factor of 28.8. The specific activity was 0.981 mg functional AAT per mg protein.

[0089] In preparation for sterile filtration, a sterile bulk assembly consisting of a 60 L bulk receiver, a Pall 0.2 micron KA1 NFP2 sterilizing filter and two (2) Millipore 0.2 Micron Aervent™ 50 vent filters was prepared. The assembly was autoclaved and used within 7 days of autoclaving. The non-sterile bulk solution was sterile-filtered with control of temperature (2-8 °C), pressure (≤ 20 psi), filtration time (≤ 120 minutes), and load including postwash (≤ 0.26 kg non-sterile bulk per cm² filter area). The sterile filtrate ultimately obtained from 687 kg of Cohn fraction IV paste contained 1.78 kg of functional AAT, corresponding to an overall yield of 67% based on the activity of the initial Cohn fraction IV 1.4 suspension, and a purification factor of 29.8. The specific activity was 1.09 mg functional AAT per mg protein. The product was >99% AAT by SDS-PAGE, and >95% monomer by size-exclusion HPLC.

[0090] AAT sterile bulk was aseptically filled into 50 ml Type I glass vials using a fill volume targeted to achieve approximately 1000 mg functional AAT activity per vial (i.e. 20.8 g ± 0.2g solution per vial), and the vial contents were frozen and lyophilized.

| Table 4 |
|-------------------------|---|---|---|---|---|---|---|---|
| Fr. IV1,4 Extract | Post- Aerosil | Filtrate | IEC Eluate | HIC Effluent | DF HIC Conc. | Diluted, Pasteur. | YM100 Filtrate | Non-St erile Bulk |
| No. of Batches | 9 | 9 | 9 | 9 | 3 | 3 | 2 | 2 | 1 | 1 |
| Yield (g AAT; total for all batches) | 2658 | 1833* | 1557* | 2241 | 2090 | 2096 | 1941 | 1960 | 1822 | 1780 |
| Overall Yield from Extract | 100% | 69% | 59% | 84% | 79% | 79% | 73% | 74% | 69% | 67% |
| Purification Factor | 1.0 | 1.4 | 1.5 | 16.2 | 25.6 | 26.6 | 26.6 | 27.5 | 26.8 | 29.8 |
| Specific Activity** (mg/mg) | 0.037† | 0.053† | 0.056† | 0.592† | 0.937† | 0.974† | 0.973† | 1.01† | 0.981† | 1.09† |

*The AAT assay for these fractions is believed to be low, due to unidentified interfering factors.
**Specific activities are averages over the number of batches shown.
†The Bradford Protein assay was used for these fractions because they are too impure to determine protein concentration by OD200. The protein standard used in the Bradford assay was purified AAT, calibrated using an extinction coefficient for AAT of 5.3, see R. Pannell, D. Johnson, and J. Travis, Biochemistry 13:5439-5445 (1974).
‡Protein concentration by OD200 using an extinction coefficient for AAT of 5.3.

[0091] Functional AAT yields, and characteristics of the AAT fractions obtained, at each of the above steps are set out in Table 4. Modifications of the above-described methods will be obvious to those of skill in the fields of protein purification, analytical chemistry, medicine, and related fields, and such substitutions and modifications are contemplated.

REFERENCES CITED IN THE DESCRIPTION
This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

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Patentkrav

1. Alpha-1-antitrypsin (AAT) sammensætning egnet til farmaceutisk brug og omfattende:

(a) 6% eller mindre forurenende proteiner baseret på den samlede proteinvægt bestemt ved SDS-PAGE, og
(b) mindre en 0,2% IgA,

hvori den specifikke aktivitet af alpha-1-antitrypsinet er mindst 0,99 mg funktional AAT pr. mg protein, når der som ekstinktionskoefficient anvendes E\text{1}\%\text{1} cm, 280 nm

= 5,3, og

hvori mindre end 8% af sammensætningen har større molekylvægt end monomerisk AAT, og

hvori AAT er udskilt fra en blodplasmafraktion.
Figure 1