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PROTEIN C IN INFLAMMATORY  
CONDITIONS AFFECTING THE  
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(2), (4) Date: **May 15, 2008**(57) **ABSTRACT**

The present invention provides methods for the local treatment of acute and chronic extravascular pulmonary fibrin deposition and/or reducing unwanted effects associated with systemic administration of anticoagulants to a subject via airway administration to the subject by intratracheal, intra-bronchial or intraalveolar routes of human activated protein C or biologically active derivatives thereof.

# AIRWAY ADMINISTRATION OF ACTIVATED PROTEIN C IN INFLAMMATORY CONDITIONS AFFECTING THE RESPIRATORY TRACT

[0001] All patent and non-patent references cited in the application, or in the present application, are also hereby incorporated by reference in their entirety.

## FIELD OF INVENTION

[0002] The present invention provides methods to arrest acute, recurrent and chronic fibrin deposition in the airspaces of the respiratory tract, in particular the alveoli and/or bronchioli, in any disease associated with such deposition in children and adults. These diseases include inflammatory lung diseases such as acute lung injury (ALI), which may be related to direct or indirect pulmonary trauma, e.g. following ventilator therapy (Ventilatory induced Lung Injury (VILI)), inflammatory conditions like auto-immune diseases, pancreatitis, aspiration pneumonitis, inhalation of toxic fumes, acute respiratory distress syndrome (ARDS), which is a more severe manifestation of ALI, infections such as sepsis, severe sepsis and septic shock; pneumonia of whatever cause; acute and chronic bronchoalveolar diseases, fibrosing alveolitis, bronchiolitis, cystic fibrosis and also diseases with severe airway hyperreactivity, e.g. bronchial asthma and drug induced lung insufficiency, e.g. following chemotherapy like bleomycin. In the methods of the present invention, anticoagulants such as activated protein C, whether these agents are derived from plasma or prepared by recombinant DNA technology, are administered intratracheally, intrabronchially, or to the alveolar space via airway administration. These methods are useful in clinical medicine, especially critical or intensive care medicine and respiratory medicine.

## BACKGROUND OF INVENTION

[0003] The deposition of fibrin in the airspaces of the respiratory tract, particularly the alveolar and bronchiolar airspaces, is a frequent complication of systemic inflammatory conditions such as those resulting from trauma, sepsis, severe sepsis and septic shock, drug induced ALI, ARDS or pneumonitis (e.g. due to methotrexate, bleomycin or sirolimus) (Amigues L, Klouche K, Massanet P, Gaillard N, Garrigue V, Beraud J J, Mourad G. Sirolimus-associated acute respiratory distress syndrome in a renal transplant recipient. *Transplant Proc.* 2005; 37:2830-1.) and ventilator-induced lung injury (VILI) (Macintyre N R. Current issues in mechanical ventilation for respiratory failure. *Chest.* 2005; 128:561 S-567S) and lung injury secondary to mechanical ventilation (VILI) (The Acute Respiratory Distress Syndrome Network: Ventilation with lower tidal volumes as compared with traditional tidal volumes for acute lung injury and the acute respiratory distress syndrome. *N Engl J. Med.* 2000; 342:1301-1308).

[0004] These conditions lead to a systemic activation of coagulation, ultimately resulting in the deposition of fibrin in the intravascular and extravascular spaces, i.e. in the alveolar and bronchial compartments. At the same time, the inflammatory activation causes the release of pro-inflammatory cytokines such as tumor necrosis factor alpha (TNF- $\alpha$ ), interleukin 1-beta (IL-1 $\beta$ ), interleukin 6 (IL-6) and interleukin 8 (IL-8) from activated inflammatory cells. Intravascular fibrin deposition and the release of pro-inflammatory cytokines in

the airspaces of the lungs cause a tissue injury characterized by an increased permeability of the alveolar-capillary membrane with diffuse alveolar damage and the accumulation in the alveoli of edema fluid rich in plasma proteins, including the components of the blood coagulation system, and a reduction in surfactant production. As a result, a fibrin-rich hyaline membrane is formed in the alveolar ducts and airspaces. In a later phase, a massive infiltration of neutrophils and other inflammatory cells occurs, followed by organization of the exudates and fibrosis. This pathologic sequence has been described and reviewed by Bellingan G J, 2002: "The pathogenesis of ALI/ARDS", *Thorax* 57:540-546. The clinical conditions corresponding to this pathology are called ALI or ARDS, differing only in that ARDS is more severe and characterized by greater hypoxemia such that the ratio of arterial  $P_{O_2}$  to inspired oxygen fraction ( $P_{aO_2}/F_{I_{O_2}} \leq 200$  mmHg. ALI and ARDS occur as part of the systemic inflammatory response syndrome (SIRS), which can be due infective or non-infective causes such as pancreatitis or direct or indirect pulmonary trauma; when the cause of SIRS is infective, it is called sepsis, which, when associated with organ dysfunction, is defined as severe sepsis, and when associated with significant hypotension, as septic shock.

[0005] A similar sequence of pathological events occurs in pneumonias due to a variety of causes, including viral, bacterial and fungal agents, e.g. *Pneumocystis carinii* (PCP) pneumonia, bronchiolitis (e.g. secondary to viral pneumonitis and/or pulmonary graft-versus-host disease (GVHD)), leading also to alveolar exudates and fibrin deposition in those regions of the lung affected by the inflammatory process.

[0006] It has been pointed out (e.g. by Levi M et al., 2003: "Bronchoalveolar coagulation and fibrinolysis in endotoxemia and pneumonia", *Crit. Care Med* 31:S238-S242) that the lung is particularly susceptible to fibrin deposition in sepsis, showing this phenomenon to a greater degree than other organs. The extensive local fibrin deposition suggests that local activation of coagulation or perturbation of local physiologic regulatory systems may be involved. In the bronchoalveolar compartment, tissue factor (TF), expressed locally on alveolar macrophage cells and on the epithelium, seems to have a pivotal role in the initiation of coagulation, while physiologic anticoagulation due to antithrombin and the protein C system is dysfunctional. It has been documented that the protein C system is markedly disrupted in patients with ALI/ARDS from both septic and non-septic causes and there is evidence of both circulatory and intra-alveolar derangements in the protein C pathway in ALI/ARDS (Ware L B et al., 2003: "Protein C and thrombomodulin in human acute lung injury", *Am J Physiol Lung Cell Mol Physiol* 285:L514-L521).

[0007] At the same time, there is a marked depression of local fibrinolysis, i.e. coagulation is locally upregulated in the injured lung, while fibrinolytic activity is markedly depressed.

## SUMMARY OF INVENTION

[0008] The aim of the present invention is to improve the treatment of ALI, ARDS, pneumonia and inflammatory pulmonary diseases by addressing the local pulmonary activation of the coagulation system and the local deficiency of anticoagulatory mechanisms, by applying the relevant anticoagulants, or agents capable of blocking the local initiation of coagulation, by local administration into the airways. In

this way a high local concentration of these agents can be achieved in the affected airways, so that extravascular fibrin deposition can be more effectively inhibited than by the systemic (intravenous) administration of the same agents, but avoiding or reducing systemic adverse effects. The airway administration of these agents can be given alone or as a supplement to intravenous administration of the same or other agents. Because of the “cross-talk” between coagulation and inflammation, the airway administration of these agents is also expected to modulate local pulmonary inflammation, by reducing local thrombin activation and in certain cases also by direct anti-inflammatory action.

**[0009]** An aspect of the present invention relates to a method for reducing extravascular fibrin deposition in the airways, especially in the alveolar or bronchoalveolar spaces, in human subjects with inflammatory and/or infective pulmonary conditions that lead to such fibrin deposition, the method comprising the administration via the airway of anticoagulants, whether purified from plasma or obtained by recombinant DNA technology.

#### DETAILED DESCRIPTION OF THE INVENTION

**[0010]** The present invention relates to the airway administration, by any appropriate method including, but not limited to, intratracheal, intrabronchial or intraalveolar administration, to a human subject inclusive of both adults and children, of purified or concentrated human activated protein C (APC), or derivatives thereof, however prepared, to prevent or reduce extravascular fibrin deposition in the airways, especially the alveolar or bronchoalveolar airspaces. Such fibrin deposition may result from an acute condition, a recurrent condition or a chronic condition and may be due to a variety of causes, including but not limited to trauma, direct or indirect, inflammation or infection, due to drug induced fibrin deposition in airways and lung interstitium, congenital diseases like cystic fibrosis, or due to a combination of such possible causes. For example, it is believed that the methods of the present invention will be useful in treating the alveolar fibrin deposition characteristic of ALI or ARDS arising in a large proportion of patients with sepsis of varying degrees, in patients with severe pneumonias, bronchiolitis obliterans and in fibrosing alveolitis.

#### DEFINITIONS

**[0011]** Affinity: the strength of binding between receptors and their ligands, for example between an antibody and its antigen.

**[0012]** Amino Acid Residue: That part of the amino acid which is present in the polypeptide chain in which the amino acid is linked to other amino acids by peptide (amide) bonds. The amino acid residues described herein are preferably in the “L” isomeric form. However, the amino acid encompasses every amino acid such as L-amino acid, D-amino acid, alpha-amino acid, beta-amino acid, gamma-amino acid, natural amino acid and synthetic amino acid or the like, as long as the desired functional property is retained by the polypeptide. Further included are natural or synthetic amino acids which have been modified. NH<sub>2</sub> refers to the free amino group present at the amino terminus of a polypeptide. COOH refers to the free carboxy group present at the carboxy terminus of a polypeptide. Standard polypeptide abbreviations for amino acid residues are used herein.

**[0013]** It should be noted that all amino acid residue sequences represented herein by formulae have a left-to-right orientation in the conventional direction of amino terminus to carboxy terminus. Furthermore, it should be noted that a dash at the beginning or end of an amino acid residue sequence indicates a peptide bond to a further sequence of one or more amino acid residues or a covalent bond to an amino-terminal group such as NH<sub>2</sub> or acetyl or to a carboxy-terminal group such as COOH.

**[0014]** Modified amino acid: an amino acid wherein an arbitrary group thereof is chemically modified. In particular, a modified amino acid chemically modified at the alpha-carbon atom in an alpha-amino acid is preferable.

**[0015]** Polypeptide: The phrase polypeptide refers to a molecule comprising amino acid residues which do not contain linkages other than amide linkages between adjacent amino acid residues.

#### APC Molecule

**[0016]** The present invention relates to the use of APC molecule(s) in the manufacture of a medicament for the local treatment of acute and extravascular pulmonary fibrin deposition. The term “APC molecule” is used herein to refer to any molecule capable of binding to and proteolytically destroying the coagulation factors Va and VIIIa and capable of binding to and neutralizing PAI-1 and/or binding specifically to the protein C receptor (Endothelial protein C Receptor EPCR)). Methods for assaying the functional activity of the APC molecules for use in the present invention include those described by Gruber A, Griffin JH. Direct detection of activated protein C in blood from human subjects. *Blood*. 1992 May 1; 79(9): 2340-8, and Strandberg K, Kjellberg M, Knebel R, Lilja H, Stenflo J. A sensitive immunochemical assay for measuring the concentration of the activated protein C-protein C inhibitor complex in plasma: use of a catcher antibody specific for the complexed/cleaved form of the inhibitor. *Thromb Haemost*. 2001; 86(2):604-10.

**[0017]** It is to be understood that the activity of the APC molecules for use in the present invention can be less potent or more potent than native APC.

**[0018]** APC is a vitamin K-dependent serine proteinase and is the activated form of the protein C. APC has both anticoagulant and profibrinolytic properties. Human protein C is a polypeptide of 461 amino acids, which undergoes a number of post-translational modifications to create the activated protein C. These modifications include 1) cleavage of a 42 amino acid signal sequence; 2) cleavage of lysine and arginine residues (positions 156 and 157) to make a 2-chain inactive zymogen (a155 amino acid residue light chain attached via a disulfide bridge to a 262 amino acid residue heavy chain); 3) vitamin K-dependent carboxylation of nine glutamic acid residues located within the aminoterminal 45 residues (gladomain); an, 4 carbohydrate attachment at four sites (one in the light chain and three in the heavy chain). Finally, the 2-chain zymogen may be activated by removal of a dodecapeptide at the N-terminus of the heavy chain.

**[0019]** The present invention further includes the use of recombinant or synthetically or transgenically produced human APC polypeptides. Thus in one embodiment, the APC molecule is a homologue of APC.

**[0020]** The human activated protein C that is intended to be administered according to the present invention comprises naturally occurring human activated protein C, or biologically active analogues of the same, whether prepared from

plasma or recombinantly or transgenically or synthetically produced. Recombinant activated protein C may incorporate modifications (e.g. amino acid substitutions and/or deletions and/or additions of heterologous amino acid sequences), which may result in analogues with enhanced biologic activity. For example, the APC can be produced using eukaryotic cell culture systems (e.g. human kidney 293, HEPG-2, LLC-MK2, CHO or AV12 cells), transgenic animals, transgenic plants, or in vitro systems. In these systems, the protein can be produced as an inactive precursor, which, after purification, is activated by thrombin cleavage and formulated for administration. Alternatively, the APC can be produced by direct secretion of the activated form of protein C.

**[0021]** Details of producing, purifying, activating, and formulating APC are known in the art and are described, for example, in U.S. Pat. No. 6,156,734, which is incorporated by reference herein in its entirety. Also, APC genes and plasmids that can be used in these methods are described in U.S. Pat. Nos. 4,981,952; 4,775,624; and 4,992,373, which are also incorporated by reference herein. Activated protein C can also be obtained from commercial sources. For instance, a specific example of an APC that can be used in the invention is produced by Eli Lilly and Company, under the name of XIGRIS™ (activated recombinant human protein C).

**[0022]** In one preferred embodiment of the present invention, the APC molecule is an APC analogue. An “APC analogue” is defined as a molecule having one or more (such as 20 or fewer, for example 17 or fewer, such as 15 or fewer, for example 13 or fewer, such as 11 or fewer, for example 9 or fewer, such as 7 or fewer, for example 5 or fewer, such as 3 or fewer, for example 2 or fewer, such as 1 or fewer) amino acid substitutions, deletions, inversions, or additions relative to APC and may include D-amino acid forms.

**[0023]** APC analogues also have been described in WO 01/57193 and include APC analogues which are created from protein C analogues containing at least two of the following amino acid substitutions in SEQ ID NO 1: His at position 10 is substituted with Gln; Ser at position 11 is substituted with Gly; Ser at position 12 is substituted with Lys; Gln at position 32 is substituted with Glu; Asn at position 33 is substituted with Asp or Phe; and, the amino acid at position 194, 195, 228, 249, 254, 302, or 316 is substituted with an amino acid selected from Ser, Ala, Thr, His, Lys, Leu, Arg, Asn, Asp, Glu, Gly, and Gln, or the amide form thereof, and pharmaceutically-acceptable salts thereof.

**[0024]** Preferred APC molecules used in the present invention also include analogues of APC in which one or more amino acids which are not present in the original sequence are added or deleted, and derivatives thereof. Such analogues are, for example, described in EP 0 946 715, where the protein C Gla domain was replaced by Gla domains of other vitamin K-dependent polypeptides, such as factor VII, factor X and prothrombin.

**[0025]** In one embodiment of the present invention the APC analogue exhibits an enhanced anticoagulant activity compared with the wild-type protein. In a preferred embodiment the analogue has a higher binding affinity for its substrates, such as for example FVa and FVIIIa, than the wild-type protein. In yet another embodiment the APC analogue has a lower binding affinity for its inhibitors, such as for example protein C inhibitor,  $\alpha$ -antitrypsin or plasminogen activator inhibitor-1, than the wild-type protein. In one embodiment of the present invention the APC analogue exhibits an enhanced profibrinolytic activity compared with the wild-type protein.

In a preferred embodiment the analogue has a higher binding affinity to PAI-1 than the wild-type protein. In another embodiment the APC analogue is capable of inhibiting PAI-1. In a further embodiment the modifications result in a stabilization of the APC analogue. Such analogues are for example described in patent applications WO 99/20767 and WO 03/073980, which are hereby incorporated by reference in their entirety. APC analogues can be used according to the present invention alone or in combination with other APC analogues and or homologues and/or derivatives and/or conjugates. For example an APC analogue with a higher binding affinity for FVa and/or FVIIIa can be used in combination with an APC homologue capable binding selectively to and inhibiting PAI-1.

**[0026]** In another preferred embodiment of the present invention, the APC molecule is an APC derivative. An “APC derivative” is defined as a molecule having the amino acid sequence of APC or of an APC analogue, but additionally comprises chemical modification of one or more of its amino acid side groups, alpha-carbon atoms, terminal amino group, or terminal carboxylic acid group. A chemical modification includes, but is not limited to, adding chemical moieties, creating new bonds, and removing chemical moieties. Modifications at amino acid side groups include, without limitation, acylation of lysine epsilon-amino groups, N-alkylation of arginine, histidine, or lysine, alkylation of glutamic or aspartic carboxylic acid groups, and deamidation of glutamine or asparagine. Modifications of the terminal amino include, without limitation, the des-amino, N-lower alkyl, N-di-lower alkyl, and N-acyl modifications. Modifications of the terminal carboxy group include, without limitation, the amide, lower alkyl amide, dialkyl amide, and lower alkyl ester modifications. Lower alkyl is C1-C4 alkyl. Furthermore, one or more side groups, or terminal groups, may be protected by protective groups known to the ordinarily-skilled protein chemist. The alpha-carbon of an amino acid may be mono- or dimethylated.

#### Homologues of APC Molecules

**[0027]** A homologue of one or more of the sequences specified herein may vary in one or more amino acids as compared to the sequences defined, but is capable of performing the same function, i.e. a homologue may be envisaged as a functional equivalent of a predetermined sequence.

**[0028]** Thus, in one preferred embodiment of the present invention, the APC molecule is a homologue of any of the molecules disclosed herein, such as a homolog of any of the molecules selected from the group consisting of:

**[0029]** APC

**[0030]** XIGRIS

**[0031]** Drotrecogin alfa activated

**[0032]** Thus, in one preferred embodiment of the present invention, the APC molecule is a peptide containing one or more amino acid substitutions, inversion, additions and/or deletions, compared with any of the molecules disclosed herein, such as a molecule selected from the group consisting of:

**[0033]** APC

**[0034]** XIGRIS

**[0035]** Drotrecogin alfa

**[0036]** In one embodiment, the number of substitutions, deletions, or additions is 20 amino acids or less, such as 15 amino acids or less, for example 10 amino acids or less, such as 9 amino acids or less, for example 8 amino acids or less,

such as 7 amino acids or less, for example 6 amino acids or less, such as 5 amino acids or less, for example 4 amino acids or less, such as 3 amino acids or less, for example 2 amino acids or less (such as 1), or any integer in between these amounts. In one aspect of the invention, the substitutions include one or more conservative substitutions, such as 20 or fewer conservative substitutions, for example 18 or fewer, such as 16 or fewer, for example 14 or fewer, such as 12 or fewer, for example 10 or fewer, such as 8 or fewer, for example 6 or fewer, such as 4 or fewer, for example 3 or fewer, such as 2 or fewer conservative substitutions. A “conservative” substitution denotes the replacement of an amino acid residue by another, related amino acid residue belonging to the same group of amino acids, such as those with a hydrophobic side chain, those with an aromatic side chain, those with a basic side-chain, those with an acidic side chain, those with a hydroxyl side chain and those with a non-ionized polar side chain. Examples of conservative substitution include the substitution of one hydrophobic residue, such as isoleucine, valine, leucine or methionine for another, or the substitution of one basic residue for another, such as the substitution of arginine for lysine, or the substitution of one acidic residue for another, such as glutamic acid for aspartic acid, or the substitution of one non-ionized polar residue for another, such as the substitution of glutamine for asparagine, and the like. The following table lists illustrative, but non-limiting, conservative amino acid substitutions.

| Original residue | Exemplary substitutions |
|------------------|-------------------------|
| Ala              | Ser, Thr, Val, Gly      |
| Arg              | Lys                     |
| Asn              | His, Ser                |
| Asp              | Glu, Asn                |
| Cys              | Ser                     |
| Gln              | Asn, His                |
| Glu              | Asp, Glu                |
| Gly              | Ala, Ser                |
| His              | Asn, Gln                |
| Ile              | Leu, Val, Thr           |
| Leu              | Ile, Val                |
| Lys              | Arg, Gln, Glu, Thr      |
| Met              | Leu, Ile, Val           |
| Phe              | Leu, Tyr                |
| Ser              | Thr, Ala, Asn           |
| Thr              | Ser, Ala                |
| Trp              | Arg, Ser                |
| Tyr              | Phe                     |
| Val              | Ile, Leu, Ala           |
| Pro              | Ala                     |

[0037] Other APC homologues suitable for the uses and methods of the present invention are peptide sequences having greater than 50 percent sequence identity, and preferably greater than 90 percent sequence identity (such as greater than 91% sequence identity, for example greater than 92% sequence identity, such as greater than 93% sequence identity, for example greater than 94% sequence identity, such as greater than 95% sequence identity, for example greater than 96% sequence identity, such as greater than 97% sequence identity, for example greater than 98% sequence identity, such as greater than 99% sequence identity, for example greater than 99.5% sequence identity), to any of the molecules disclosed herein, such as a molecule selected from the group consisting of:

[0038] APC

[0039] XIGRIS

[0040] Drotrecogin alfa

[0041] As used herein, sequence identity refers to a comparison made between two molecules using standard algorithms well known in the art. The preferred algorithm for calculating sequence identity for the present invention is the Smith-Waterman algorithm, where the reference sequence is used to define the percentage identity of polypeptide homologs over its length. The choice of parameter values for matches, mismatches, and inserts or deletions is arbitrary, although some parameter values have been found to yield more biologically realistic results than others. One preferred set of parameter values for the Smith-Waterman algorithm is set forth in the “maximum similarity segments” approach, which uses values of 1 for a matched residue and  $-1/3$  for a mismatched residue (a residue being either a single nucleotide or single amino acid) (Waterman, Bull. Math. Biol. 46, 473-500 (1984)). Insertions and deletions (indels), x, are weighted as

$$x_k = 1 + k/3,$$

where k is the number of residues in a given insert or deletion (Id.).

[0042] For instance, a sequence that is identical to a 42 amino acid residue sequence, except for 18 amino acid substitutions and an insertion of 3 amino acids, would have a percent identity given by:

$$[(1 \times 42 \text{ matches}) - (1/3 \times 18 \text{ mismatches}) - (1 + 3/3 \text{ indels})]/42 = 81\% \text{ identity}$$

[0043] In one preferred embodiment of the present invention, truncations at the end of the molecule are not taken into account when calculating sequence identity (i.e., if one molecule is longer than the other, only the overlapping lengths of the molecules are used in the sequence identity analysis); in another preferred embodiment of the present invention, truncations are counted as deletions.

[0044] An APC homologue may include D-amino acid forms and may be a molecule having one or more amino acid substitutions, deletions, inversions, or additions relative to any of the molecules disclosed herein, such as a molecule selected from the group consisting of:

[0045] APC

[0046] XIGRIS

[0047] Drotrecogin alfa

[0048] In one preferred embodiment of the present invention, the APC molecule is a peptide containing one or more amino acid substitutions, inversion, additions or deletions, compared with APC. In one embodiment, the number of substitutions, deletions, or additions is 20 amino acids or less,

such as 15 amino acids or less, for example 10 amino acids or less, such as 9 amino acids or less, for example 8 amino acids or less, such as 7 amino acids or less, for example 6 amino acids or less, such as 5 amino acids or less, for example 4 amino acids or less, such as 3 amino acids or less, for example 2 amino acids or less (such as 1), or any integer in between these amounts. In one aspect of the invention, the substitutions include one or more conservative substitutions. Examples of suitable conservative substitutions are given above.

**[0049]** Other APC homologues suitable for the uses and methods of the present invention are peptide sequences derived from protein C sequences having greater than 50 percent sequence identity, and preferably greater than 90 percent sequence identity (such as greater than 91% sequence identity, for example greater than 92% sequence identity, such as greater than 93% sequence identity, for example greater than 94% sequence identity, such as greater than 95% sequence identity, for example greater than 96% sequence identity, such as greater than 97% sequence identity, for example greater than 98% sequence identity, such as greater than 99% sequence identity, for example greater than 99.5% sequence identity), to (1) SEQ ID NO:1 and/or (2) to truncated sequences thereof. As used herein, sequence identity refers to a comparison made between two molecules using standard algorithms well known in the art. The preferred algorithm for calculating sequence identity for the present invention is the Smith-Waterman algorithm, as described above.

**[0050]** An APC homologue may also be a molecule having one or more amino acid substitutions, deletions, inversions, or additions relative to human APC and may include D-amino acid forms.

**[0051]** In another embodiment of the present invention, said homologue of any of the pre-determined sequences herein, such as SEQ ID NO: 1 may be defined as:

**[0052]** i) homologues comprising an amino acid sequence capable of binding selectively to the coagulation factors Va and VIIIa, and/or

**[0053]** ii) homologues having a substantially similar or higher binding affinity to the coagulation factors Va and VIIIa than human APC, and/or

**[0054]** iii) homologues comprising an amino acid sequence capable of binding selectively to and inhibiting PAI-1 and/or

**[0055]** iv) homologues having a substantially similar or higher binding affinity to PAI-1 and/or

**[0056]** v) homologues having a substantially similar or higher binding affinity to endothelial protein C receptor (EPCR) and/or

**[0057]** vi) homologues having substantially similar, higher or lower half life after deposition in the airways

#### Chemically Derivatized APC Molecules

**[0058]** It is further understood that APC molecules suitable for use in the present invention may be chemically derivatized or altered, for example, peptides with non-natural amino acid residues (e.g., taurine residue, beta- and gamma-amino acid residues and D-amino acid residues), C-terminal functional group modifications, such as amides, esters, and C-terminal ketone modifications and N-terminal functional group modi-

fications, such as acylated amines, Schiff bases, or cyclization, such as found, for example, in the amino acid pyroglutamic acid.

#### APC Molecule Conjugates

**[0059]** The APC molecules of the present invention may also be modified with non-polypeptide moieties, such as PEG or sugar moieties, to yield APC compounds that have an increased resistance to inactivation by e.g. human plasma and  $\alpha$ 1-antitrypsin, therefore displaying an increased in vivo half-life. Preferred examples include protein C conjugates, wherein at least one additional in vivo N-glycosylation site has been introduced.

**[0060]** Suitable examples are described in U.S. Pat. No. 6,933,367 ("Protein C or activated protein C-like molecules"), the contents of which are incorporated by reference herein.

**[0061]** Thus, in one embodiment of the present invention, the APC molecule used is an N-glycosylated APC polypeptide or an analogue thereof.

**[0062]** In another embodiment of the present invention, the APC molecule is an O-glycosylated APC polypeptide or an analogue thereof.

**[0063]** In a further embodiment, the APC molecule, wherein at least one additional N-glycosylation site has been introduced.

**[0064]** In another embodiment of the present invention, the APC molecule used is an APC polypeptide or an analogue thereof further containing and N-linked or O-linked fatty acid.

#### APC Fragments

**[0065]** In one embodiment the APC molecule can be a APC fragment. A fragment is a portion of APC, APC homologue or APC derivative. Examples of fragments include EGF-like domains 1 or 2, or the serine protease domain or deletions of the N-terminus or the C-terminus of the natural APC peptide or both. Fragments of APC comprise at least 20 consecutive amino acids from SEQ ID NO: 1. For example a fragment can be 20 amino acids or more, such as 25 amino acids or more, for example 30 amino acids or more, such as 50 amino acids or more, for example 100 amino acids or more, such as 150 amino acids or more, for example 200 amino acids or more, such as 250 amino acids or more, for example 300 amino acids or more, such as 350 amino acids or more, for example 400 amino acids or more, such as 450 amino acids or more, for example 460 amino acids in length or any integer in between these amounts.

#### Intratracheal, Intrabronchial or Intraalveolar Administration

**[0066]** Methods of administration include, but are not limited to, spraying, lavage, inhalation, flushing or installation, using as fluid a physiologically acceptable composition in which the blood coagulation factor or factors have been dissolved. When used herein the terms "intratracheal, intrabronchial or intraalveolar administration" include all forms of such administration whereby the coagulation factor is applied into the trachea, the bronchi or the alveoli, respectively, whether by the instillation of a solution of the factor, by applying the factor in a powder form, or by allowing the factor to reach the relevant part of the airway by inhalation of the factor as an aerosolized or nebulized solution or powder or gel, with or without added stabilizers or other excipients.

[0067] In another embodiment, intratracheal, intrabronchial or intraalveolar administration does not include inhalation of the product but the instillation or application of a solution of the factor or a powder or a gel containing the factor into the trachea or lower airways.

[0068] Methods of intrabronchial/alveolar administration include, but are not limited to, bronchoalveolar lavage (BAL) administration according to methods well known to those skilled in the art, using as a lavage fluid a physiologically acceptable composition in which the APC and/or a APC homolog and/or derivative and/or conjugate has been dissolved, or indeed by any other effective form of intrabronchial administration including the use of nebulized powders containing the anticoagulant in dry form, with or without excipients, or the direct application of the anticoagulant in solution or powder or gel form during bronchoscopy. Methods of intratracheal administration include, but are not limited to, blind tracheal washing with a similar solution of dissolved activated protein C, or the inhalation of nebulized aerosolized fluid droplets containing the dissolved activated protein C obtained by use of any nebulizing apparatus adequate for this purpose.

[0069] The present invention provides a useful new addition to the methods of treating ALI, ARDS, pneumonia and other conditions associated with bronchoalveolar fibrin deposition. Furthermore, the administration of the anticoagulants via the airway is expected to avoid the unwanted hemorrhagic adverse effects of systemic administration of anticoagulants such as APC, whose intravenous use is associated with a significant incidence of internal hemorrhage including cerebral hemorrhage. At the same time, the application of the anticoagulants via the airway is expected to potentiate their effect on extravascular fibrin deposition in the lungs when compared with their systemic administration. It is expected that the total dosage of an anticoagulant and anti-inflammatory agent such as APC may be used alone locally within the air-spaces or may be divided between the conventional intravenous route and the airway route of the present invention to obtain the optimal balance between the systemic and local pulmonary effects of the treatment, and a reduced incidence of drug adverse effect, for example in patients with severe sepsis, septic shock and ARDS. Furthermore, the time interval ("window of opportunity") during which the intravenous use of APC is likely to be beneficial is limited. A longer time interval of drug response may be expected when the agent is used in the post-septic phase or even in the late ARDS phase dominated by alveolar fibrin deposition, e.g. as seen in ALI and ARDS.

[0070] A preferred embodiment of the present invention comprises local intrabronchial administration to human patients with ARDS of APC by means of bronchoalveolar lavage with lavage fluid (e.g. 25 ml to 100 ml of isotonic saline) in which a suitable dose (e.g. 2 mg to 5 mg or more) of APC has been dissolved. This administration is repeated at intervals during one or more days depending on the duration of either early or late phases of ALI or ARDS. As supplementary or combinatorial treatments in patients fulfilling the indications for intravenous administration of APC, APC can also be given by intravenous infusion.

[0071] Other preferred methods of administration may include using the following devices:

- [0072] 1. Pressurized nebulizers using compressed air/oxygen mixture
- [0073] 2. Ultrasonic nebulizers

[0074] 3. Electronic micropump nebulizers (e.g. Aeroneb Professional Nebulizer)

[0075] 4. Metered dose inhaler (MDI)

[0076] 5. Dry powder inhaler systems (DPI),

The aerosol may be delivered by via a) facemasks or b) via endotracheal tubes in intubated patients during mechanical ventilation (device 1, 2 and 3). The devices 4 and 5 can also be used by the patient without assistance provided that the patient is able to self-activate the aerosol device.

[0077] Preferred concentrations for a solution comprising APC and/or homologues and/or derivatives of APC are in the range of 0.1 µg to 10000 µg active ingredient per ml solution. Using monomeric forms of the compounds, the suitable concentrations are often in the range of from 0.1 µg to 5000 µg per ml solution, such as in the range of from about 0.1 µg to 3000 µg per ml solution, and especially in the range of from about 0.1 µg to 1000 µg per ml solution, such as in the range of from about 0.1 µg to 250 µg per ml solution. A preferred concentration would be from about 0.1 to about 5.0 mg, preferably from about 0.3 mg to about 3.0 mg, such as from about 0.5 to about 1.5 mg and especially in the range from 0.8 to 1.0 mg per ml solution. Using multimeric forms of the compounds, the suitable concentrations are often in the range of from 0.1 µg to 1000 µg per ml solution, such as in the range of from about 0.1 µg to 750 µg per ml solution, and especially in the range of from about 0.1 µg to 500 µg per ml solution, such as in the range of from about 0.1 µg to 250 µg per ml solution. A preferred concentration would be from about 0.1 to about 5.0 mg, preferably from about 0.3 mg to about 3.0 mg, such as from about 0.5 to about 1.5 mg and especially in the range from 0.8 to 1.0 mg per ml solution.

#### Indications

[0078] One aspect of the present invention relates to a method of treating or preventing extravascular fibrin deposition in the airways. Thus, the present invention relates to the treatment of individuals suffering from, or at risk of suffering from, extravascular fibrin depositions caused by an inflammatory lung disease.

[0079] In a preferred aspect the inflammatory lung disease is selected from the group consisting of:

- [0080] ALI
- [0081] ARDS
- [0082] pneumonia
- [0083] acute bronchoalveolar disease
- [0084] chronic bronchoalveolar disease
- [0085] fibrosing alveolitis or
- [0086] bronchial asthma
- [0087] Alveolitis
- [0088] Bronchiolitis
- [0089] Bronchiolitis obliterans-organizing pneumonia (BOPA)
- [0090] Graft-versus-host disease (GVHD)
- [0091] *Pneumocystis carinii* pneumonia (PCP)
- [0092] Pneumonitis, e.g. aspiration pneumonitis
- [0093] Drug-induced pneumonitis (e.g. due to methotrexate, bleomycin or sirolimus)
- [0094] Fibrosing alveolitis, acute or chronic
- [0095] Cystic fibrosis
- [0096] Idiopathic pulmonary fibrosis

[0097] In another embodiment, the inflammatory lung disease is related to a condition selected from the group consisting of:

- [0098] direct or indirect pulmonary trauma
- [0099] pancreatitis

- [0100] aspiration pneumonitis
- [0101] sepsis
- [0102] severe sepsis and/or
- [0103] septic shock
- [0104] *Pneumocystis Carinii* Pneumonia (PCP) as an adjuvant prophylactic or pre-emptive therapy or as a treatment of PCP manifest ARDS induced by PCP, i.e. before or early in the PCP phase or in manifest ARDS concomitant with the anti-*Pneumocystis* antibiotic therapy with e.g. Sulfamethoxazol with Trimethoprim.

#### Pharmaceutical Compositions

[0105] Pharmaceutical compositions or formulations for use in the present invention include an APC preparation in combination with, preferably dissolved in, a pharmaceutically acceptable carrier, preferably an aqueous carrier or diluent. The pharmaceutical composition may be a solid, a liquid, a gel or an aerosol. A variety of aqueous carriers may be used, such as 0.9% saline, buffered saline, physiologically compatible buffers and the like. The compositions may be sterilized by conventional techniques well known to those skilled in the art. The resulting aqueous solutions may be packaged for use or filtered under aseptic conditions and freeze-dried, the freeze-dried preparation being dissolved in a sterile aqueous solution prior to administration.

[0106] The compositions may contain pharmaceutically acceptable auxiliary substances or adjuvants, including, without limitation, pH adjusting and buffering agents and/or tonicity adjusting agents, such as, for example, sodium acetate, sodium lactate, sodium chloride, potassium chloride, calcium chloride, etc. The formulations may contain pharmaceutically acceptable carriers and excipients including microspheres, liposomes, microcapsules, nanoparticles or the like. Conventional liposomes are typically composed of phospholipids (neutral or negatively charged) and/or cholesterol. The liposomes are vesicular structures based on lipid bilayers surrounding aqueous compartments. They can vary in their physiochemical properties such as size, lipid composition, surface charge and number and fluidity of the phospholipid bilayers. The most frequently used lipid for liposome formation are: 1,2-Dilauroyl-sn-Glycero-3-Phosphocholine (DLPC), 1,2-Dimyristoyl-sn-Glycero-3-Phosphocholine (DMPC), 1,2-Dipalmitoyl-sn-Glycero-3-Phosphocholine (DPPC), 1,2-Distearoyl-sn-Glycero-3-Phosphocholine (DSPC), 1,2-Dioleoyl-sn-Glycero-3-Phosphocholine (DOPC), 1,2-Dimyristoyl-sn-Glycero-3-Phosphoethanolamine (DMPE), 1,2-Dipalmitoyl-sn-Glycero-3-Phosphoethanolamine (DPPE), 1,2-Dioleoyl-sn-Glycero-3-Phosphoethanolamine (DOPE), 1,2-Dimyristoyl-sn-Glycero-3-Phosphate (Monosodium Salt) (DMPA), 1,2-Dipalmitoyl-sn-Glycero-3-Phosphate (Monosodium Salt) (DPPA), 1,2-Dioleoyl-sn-Glycero-3-Phosphate (Monosodium Salt) (DOPA), 1,2-Dimyristoyl-sn-Glycero-3-[Phospho-rac-(1-glycerol)] (Sodium Salt) (DMPG), 1,2-Dipalmitoyl-sn-Glycero-3-[Phospho-rac-(1-glycerol)] (Sodium Salt) (DPPG), 1,2-Dioleoyl-sn-Glycero-3-[Phospho-rac-(1-glycerol)] (Sodium Salt) (DOPG), 1,2-Dimyristoyl-sn-Glycero-3-[Phospho-L-Serine] (Sodium Salt) (DMPS), 1,2-Dipalmitoyl-sn-Glycero-3-[Phospho-L-Serine] (Sodium Salt) (DPPS), 1,2-Dioleoyl-sn-Glycero-3-[Phospho-L-Serine] (Sodium Salt) (DOPS), 1,2-Dioleoyl-sn-Glycero-3-Phosphoethanolamine-N-(glutaryl) (Sodium Salt) and 1,1',2,2'-Tetramyristoyl Cardiolipin (Ammonium Salt). Formulations composed of

DPPC in combination with other lipids or modifiers of liposomes are preferred e.g. in combination with cholesterol and/or phosphatidylcholine.

[0107] Long-circulating liposomes are characterized by their ability to extravasate at body sites where the permeability of the vascular wall is increased. The most popular way of producing long-circulating liposomes is to attach hydrophilic polymer polyethylene glycol (PEG) covalently to the outer surface of the liposome. Some of the preferred lipids are: 1,2-Dipalmitoyl-sn-Glycero-3-Phosphoethanolamine-N-[Methoxy(Polyethylene glycol)-2000] (Ammonium Salt), 1,2-Dipalmitoyl-sn-Glycero-3-Phosphoethanolamine-N-[Methoxy(Polyethylene glycol)-5000] (Ammonium Salt), 1,2-Dioleoyl-3-Trimethylammonium-Propane (Chloride Salt) (DOTAP).

[0108] Possible lipids applicable for liposomes are supplied by Avanti, Polar Lipids, Inc, Alabaster, Ala. Additionally, the liposome suspension may include lipid-protective agents which protect lipids against free-radical and lipid-peroxidative damage on storage. Lipophilic free-radical quenchers, such as alpha-tocopherol and water-soluble iron-specific chelators, such as ferrioxamine, are preferred.

[0109] A variety of methods are available for preparing liposomes, as described in, e.g., Szoka et al., Ann. Rev. Biophys. Bioeng. 9:467 (1980), U.S. Pat. Nos. 4,235,871, 4,501,728 and 4,837,028, all of which are incorporated herein by reference. Another method produces multilamellar vesicles of heterogeneous sizes. In this method, the vesicle-forming lipids are dissolved in a suitable organic solvent or solvent system and dried under vacuum or an inert gas to form a thin lipid film. If desired, the film may be redissolved in a suitable solvent, such as tertiary butanol, and then lyophilized to form a more homogeneous lipid mixture which is in a more easily hydrated powder-like form. This film is covered with an aqueous solution of the targeted drug and the targeting component and allowed to hydrate, typically over a 15-60 minute period with agitation. The size distribution of the resulting multilamellar vesicles can be shifted toward smaller sizes by hydrating the lipids under more vigorous agitation conditions or by adding solubilizing detergents such as deoxycholate.

[0110] Micelles are formed by surfactants (molecules that contain a hydrophobic portion and one or more ionic or otherwise strongly hydrophilic groups) in aqueous solution.

[0111] Common surfactants well known to one of skill in the art can be used in the micelles of the present invention. Suitable surfactants include sodium laurate, sodium oleate, sodium lauryl sulfate, octaoxyethylene glycol monododecyl ether, octoxynol 9 and PLURONIC F-127 (Wyandotte Chemicals Corp.). Preferred surfactants are nonionic polyoxyethylene and polyoxypropylene detergents compatible with IV injection such as, TWEEN-80, PLURONIC F-68, n-octyl-beta-D-glucopyranoside, and the like. In addition, phospholipids, such as those described for use in the production of liposomes, may also be used for micelle formation.

[0112] In some cases, it will be advantageous to include a compound, which promotes delivery of the active substance to its target.

#### Dosing Regimes

[0113] The preparations are administered in a manner compatible with the dosage formulation, and in such amount as will be therapeutically effective. The quantity to be administered depends on the subject to be treated, including, e.g. the weight and age of the subject, the disease to be treated and the

stage of disease. Suitable dosage ranges are per kilo body weight normally of the order of several hundred  $\mu\text{g}$  active ingredient per administration with a preferred range of from about 0.1  $\mu\text{g}$  to 10000  $\mu\text{g}$  per kilo body weight. Using monomeric forms of the compounds, the suitable dosages are often in the range of from 0.1  $\mu\text{g}$  to 5000  $\mu\text{g}$  per kilo body weight, such as in the range of from about 0.1  $\mu\text{g}$  to 3000  $\mu\text{g}$  per kilo body weight, and especially in the range of from about 0.1  $\mu\text{g}$  to 1000  $\mu\text{g}$  per kilo body weight. Using multimeric forms of the compounds, the suitable dosages are often in the range of from 0.1  $\mu\text{g}$  to 1000  $\mu\text{g}$  per kilo body weight, such as in the range of from about 0.1  $\mu\text{g}$  to 750  $\mu\text{g}$  per kilo body weight, and especially in the range of from about 0.1  $\mu\text{g}$  to 500  $\mu\text{g}$  per kilo body weight such as in the range of from about 0.1  $\mu\text{g}$  to 250  $\mu\text{g}$  per kilo body weight. A preferred dosage would be from about 0.1 to about 5.0 mg, preferably from about 0.3 mg to about 3.0 mg, such as from about 0.5 to about 1.5 mg and especially in the range from 0.8 to 1.0 mg per administration. Administration may be performed once or may be followed by subsequent administrations. The dosage will also depend on the route of administration and will vary with the age, sex and weight of the subject to be treated. A preferred dosage of multimeric forms would be in the interval 1 mg to 70 mg per 70 kilo body weight.

[0114] Suitable daily dosage ranges are per kilo body weight per day normally of the order of several hundred  $\mu\text{g}$  active ingredient per day with a preferred range of from about 0.1  $\mu\text{g}$  to 10000  $\mu\text{g}$  per kilo body weight per day. Using monomeric forms of the compounds, the suitable dosages are often in the range of from 0.1  $\mu\text{g}$  to 5000  $\mu\text{g}$  per kilo body weight per day, such as in the range of from about 0.1  $\mu\text{g}$  to 3000  $\mu\text{g}$  per kilo body weight per day, and especially in the range of from about 0.1  $\mu\text{g}$  to 1000  $\mu\text{g}$  per kilo body weight per day. Using multimeric forms of the compounds, the suitable dosages are often in the range of from 0.1  $\mu\text{g}$  to 1000  $\mu\text{g}$  per kilo body weight per day, such as in the range of from about 0.1  $\mu\text{g}$  to 750  $\mu\text{g}$  per kilo body weight per day, and especially in the range of from about 0.1  $\mu\text{g}$  to 500  $\mu\text{g}$  per kilo body weight per day, such as in the range of from about 0.1  $\mu\text{g}$  to 250  $\mu\text{g}$  per kilo body weight per day. A preferred dosage would be from about 0.1 to about 100  $\mu\text{g}$ , preferably from about 0.1  $\mu\text{g}$  to about 50  $\mu\text{g}$ , such as from about 0.3 to about 30  $\mu\text{g}$  and especially in the range from 1.0 to 10  $\mu\text{g}$  per kilo body weight per day. Administration may be performed once or may be followed by subsequent administrations. The dosage will also depend on the route of administration and will vary with the age, sex and weight of the subject to be treated. A preferred dosage of multimeric forms would be in the interval 1 mg to 70 mg per 70 kilo body weight per day.

#### Medical Packaging

[0115] The compounds used in the invention may be administered alone or in combination with pharmaceutically acceptable carriers or excipients, in either single or multiple doses. The formulations may conveniently be presented in unit dosage form by methods known to those skilled in the art.

[0116] It is preferred that the compounds according to the invention are provided in a kit. Such a kit typically contains an active compound in dosage forms for administration. A dosage form contains a sufficient amount of active compound such that a desirable effect can be obtained when administered to a subject.

[0117] Thus, it is preferred that the medical packaging comprises an amount of dosage units corresponding to the

relevant dosage regimen. Accordingly, in one embodiment, the medical packaging comprises a pharmaceutical composition comprising a compound as defined above or a pharmaceutically acceptable salt thereof and pharmaceutically acceptable carriers, vehicles and/or excipients, said packaging comprising from 1 to 7 dosage units, thereby having dosage units for one or more days, or from 7 to 21 dosage units, or multiples thereof, thereby having dosage units for one week of administration or several weeks of administration.

[0118] The dosage units can be as defined above. The medical packaging may be in any suitable form for intratracheal, intrabronchial or intraalveolar administration. In a preferred embodiment the packaging is in the form of a vial, ampule, tube, blister pack, cartridge or capsule.

[0119] When the medical packaging comprises more than one dosage unit, it is preferred that the medical packaging is provided with a mechanism to adjust each administration to one dosage unit only.

[0120] Preferably, a kit contains instructions indicating the use of the dosage form to achieve a desirable affect and the amount of dosage form to be taken over a specified time period. Accordingly, in one embodiment the medical packaging comprises instructions for administering the pharmaceutical composition.

#### EXAMPLES

##### Example 1

[0121] A 60 year-old patient developed severe ARDS with opacities mainly localized to the right lung as judged from chest radiography. Bronchoalveolar lavage (BAL) was performed in the right lower lobe bronchus via a bronchoscope with a single dose of 5 mg drotrecogin-alfa activated (Xigris) dissolved in 20 ml of normal saline. There were no apparent adverse effects and no bleeding from the airways was observed. The following day there was some clearing of the opacities in the right lower lobe which had been treated with BAL. On the other hand, the single dose of APC did not improve gas exchange as monitored by the  $\text{PaO}_2/\text{FiO}_2$  ratio.

##### Example 2

[0122] An 11 year old boy with Mb. Hodgkin 2003 initially treated with chemotherapy. A relapse was documented in 2005, where he was treated with radiation therapy and autologous bone marrow transplantation (BMT). Lung biopsy performed before the ICU admittance showed: No sign of pulmonary graft versus host disease (GvHD). No signs of Lymphoma infiltration Admitted to the ICU 9 days after the BMT for acute lung insufficiency (ALI) and continuous high fever with sepsis. Treated initially with broad spectrum empiric antibiotics. At day 4 oxygenation capacity was further reduced with a  $\text{PaO}_2/\text{FiO}_2$  ratio <200 in spite of high PEEP. Chest film showed bilateral diffuse infiltrations localized to all 4 quadrants of the lung field. Echocardiography showed no cardiac shunt or valve insufficiency with normal myocardial function No other organ dysfunction was detected. Patient has mono organ dysfunction with severe ARDS. Bronchoalveolar lavage (BAL) was without *Candida* spp. *Aspergillus*, virus or bacteria. Based on an Institutional Review Board (IRB) decision inhalation therapy with activated protein C (Drotrecogin alpha activated) in a dose of 5  $\text{mg} \times 3$  via nebulizer (Aeroneb®) was commenced. Already

after 3 doses after 12 hours later, the infiltrates on chest film was clearly reduced; concomitantly the arterial oxygen saturation increased from 85% to 95% whilst on FiO<sub>2</sub>=1.0. There were not at any time serious adverse effects, e.g. bleeding

from the airways during or after the APC inhalation. 24 hours after initiation of the inhalation therapy with APC the patient died from circulatory failure due to failing right heart ventricle, i.e. acute cor pulmonale.

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SEQUENCE LISTING

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<210> SEQ ID NO 1

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<212> TYPE: PRT

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20     25     30
Asn Val Asp Asp Thr Leu Ala Phe Trp Ser Lys His Val Asp Gly Asp
35     40     45
Gln Cys Leu Val Leu Pro Leu Glu His Pro Cys Ala Ser Leu Cys Cys
50     55     60
Gly His Gly Thr Cys Ile Asp Gly Ile Gly Ser Phe Ser Cys Asp Cys
65     70     75     80
Arg Ser Gly Trp Glu Gly Arg Phe Cys Gln Arg Glu Val Ser Phe Leu
85     90     95
Asn Cys Ser Leu Asp Asn Gly Gly Cys Thr His Tyr Cys Leu Glu Glu
100    105    110
Val Gly Trp Arg Arg Cys Ser Cys Ala Pro Gly Tyr Lys Leu Gly Asp
115    120    125
Asp Leu Leu Gln Cys His Pro Ala Val Lys Phe Pro Cys Gly Arg Pro
130    135    140
Trp Lys Arg Met Glu Lys Lys Arg Ser His Leu Lys Arg Asp Thr Glu
145    150    155    160
Asp Gln Glu Asp Gln Val Asp Pro Arg Leu Ile Asp Gly Lys Met Thr
165    170    175
Arg Arg Gly Asp Ser Pro Trp Gln Val Val Leu Leu Asp Ser Lys Lys
180    185    190
Lys Leu Ala Cys Gly Ala Val Leu Ile His Pro Ser Trp Val Leu Thr
195    200    205
Ala Ala His Cys Met Asp Glu Ser Lys Lys Leu Leu Val Arg Leu Gly
210    215    220
Glu Tyr Asp Leu Arg Arg Trp Glu Lys Trp Glu Leu Asp Leu Asp Ile
225    230    235    240
Lys Glu Val Phe Val His Pro Asn Tyr Ser Lys Ser Thr Thr Asp Asn
245    250    255
Asp Ile Ala Leu Leu His Leu Ala Gln Pro Ala Thr Leu Ser Gln Thr
260    265    270
Ile Val Pro Ile Cys Leu Pro Asp Ser Gly Leu Ala Glu Arg Glu Leu
275    280    285
Asn Gln Ala Gly Gln Glu Thr Leu Val Thr Gly Trp Gly Tyr His Ser
290    295    300
Ser Arg Glu Lys Glu Ala Lys Arg Asn Arg Thr Phe Val Leu Asn Phe

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-continued

|     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |  |
|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| 305 | 310 |     |     |     |     |     |     | 315 |     |     |     |     |     |     | 320 |     |  |
| Ile | Lys | Ile | Pro | Val | Val | Pro | His | Asn | Glu | Cys | Ser | Glu | Val | Met | Ser |     |  |
|     |     |     |     | 325 |     |     |     |     |     | 330 |     |     |     |     |     | 335 |  |
| Asn | Met | Val | Ser | Glu | Asn | Met | Leu | Cys | Ala | Gly | Ile | Leu | Gly | Asp | Arg |     |  |
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| His | Gly | Thr | Trp | Phe | Leu | Val | Gly | Leu | Val | Ser | Trp | Gly | Glu | Gly | Cys |     |  |
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| Asp | Trp | Ile | His | Gly | His | Ile | Arg | Asp | Lys | Glu | Ala | Pro | Gln | Lys | Ser |     |  |
|     |     |     |     | 405 |     |     |     |     |     | 410 |     |     |     |     |     | 415 |  |
| Trp | Ala | Pro |     |     |     |     |     |     |     |     |     |     |     |     |     |     |  |

1. A method for treating or preventing extravascular fibrin deposition in the airways, in particular the alveolar or bronchoalveolar spaces, in a human subject comprising administering to the subject human activated protein C or a biologically active derivative thereof, via the airway by means of intratracheal, intrabronchial or intraalveolar administration.

2. The method of claim 1, wherein the fibrin deposition is caused by an inflammatory lung disease.

3. The method of claim 2, wherein the inflammatory lung disease is selected from the group consisting of: ALI, ARDS, pneumonia, acute bronchoalveolar diseases, chronic bronchoalveolar diseases, fibrosing alveolitis or bronchial asthma.

4. The method of any of the preceding claims, wherein the inflammatory lung disease is related to a condition selected from the group consisting of: direct or indirect pulmonary trauma, pancreatitis, aspiration pneumonitis, sepsis, severe sepsis and/or septic shock and/or drug induced acute lung insufficiency.

5. The method of any of the preceding claims, wherein the activated protein C is administered by bronchoalveolar lavage with a solution of the activated protein C.

6. The method of any of the preceding claims, wherein the activated protein C is administered by blind tracheal washing with a solution of the activated protein C.

7. The method of any of the preceding claims, wherein the activated protein C is administered by causing the inhalation of a nebulized solution of the activated protein C.

8. The method of any of the preceding claims, wherein the activated protein C is administered by causing the inhalation of the activated protein C in nebulized powder form.

9. The method of any of the preceding claims, wherein the activated protein C is administered by direct application of the activated protein C during bronchoscopy.

10. The method of any of claims 1 through 9 wherein the human is an adult.

11. The method of any of claims 1 through 9 wherein the human is a child.

12. The method of any of the preceding claims, wherein the APC is administered in an amount of from 0.1 µg/kg to about 10 mg/kg body weight per day.

13. The method of any of claims 1 through 9 further comprising administering to the human subject an anticoagulant systemically as an adjuvant therapy.

14. The method of claim 13 wherein the human is an adult.

15. The method of claim 13 wherein the human is a child.

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