

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 May 2003 (15.05.2003)

PCT

(10) International Publication Number
WO 03/039582 A1

- (51) International Patent Classification⁷: **A61K 38/36**, 38/48
- (21) International Application Number: PCT/DK02/00752
- (22) International Filing Date:
8 November 2002 (08.11.2002)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
PA 2001 01667 9 November 2001 (09.11.2001) DK
60/333,572 27 November 2001 (27.11.2001) US
- (71) Applicant (for all designated States except US): **NOVO NORDISK A/S** [DK/DK]; Novo Allé, DK-2880 Bagsvaerd (DK).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): **RØJKJÆR, Rasmus** [DK/DK]; Folkevej 30A, DK-2820 Gentofte (DK).
- (74) Common Representative: **NOVO NORDISK A/S**; Corporate Patents, Novo Allé, DK-2880 Bagsvaerd (DK).
- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).
- Published:**
— with international search report
— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*



WO 03/039582 A1

(54) Title: PHARMACEUTICAL COMPOSITION COMPRISING A FACTOR VII POLYPEPTIDE AND EPSILON-AMINOCAPRONIC ACID

(57) Abstract: The present invention relates to a composition comprising factor VII or a factor VII-related polypeptide and epsilon-aminocaproic acid, and the use thereof for treating bleeding episodes.

PHARMACEUTICAL COMPOSITION COMPRISING A FACTOR VII POLYPEPTIDE AND EPSILON-AMINOCAPRONIC ACID

FIELD OF THIS INVENTION

5 The present invention relates to a pharmaceutical composition comprising a factor VII-related polypeptide and epsilon-aminocaproic acid. The invention also relates to the use of a combination of a factor VII-related polypeptide, and epsilon-aminocaproic acid for the manufacture of a medicament for treatment of subjects suffering from bleeding episodes, or prevention hereof. The invention further relates to use of factor VII in combination with epsilon-aminocaproic acid for the manufacture of a medicament for treatment of non-haemophilic
10 bleeding episodes. The invention also relates to methods of treatment.

BACKGROUND OF THE INVENTION

 Haemostasis is initiated by the formation of a complex between tissue factor (TF) being
15 exposed to the circulating blood following an injury to the vessel wall, and FVIIa which is present in the circulation in an amount corresponding to about 1% of the total FVII protein mass. This complex is anchored to the TF-bearing cell and activates FX into FXa and FIX into FIXa on the cell surface. FXa activates prothrombin to thrombin, which activates FVIII, FV, FXI and FXIII. Furthermore, the limited amount of thrombin formed in this initial step of haemostasis also activates
20 the platelets. Following the action of thrombin on the platelets these change shape and expose charged phospholipids on their surface. This activated platelet surface forms the template for the further FX activation and the full thrombin generation. The further FX activation on the activated platelet surface occurs via a FIXa-FVIIIa complex formed on the surface of the activated platelet, and FXa then converts prothrombin into thrombin while still on the surface. Thrombin
25 then converts fibrinogen into fibrin which is insoluble and which stabilizes the initial platelet plug. This process is compartmentalized, i.e., localised to the site of TF expression or exposure, thereby minimizing the risk of a systemic activation of the coagulation system. The insoluble fibrin forming the plug is furthermore stabilised by FXIII-catalysed cross-linking of the fibrin fibres.

 FVIIa exists in plasma mainly as a single-chain zymogen, which is cleaved by FXa into its
30 two-chain, activated form, FVIIa. Recombinant activated factor VIIa (rFVIIa) has been developed as a pro-haemostatic agent. The administration of rFVIIa offers a rapid and highly effective pro-haemostatic response in haemophilic subjects with bleedings who cannot be treated with coagulation factor products due to antibody formation. Also bleeding subjects with a factor VII deficiency or subjects having a normal coagulation system but experiencing excessive bleeding
35 can be treated successfully with FVIIa. In these studies, no unfavourable side effects of rFVIIa (in particular the occurrence of thromboembolism) has been encountered.

Extra exogenously administered FVIIa increases the formation of thrombin on the activated platelet surface. This occurs in haemophilic subjects lacking FIX or FVIII and therefore missing the most potent pathway for full thrombin formation. Also in the presence of a lowered number of platelets or platelets with a defect function, extra FVIIa increases the thrombin formation.

Commercial preparations of recombinant human FVIIa are sold as NovoSeven® (Novo Nordisk A/S, Denmark). Novoseven® is indicated for treatment of bleeding episodes in haemophilia A and B patients. Novoseven® is the only recombinant FVIIa available on the market for effective and reliable treatment of bleeding episodes.

Epsilon-aminocaproic acid (ϵ -aminocaproic acid, EACA) and its analogue, epsilon-aminocaproic acid (TA) (Trade name in UK "Cyclokapron") are derivatives of the amino acid lysine. Both of these drugs inhibit the proteolytic activity of plasmin and the conversion of plasminogen to plasmin by plasminogen activators. Plasmin cleaves fibrinogen and a series of other proteins involved in coagulation. Epsilon-aminocaproic acid is 6 to 10 times more potent than ϵ -aminocaproic acid.

Epsilon-aminocaproic acid is usually given in tablet form at a typical dose of 3 or 4 grams (in divided doses) daily for an adult. Gastrointestinal upset (nausea, vomiting and diarrhoea) may rarely occur as a side-effect, but these symptoms usually resolve if the dosage is reduced. It may also be given by intravenous injection, but it must be infused slowly as rapid injection may result in dizziness and hypotension. A syrup formulation is also available for paediatric use: the syrup contains 500 mg epsilon-aminocaproic acid in each 5ml, and the usual dose for children is 25 mg/kg up to three times daily. The drug may be of particular use in controlling oral bleeding associated with eruption of teeth. The drug is excreted by the kidneys, and the dose must be reduced if there is renal impairment in order to avoid toxic accumulation.

It is well known that subjects who bleed excessively in association with surgery or major trauma and need blood transfusions develop more complications than those who do not experience any bleeding. However, also moderate bleedings requiring the administration of human blood or blood products (platelets, leukocytes, plasma-derived concentrates for the treatment of coagulation defects, etc.) may lead to complications associated with the risk of transferring human viruses (hepatitis, HIV, parvovirus, and other, by now unknown viruses). Extensive bleedings requiring massive blood transfusions may lead to the development of multiple organ failure including impaired lung and kidney function. Once a subject has developed these serious complications a cascade of events involving a number of cytokines and inflammatory reactions is started making any treatment extremely difficult and unfortunately often unsuccessful. Therefore a major goal in surgery as well as in the treatment of major tissue damage is to avoid or minimise the bleeding. To avoid or minimise such bleeding it is of importance to ensure the formation of sta-

ble and solid haemostatic plugs that are not easily dissolved by fibrinolytic enzymes. Furthermore, it is of importance to ensure quick and effective formation of such plugs or clots.

Today, subjects experiencing bleeding episodes, including trauma victims and subjects bleeding in association with surgery, are often treated with several injections or infusions of FVIIa since the short half-life of FVIIa (2.5 hours) may require more than one administration to maintain a certain level of haemostatic ability. A faster arrest of bleedings would be an important benefit to such subjects. So would a reduction in the number of administrations needed to stop bleeding and maintain haemostasis.

10 European Patent No. 225.160 (Novo Nordisk) concerns compositions of FVIIa and methods for the treatment of bleeding disorders not caused by clotting factor defects or clotting factor inhibitors.

European Patent No. 82.182 (Baxter Travenol Lab.) concerns a composition of factor VIIa for use in counteracting deficiencies of blood clotting factors or the effects of inhibitors to blood clotting factors in a subject.

International Patent Publication No. WO 93/06855 (Novo Nordisk) concerns the topical application of FVIIa.

Hirawat et al., Blood (16 Nov 2000), 96 (11, part 2), p. 84 concerns administration of FVIIa and epsilon-capronic acid to a FVII-deficient patient.

20 Ciavarella et al., Haemostasis 1996:150-154, concerns administration of recombinant factor VIIa (NovoSeven) in the treatment of two patients with Type III von Willebrand's Disease and an inhibitor against von Willebrand Factor.

There is still a need in the art for improved treatment of subjects experiencing bleeding episodes, including subjects where the bleeding episodes are due to surgery, trauma, or other forms of tissue damage; induced coagulopathy, including coagulopathy in multi-transfused subjects; congenital or acquired coagulation or bleeding disorders, including diminished liver function ("liver disease"); defective platelet function or decreased platelet number; lacking or abnormal essential clotting "compounds" (e.g., platelets or von Willebrand factor protein); increased fibrinolysis; anticoagulant therapy or thrombolytic therapy; or stem cell transplantation.

30 There remains a need in the art for an improved, reliable and widely applicable method of enhancing coagulation, enhancing or ensuring formation of stable haemostatic plugs, or enhancing convenience for the treated subject, or achieving full haemostasis in subjects, in particular in subjects having an impaired thrombin generation. There is also a need for methods wherein the time to bleeding arrest is shortened.

SUMMARY OF THE INVENTION

One object of the present invention is to provide compositions, which can effectively be used in the treatment or prophylaxis of bleeding episodes and coagulation disorders.

A second object of the present invention is to provide compositions in single-unit dosage form, which can effectively be used in the treatment or prophylaxis of bleeding episodes or as a procoagulant. Another object of the present invention is to provide compositions,
5 methods of treatment or kits exhibiting a synergistic effect.

A further object of the present invention is to provide compositions, methods of treatment or kits exhibiting no substantial side effects, such as a high level of systemic activation of the coagulation system.

10 Other objects of the present invention will become apparent upon reading the present description.

In a first aspect the invention provides a pharmaceutical composition comprising a factor VII-related polypeptide and epsilon-aminocaproic acid.

15 In a second aspect, the invention provides the use of a factor VII-related polypeptide in combination with a epsilon-aminocaproic acid for the manufacture of a medicament for treating bleeding episodes in a subject.

In different embodiments thereof, the medicaments are for reducing time needed to obtain full haemostasis, reducing time needed to maintain haemostasis, reducing clotting time, prolonging the clot lysis time, and increasing clot strength.

20 In different embodiments, the medicaments are for treatment of subjects experiencing bleeding episodes due to surgery, trauma, or other forms of tissue damage; coagulopathy, including coagulopathy in multi-transfused subjects; congenital or acquired coagulation or bleeding disorders, including decreased liver function ("liver disease"); defective platelet function or decreased platelet number; lacking or abnormal essential clotting "compounds" (e.g., platelets or von Willebrand factor protein); increased fibrinolysis; anticoagulant therapy or thrombolytic therapy; stem cell transplantation. In one series of embodiments, the bleedings occur in organs
25 such as the brain, inner ear region, eyes, liver, lung, tumour tissue, gastrointestinal tract; in another series of embodiments, it is diffuse bleeding, such as in haemorrhagic gastritis and profuse uterine bleeding. In another series of embodiments, the bleeding episodes are bleeding in connection with surgery or trauma in subjects having acute haemarthroses (bleedings in joints),
30 chronic haemophilic arthropathy, haematomas, (e.g., muscular, retroperitoneal, sublingual and retropharyngeal), bleedings in other tissue, haematuria (bleeding from the renal tract), cerebral haemorrhage, surgery (e.g., hepatectomy), dental extraction, and gastrointestinal bleedings (e.g., UGI bleeds). In one embodiment, the medicament is for treating bleeding episodes due to
35 trauma, or surgery, or lowered count or activity of platelets, in a subject.

In a further aspect, the invention provides a method for treating bleeding episodes in a subject, the method comprising administering to a subject in need thereof a first amount of a

preparation of a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amount together are effective to treat bleedings.

In a further aspect, the invention provides a method for reducing clotting time in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to reduce clotting time.

In a further aspect, the invention provides a method to enhance haemostasis in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to enhance haemostasis.

In a further aspect, the invention provides a method for prolonging the clot lysis time in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to prolong the clot lysis time.

In a further aspect, the invention provides a method for increasing clot strength in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to increase clot strength.

In a further aspect, the invention provides the use of factor VII in combination with epsilon-aminocaproic acid for the manufacture of a medicament for treating bleeding episodes in non-haemophilic subjects.

In different embodiments thereof, the medicaments are for reducing time needed to obtain full haemostasis, reducing time needed to maintain haemostasis, reducing clotting time, prolonging the clot lysis time, and increasing clot strength.

In different embodiments, the medicaments are for treatment of non-haemophilic subjects experiencing bleeding episodes due to surgery, trauma, or other forms of tissue damage; coagulopathy, including coagulopathy in multi-transfused subjects; defective platelet function or decreased platelet number; increased fibrinolysis; anticoagulant therapy or thrombolytic therapy; stem cell transplantation. In one series of embodiments, the bleedings occur in organs such as the brain, inner ear region, eyes, liver, lung, tumour tissue, gastrointestinal tract; in another series of embodiments, it is diffuse bleeding, such as in haemorrhagic gastritis and profuse uterine bleeding. In one embodiment, the medicament is for treating bleeding episodes due to co-

agulopathy or lowered count or activity of platelets; in another, the bleeding episodes are due to anticoagulant treatment.

In one aspect, the invention provides a method for treating bleeding episodes in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amount together are effective to treat bleedings.

In one aspect, the invention provides a method for reducing clotting time in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to reduce clotting time.

In one aspect, the invention provides a method to enhance haemostasis in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to enhance haemostasis.

In one aspect, the invention provides a method for prolonging the clot lysis time in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to prolong the clot lysis time.

In one aspect, the invention provides a method for increasing clot strength in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to increase clot strength.

In one aspect, the invention provides a kit containing a treatment for non-haemophilic bleeding episodes comprising (a) An effective amount of factor VII and an effective amount of epsilon-aminocaproic acid and a pharmaceutically acceptable carrier in a single-unit dosage form; and (b) Container means for containing said single-unit dosage form.

In one series of embodiments of the methods, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered in single-unit dosage form.

In another series of embodiments the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered in the form of a first-unit dosage form comprising a preparation of factor VII or a factor VII-related polypeptide and a second-unit dosage form comprising a preparation of epsilon-aminocaproic acid. In a series of embodiments thereof, the

first-unit dosage form and the second-unit dosage form are administered with a time separation of no more than 15 minutes.

In a further aspect, the invention provides a kit containing a treatment for bleeding episodes comprising (a) An effective amount of a factor VII-related polypeptide, and an effective amount of epsilon-aminocaproic acid, and a pharmaceutically acceptable carrier in a single-unit dosage form; and (a) Container means for containing said single-unit dosage form.

In one series of embodiments of the invention the factor VII-related polypeptide is a factor VII amino acid sequence variant. In one embodiment the ratio between the activity of the factor VII-related polypeptide and the activity of native human factor VIIa (wild-type FVIIa) is at least about 1.25 when tested in the "In Vitro Hydrolysis Assay" as described in the present description.

In one embodiment factor VII is human factor VII. In one embodiment the factor VII is bovine, porcine, canine, equine, murine or salmon factor VII. In another embodiment the factor VII is recombinantly made. In another embodiment the factor VII is derived from plasma. In a preferred embodiment the factor VII is recombinant human factor VII. In one series of embodiments of the invention the factor VII or factor VII-related polypeptide is in its activated form. In one preferred embodiment of the invention the factor VII is recombinant human factor VIIa.

In one embodiment the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are present in a ratio by mass of between about 100:1 and about 1:100 (w/w factor VII:epsilon-aminocaproic acid).

In one embodiment, the factor VII-related polypeptides are amino acid sequence variants having no more than 20 amino acids replaced, deleted or inserted compared to wild-type factor VII (i.e., a polypeptide having the amino acid sequence disclosed in U.S. Patent No. 4,784,950), In another embodiment, the factor VII variants have no more than 15 amino acids replaced, deleted or inserted; in other embodiments, the factor VII variants have no more than 10 amino acids, such as 8, 6, 5, or 3 amino acids, replaced, deleted or inserted compared to wild-type factor VII. In one embodiment, the factor VII variants are selected from the list of L305V-FVIIa, L305V/M306D/D309S-FVIIa, L305I-FVIIa, L305T-FVIIa, F374P-FVIIa, V158T/M298Q-FVIIa, V158D/E296V/M298Q-FVIIa, K337A-FVIIa, M298Q-FVIIa, V158D/M298Q-FVIIa, L305V/K337A-FVIIa, V158D/E296V/M298Q/L305V-FVIIa, V158D/E296V/M298Q/K337A-FVIIa, V158D/E296V/M298Q/L305V/K337A-FVIIa, K157A-FVII, E296V-FVII, E296V/M298Q-FVII, V158D/E296V-FVII, V158D/M298K-FVII, and S336G-FVII

In a further embodiment, the factor VII-related polypeptides have increased tissue factor-independent activity compared to native human coagulation factor VIIa. In another embodiment, the increased activity is not accompanied by changes in the substrate specificity. In

another embodiment of the invention, the binding of the factor VII-related polypeptides to tissue factor are not impaired and the factor VII-related polypeptides have at least the activity of wild-type factor VIIa when bound to tissue factor.

In one embodiment, the clotting time is reduced in mammalian blood. In another
5 embodiment the haemostasis is enhanced in mammalian blood. In another embodiment the clot lysis time is prolonged in mammalian blood. In another embodiment the clot strength is increased in mammalian blood. In one embodiment, the mammalian blood is human blood. In another embodiment, the mammalian blood is normal human blood; in one embodiment, the blood is blood from a subject having an impaired thrombin generation. In one embodiment, the
10 blood is blood from a subject having a deficiency of one or more coagulation factors; in another embodiment, the blood is blood from a subject having inhibitors against one or more coagulation factors; in one embodiment, the blood is from a subject having a lowered concentration of fibrinogen. In one series of embodiments, the blood is plasma.

In one embodiment of the invention, the factor VII or factor VII-related polypeptide and
15 the epsilon-aminocaproic acid are the sole haemostatic agents contained in the composition. In another embodiment, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid polypeptide are the sole active haemostatic agents contained in the composition. In another embodiment, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are the sole coagulation factors administered to the subject. In one embodi-
20 ment of the invention, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are the sole active agents administered to the patient. In one embodiment, the composition is substantially free of thrombin or prothrombin; in another embodiment, the composition is substantially free of FX; in another embodiment, the composition is substantially free of FXa.

25 In another embodiment, the pharmaceutical composition is formulated for intravenous administration, preferably injection or infusion, in particular injection. In one embodiment, the composition contains at least one pharmaceutical acceptable excipients or carrier.

In one embodiment of the invention, the composition is in single-unit dosage form wherein the single-unit dosage form contains both compounds. In one embodiment of the
30 invention, the composition is in the form of a kit-of-parts comprising a preparation of factor VII or a factor VII-related polypeptide as a first-unit dosage form and a preparation of epsilon-aminocaproic acid as a second-unit dosage form, and comprising container means for containing said first and second unit dosage forms. In one embodiment the composition or kit, as applicable, further contains directions for the administration of the composition or separate
35 components, respectively.

In one embodiment of the invention, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered in single-dosage form. In one embodiment

of the invention, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered in the form of a first-unit dosage form comprising a preparation of factor VII or a factor VII-related polypeptide and a second-unit dosage form comprising a preparation of epsilon-aminocaproic acid.

5 In one embodiment of the invention, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered simultaneously. In another embodiment, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered sequentially. In one embodiment, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered with a time separation of no more than 15
10 minutes, preferably 10, more preferred 5, more preferred 2 minutes. In one embodiment, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are administered with a time separation of more than 15 minutes, preferably up to 2 hours, more preferred from 1 to 2 hours, more preferred up to 1 hour, more preferred from 30 minutes to 1 hour, more preferred up to 30 minutes, more preferred from 15 to 30 minutes.

15 In one embodiment, the amount of the factor VII or factor VII-related polypeptide is an amount from about 0.05 mg/day to about 500 mg/day (70-kg subject). In one embodiment, the amount of epsilon-aminocaproic acid ranges from about 0.05 mg to about 6000 mg/day, e.g., from about 1 mg to about 5000 mg/day, or, e.g., from about 5 mg to about 5000 mg/day (70-kg subject).

20 In one embodiment the factor VII or factor VII-related polypeptide and epsilon-aminocaproic acid are present in a ratio by mass of between about 100:1 and about 1:100 (w/w factor VII:epsilon-aminocaproic acid)

In one embodiment of the present invention, the pharmaceutical composition is in single-unit dosage form and consists essentially of a preparation of factor VII or a factor VII-
25 related polypeptide, and a preparation of epsilon-aminocaproic acid, and one or more of the components selected from the list of pharmaceutical acceptable carriers, stabilizers, detergents, neutral salts, antioxidants, preservatives, and protease inhibitors.

In another embodiment of the present invention, the pharmaceutical composition is in the form of a kit-of-parts with the first-unit dosage form consisting essentially of a preparation
30 of factor VII or a factor VII-related polypeptide, and one or more of the components selected from the list of pharmaceutical acceptable carriers, stabilizers, detergents, neutral salts, antioxidants, preservatives, and protease inhibitors; and with the second-unit dosage form consisting essentially of a preparation of epsilon-aminocaproic acid and one or more of the components selected from the list of pharmaceutical acceptable carriers, stabilizers, detergents,
35 neutral salts, antioxidants, preservatives, and protease inhibitors.

In a further embodiment, the subject is a human; in another embodiment, the subject has an impaired thrombin generation; in one embodiment, the subject has a lowered plasma

concentration of fibrinogen (e.g., a multi-transfused subject); in one embodiment, the subject has a lowered plasma concentration of factor VIII or FIX; in one embodiment, the subject has coagulopathy; in one embodiment, the subject has a lowered number or activity of platelets..

5 In another aspect, the invention concerns a method to enhance haemostasis in a subject suffering from a factor VII responsive syndrome compared to when the subject is treated with factor VII as the only coagulation protein, the method comprising administering to the subject in need thereof a first amount of a preparation of factor VII or a factor VII-related polypeptide, and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective to enhance haemostasis.

10 In another aspect, the invention concerns a method to enhance formation of thrombin in a subject, the method comprising administering to the subject in need thereof a first amount of a preparation of factor VII or a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective to enhance formation of thrombin.

15 In another aspect, the invention concerns a method to enhance formation of thrombin in a subject suffering from a factor VII responsive syndrome compared to when the subject is treated with factor VII as the only coagulation protein, the method comprising administering to the subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective to enhance formation of thrombin.

20 In another aspect, the invention concerns a method for reducing the number of administrations of coagulation factor protein needed to accomplish haemostasis in a subject suffering non-haemophilic bleeding episode compared to the number of administrations needed when factor VII is administered to the subject as the only coagulation factor protein, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective to reduce the number of administrations of coagulation factor protein.

25 In another aspect, the invention concerns a method for reducing the number of administrations of coagulation factor protein needed to accomplish haemostasis in a subject suffering from a factor VII responsive syndrome compared to the number of administrations needed when factor VII is administered to the subject as the only coagulation factor protein, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective to reduce the number of administrations of coagulation factor protein.

30
35

In another aspect, the invention concerns a method of treating bleedings in a subject suffering from a factor VII responsive syndrome, the method comprising administering to the subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective in treating bleedings.

In another aspect, the invention concerns a method of enhancing and maintaining haemostasis in a subject suffering from a non-haemophilic bleeding episode, the method comprising administering to the subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective in treating bleedings

In another aspect, the invention concerns a method of enhancing and maintaining haemostasis in a subject suffering from a factor VII responsive syndrome, the method comprising administering to the subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective in treating bleedings.

In one embodiment, the factor VII is human recombinant factor VIIa (rFVIIa). In another embodiment, the rFVIIa is NovoSeven® (Novo Nordisk A/S, Bagsvaerd, Denmark).

In another aspect, the invention relates to the use of factor VII or a factor VII-related polypeptide in combination with a epsilon-aminocaproic acid for the manufacture of a medicament for enhancing fibrin clot formation in mammalian plasma.

In another aspect, the invention relates to a method of enhancing fibrin clot formation in a subject, which method comprises administering to a subject in need thereof a first amount of a preparation of factor VII or a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amounts together are effective in treating bleedings.

LIST OF FIGURES

Figure 1: Addition of FVIIa results in a dose-dependent prolongation of the clot lysis time. This effect was optimal at 10 nM FVIIa.

Figure 2: In the presence of 10 nM FVIIa, addition of epsilon-Aminocaproic Acid resulted in a further prolongation of the clot lysis time. The effect was dose-dependent and optimal at 1 μ M epsilon-Aminocaproic Acid.

DETAILED DESCRIPTION OF THIS INVENTION

Subjects, who bleed excessively in association with surgery or major trauma thus needing blood transfusions, develop more complications than those who do not experience any bleeding. However, also moderate bleedings may lead to complications if they require the ad-

ministration of human blood or blood products (platelets, leukocytes, plasma-derived concentrates for the treatment of coagulation defects, etc.) because this is associated with the risk of transferring human viruses (e.g., hepatitis, HIV, parvovirus, or other, by now unknown viruses) as well as non-viral pathogens. Extensive bleedings requiring massive blood transfusions may lead to the development of multiple organ failure including impaired lung and kidney function. Once a subject has developed these serious complications a cascade of events involving a number of cytokines and inflammatory reactions is started making any treatment extremely difficult and unfortunately often unsuccessful. A patient experiencing a major loss of blood becomes clinically unstable. Such patients are in risk of experiencing atrial fibrillation, which may lead to a fatal stop of cardiac activity; impaired renal function; or fluid extravasations in lungs (so-called "wet lungs" or ARDS). Therefore, a major goal in surgery as well as in the treatment of major tissue damage is to avoid or minimise the bleeding. To avoid or minimize such unwanted bleedings it is important to ensure formation of stable and solid haemostatic plugs that are not readily dissolved by fibrinolytic enzymes. Furthermore, it is of importance to ensure quick and effective formation of such plugs or clots.

Subjects with thrombocytopenia (lowered count or activity of platelets) also have an impaired thrombin generation as well as a defective stabilization of the fibrin plugs resulting in haemostatic plugs prone to premature dissolution. Furthermore, subjects subjected to major trauma or organ damage and who, as a consequence, have obtained frequent blood transfusions often have lowered platelet counts as well as lowered levels of fibrinogen, factor VIII, and other coagulation proteins. These subjects experience an impaired (or lowered) thrombin generation. These subjects, therefore, have a defective, or less efficient, haemostasis leading to the formation of fibrin plugs that are easily and prematurely dissolved by proteolytic enzymes, such enzymes in addition being extensively released in situations characterized by extensive trauma and organ damage.

Bleedings in tissues may also lead to the formation of haematomas. The sizes of (in particular intercranial and spinal) haematomas are closely correlated to the extent of loss of neurological function, rehabilitation difficulties, and/or the severity and degree of permanent impairments of neurological function following rehabilitation. The most severe consequences of haematomas are seen when they are located in the brain where they may even lead to the death of the patient.

Thus, major objectives in treatment of bleedings are to obtain haemostasis in a minimum of time, thus keeping the blood loss at a minimum.

The present invention thus provides beneficial compositions, uses and methods of treatment for treatment of bleeding episodes in subjects in need of such treatment. The compositions, uses and methods may be associated with beneficial effects such as less blood loss before haemostasis is obtained, less blood needed during surgery, blood pressure kept at an acceptable

level until haemostasis is obtained, faster stabilisation of blood pressure, shorter recovery time for the treated patient, shorter rehabilitation time for the treated patient, diminished formation of haematomas or formation of smaller haematomas, including haematomas in the brain, faster arrest of bleedings, reduction in the number of administrations needed to stop bleeding and
5 maintain haemostasis.

The administration of a preparation of factor VII or a factor VII-related polypeptide, e.g., factor VIIa, in combination with a preparation of epsilon-aminocaproic acid provides a shortened clotting time, a firmer clot and an increased resistance to fibrinolysis compared to the clotting time, clot firmness and resistance when either factor VIIa or epsilon-aminocaproic acid
10 is administered alone.

The administration of a preparation of factor VII or a factor VII-related polypeptide, e.g., factor VIIa, in combination with a preparation of epsilon-aminocaproic acid also provides for a reduced time to obtain bleeding arrest and a reduced number of administrations to maintain haemostasis compared to the situation when either factor VIIa or epsilon-aminocaproic
15 acid is administered alone. The present invention provides a beneficial effect of simultaneous or sequential dosing of a preparation of epsilon-aminocaproic acid and a preparation of factor VII or a factor VII-related polypeptide. The pharmaceutical composition according to the present invention may be in the form of a single composition or it may be in the form of a multi-
20 component kit (kit-of-parts). The composition according to the present invention is useful as a therapeutic and prophylactic procoagulant in mammals, including primates such as humans. The present invention further provides a method for treating (including prophylactically treating or preventing) bleeding episodes in a subject, including a human being.

Whenever, a first or second or third, etc., unit dose is mentioned throughout this specification this does not indicate the preferred order of administration, but is merely done for
25 convenience purposes.

A combination of a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid is an advantageous product ensuring short clotting times, rapid formation of haemostatic plugs, and formation of stable haemostatic plugs. It has been found by the present inventor that a combination of factor VII or a factor VII-related
30 polypeptide and a epsilon-aminocaproic acid is an advantageous product ensuring the formation of solid, stable and quickly formed haemostatic plugs.

The present inventors have shown that a combination of factor VIIa and epsilon-aminocaproic acid can increase the firmness of the clot more effectively than either factor VIIa or epsilon-aminocaproic acid alone. It has also been shown that combination of factor VII or a
35 factor VII-related polypeptide and a epsilon-aminocaproic acid can prolong the in vitro clot lysis time in normal human plasma more effectively than either factor VIIa or epsilon-aminocaproic acid alone. It has also been shown that combination of factor VII or a factor VII-related

polypeptide and a epsilon-aminocaproic acid can prolong the half-clot lysis time in normal human plasma more effectively than either factor VIIa or epsilon-aminocaproic acid alone. It has also been shown that combination of factor VII or a factor VII-related polypeptide and a epsilon-aminocaproic acid can protect the clot from fibrinolysis, in particular tPA-mediated fibrinolysis, in normal human plasma more effectively than either factor VIIa or epsilon-aminocaproic acid alone. Thus, by enhancing coagulation a more effective treatment of bleeding in subjects can be obtained.

Without wishing to be bound by theory, it is believed that the full thrombin generation is necessary for a solid, stabile haemostatic plug to be formed, and thereby for the maintenance of haemostasis. The fibrin structure of such a plug is dependent on both the amount of thrombin formed and the rate of the initial thrombin generation. In the presence of an impaired thrombin generation a porous fibrin plug, which is highly permeable, is being formed. The fibrinolytic enzymes normally present on the fibrin surface easily dissolve such a fibrin plug. The formation of a stable fibrin plug is also dependent on the presence of factor XIIIa, which is being activated by thrombin and therefore also dependent on the full thrombin generation. Furthermore, the recently described thrombin activatable fibrinolytic inhibitor, TAFI, requires rather high thrombin amounts for its activation. In the presence of a not fully adequate thrombin formation the TAFI may therefore not be activated resulting in the formation of a haemostatic plug, which is easier than normally dissolved by the normal fibrinolytic activity. In situations with lowered number of platelets, thrombocytopenia, a faster thrombin generation is initiated by the administration of exogenous extra factor VIIa. However, the total thrombin generation is not normalised by factor VIIa even in high concentrations.

In subjects with lowered plasma concentrations of fibrinogen (multi-transfused subjects as a consequence of multiple trauma or extensive surgery) full thrombin activation does not occur. A more effective haemostasis is then obtained by the administration of a combination of factor VII or a factor VII-related polypeptide, and a epsilon-aminocaproic acid.

Subjects with thrombocytopenia have an impaired thrombin generation as well as a defective stabilization of the fibrin plugs resulting in haemostatic plugs prone to premature dissolution. Furthermore, subjects subjected to major trauma or organ damage and who, as a consequence, have obtained frequent blood transfusions often have lowered platelet counts as well as lowered levels of fibrinogen, factor VIII, and other coagulation proteins. These subjects experience an impaired (or lowered) thrombin generation. In addition, their lowered fibrinogen level interfere negatively with the activation of factor XIII. These subjects, therefore, have a defective, or less efficient, haemostasis leading to the formation of fibrin plugs which are easily and prematurely dissolved by proteolytic enzymes, such enzymes in addition being extensively released in situations characterized by extensive trauma and organ damage.

In order to facilitate the formation of fully stabilized plugs with full capacity to maintain haemostasis in a subject, a composition according to the invention is administered. This composition is especially beneficial in subjects with a lowered number of platelets and in subjects with lowered plasma levels of fibrinogen and/or other coagulation proteins.

5

Factor VII Polypeptides:

In practicing the present invention, any factor VII polypeptide may be used that is effective in preventing or treating bleeding. This includes factor VII polypeptides derived from blood or plasma, or produced by recombinant means.

10

The present invention encompasses factor VII polypeptides, such as, e.g., those having the amino acid sequence disclosed in U.S. Patent No. 4,784,950 (wild-type human factor VII). In some embodiments, the factor VII polypeptide is human factor VIIa, as disclosed, e.g., in U.S. Patent No. 4,784,950 (wild-type factor VII). In one series of embodiments, factor VII polypeptides include polypeptides that exhibit at least about 10%, preferably at least about 30%, more preferably at least about 50%, and most preferably at least about 70%, of the specific biological activity of human factor VIIa. In one series of embodiments, factor VII polypeptides include polypeptides that exhibit at least about 90%, preferably at least about 100%, preferably at least about 120%, more preferably at least about 140%, and most preferably at least about 160%, of the specific biological activity of human factor VIIa. In one series of embodiments, factor VII polypeptides include polypeptides that exhibit at least about 70 %, preferably at least about 80 %, more preferably at least about 90 %, and most preferable at least about 95 %, of identity with the sequence of wild-type factor VII as disclosed in U.S. Patent No. 4,784,950.

15

20

25

30

35

As used herein, "factor VII polypeptide" encompasses, without limitation, factor VII, as well as factor VII-related polypeptides. The term "factor VII" is intended to encompass, without limitation, polypeptides having the amino acid sequence 1-406 of wild-type human factor VII (as disclosed in U.S. Patent No. 4,784,950), as well as wild-type factor VII derived from other species, such as, e.g., bovine, porcine, canine, murine, and salmon factor VII, said factor VII derived from blood or plasma, or produced by recombinant means. It further encompasses natural allelic variations of factor VII that may exist and occur from one individual to another. Also, degree and location of glycosylation or other post-translation modifications may vary depending on the chosen host cells and the nature of the host cellular environment. The term "factor VII" is also intended to encompass factor VII polypeptides in their uncleaved (zymogen) form, as well as those that have been proteolytically processed to yield their respective bioactive forms, which may be designated factor VIIa. Typically, factor VII is cleaved between residues 152 and 153 to yield factor VIIa.

"Factor VII-related polypeptides" include, without limitation, factor VII polypeptides that have either been chemically modified relative to human factor VII and/or contain one or

more amino acid sequence alterations relative to human factor VII (i.e., factor VII variants), and/or contain truncated amino acid sequences relative to human factor VII (i.e., factor VII fragments). Such factor VII-related polypeptides may exhibit different properties relative to human factor VII, including stability, phospholipid binding, altered specific activity, and the like.

5 The term "factor VII-related polypeptides" are intended to encompass such polypeptides in their uncleaved (zymogen) form, as well as those that have been proteolytically processed to yield their respective bioactive forms, which may be designated "factor VIIa-related polypeptides" or "activated factor VII-related polypeptides"

As used herein, "factor VII-related polypeptides" encompasses, without limitation, polypeptides exhibiting substantially the same or improved biological activity relative to wild-type human factor VII, as well as polypeptides in which the factor VIIa biological activity has been substantially modified or reduced relative to the activity of wild-type human factor VIIa. These polypeptides include, without limitation, factor VII or factor VIIa that has been chemically modified and factor VII variants into which specific amino acid sequence alterations have been introduced that modify or disrupt the bioactivity of the polypeptide.

10 It further encompasses polypeptides with a slightly modified amino acid sequence, for instance, polypeptides having a modified N-terminal end including N-terminal amino acid deletions or additions, and/or polypeptides that have been chemically modified relative to human factor VIIa.

20 Factor VII-related polypeptides, including variants of factor VII, whether exhibiting substantially the same or better bioactivity than wild-type factor VII, or, alternatively, exhibiting substantially modified or reduced bioactivity relative to wild-type factor VII, include, without limitation, polypeptides having an amino acid sequence that differs from the sequence of wild-type factor VII by insertion, deletion, or substitution of one or more amino acids.

25 Factor VII-related polypeptides, including variants, encompass those that exhibit at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 100%, at least about 110%, at least about 120%, or at least about 130%, of the specific activity of wild-type factor VIIa that has been produced in the same cell type, when tested in one or more of a clotting assay, proteolysis assay, or TF binding assay as described above.

30 Factor VII-related polypeptides, including variants, having substantially the same or improved biological activity relative to wild-type factor VIIa encompass those that exhibit at least about 25%, preferably at least about 50%, more preferably at least about 75%, more preferably at least about 100%, more preferably at least about 110%, more preferably at least about 120%, and most preferably at least about 130% of the specific activity of wild-type factor VIIa that has

35

been produced in the same cell type, when tested in one or more of a clotting assay, proteolysis assay, or TF binding assay as described above.

Factor VII-related polypeptides, including variants, having substantially reduced biological activity relative to wild-type factor VIIa are those that exhibit less than about 25%, preferably less than about 10%, more preferably less than about 5% and most preferably less than about 1% of the specific activity of wild-type factor VIIa that has been produced in the same cell type when tested in one or more of a clotting assay, proteolysis assay, or TF binding assay as described above. factor VII variants having a substantially modified biological activity relative to wild-type factor VII include, without limitation, factor VII variants that exhibit TF-independent factor X proteolytic activity and those that bind TF but do not cleave factor X.

In some embodiments the factor VII polypeptides are factor VII-related polypeptides, in particular variants, wherein the ratio between the activity of said factor VII polypeptide and the activity of native human factor VIIa (wild-type FVIIa) is at least about 1.25 when tested in the "In Vitro Hydrolysis Assay" (see "Assays", below); in other embodiments, the ratio is at least about 2.0; in further embodiments, the ratio is at least about 4.0. In some embodiments of the invention, the factor VII polypeptides are factor VII-related polypeptides, in particular variants, wherein the ratio between the activity of said factor VII polypeptide and the activity of native human factor VIIa (wild-type FVIIa) is at least about 1.25 when tested in the "In Vitro Proteolysis Assay" (see "Assays", below); in other embodiments, the ratio is at least about 2.0; in further embodiments, the ratio is at least about 4.0; in further embodiments, the ratio is at least about 8.0.

In some embodiments, the factor VII polypeptide is human factor VII, as disclosed, e.g., in U.S. Patent No. 4,784,950 (wild-type factor VII). In some embodiments, the factor VII polypeptide is human factor VIIa. In one series of embodiments, the factor VII polypeptides are factor VII-related polypeptides that exhibits at least about 10%, preferably at least about 30%, more preferably at least about 50%, and most preferably at least about 70%, of the specific biological activity of human factor VIIa. In some embodiments, the factor VII polypeptides have an amino acid sequence that differs from the sequence of wild-type factor VII by insertion, deletion, or substitution of one or more amino acids.

Non-limiting examples of factor VII variants having substantially the same or better biological activity compared to wild-type factor VIIa include, but are not limited to, those described in Danish Patent Applications Nos. PA 2000 00734 and PA 2000 01360 (corresponding to WO 01/83725), and PA 2000 01361 (corresponding to WO 02/22776). Non-limiting examples of factor VII variants having substantially the same or improved biological activity as wild-type factor VII include S52A-FVII, S60A-FVII (Iino et al., Arch. Biochem. Biophys. 352: 182-192, 1998); L305V-FVII, L305V/M306D/D309S-FVII, L305I-FVII, L305T-FVII, F374P-FVII, V158T/M298Q-FVII, V158D/E296V/M298Q-FVII, K337A-FVII, M298Q-FVII, V158D/M298Q-FVII, L305V/K337A-FVII,

V158D/E296V/M298Q/L305V-FVII, V158D/E296V/M298Q/K337A-FVII, V158D/E296V/M298Q/L305V/K337A-FVII, K157A-FVII, E296V-FVII, E296V/M298Q-FVII, V158D/E296V-FVII, V158D/M298K-FVII, and S336G-FVII; FVIIa variants exhibiting increased proteolytic stability as disclosed in U.S. Patent No. 5,580,560; factor VIIa that has been
5 proteolytically cleaved between residues 290 and 291 or between residues 315 and 316 (Mollerup et al., *Biotechnol. Bioeng.* 48:501-505, 1995); and oxidized forms of factor VIIa (Kornfelt et al., *Arch. Biochem. Biophys.* 363:43-54, 1999). Non-limiting examples of factor VII variants having substantially reduced or modified biological activity relative to wild-type factor VII include R152E-FVIIa (Wildgoose et al., *Biochem* 29:3413-3420, 1990), S344A-FVIIa (Kazama et al., *J. Biol. Chem.* 270:66-72, 1995), FFR-FVIIa (Holst et al., *Eur. J. Vasc. Endovasc. Surg.* 15:515-520,
10 1998), and factor VIIa lacking the Gla domain, (Nicolaisen et al., *FEBS Letts.* 317:245-249, 1993). Non-limiting examples of chemically modified factor VII polypeptides and sequence variants are described, e.g., in U.S. Patent No. 5,997,864.

The biological activity of factor VIIa in blood clotting derives from its ability to (i) bind
15 to tissue factor (TF) and (ii) catalyze the proteolytic cleavage of factor IX or factor X to produce activated factor IX or X (factor IXa or Xa, respectively).

For purposes of the invention, biological activity of factor VII polypeptides ("factor VII biological activity") may be quantified by measuring the ability of a preparation to promote blood clotting using factor VII-deficient plasma and thromboplastin, as described, e.g., in U.S.
20 Patent No. 5,997,864. In this assay, biological activity is expressed as the reduction in clotting time relative to a control sample and is converted to "factor VII units" by comparison with a pooled human serum standard containing 1 unit/ml factor VII activity. Alternatively, factor VIIa biological activity may be quantified by

- (i) Measuring the ability of factor VIIa or a factor VIIa -related polypeptide to produce
25 activated factor X (factor Xa) in a system comprising TF embedded in a lipid membrane and factor X. (Persson et al., *J. Biol. Chem.* 272:19919-19924, 1997);
- (ii) Measuring factor X hydrolysis in an aqueous system ("In Vitro Proteolysis Assay", see below);
- (iii) Measuring the physical binding of factor VIIa or a factor VIIa -related polypeptide to TF
30 using an instrument based on surface plasmon resonance (Persson, *FEBS Letts.* 413:359-363, 1997); and
- (iv) Measuring hydrolysis of a synthetic substrate by factor VIIa and/or a factor VIIa -related polypeptide ("In Vitro Hydrolysis Assay", see below); and
- (v) Measuring generation of thrombin in a TF-independent in vitro system.

The term "factor VII biological activity" or "factor VII activity" is intended to include the ability to generate thrombin; the term also includes the ability to generate thrombin on the surface of activated platelets in the absence of tissue factor.

5 A factor VIIa preparation that may be used according to the invention is, without limitation, NovoSeven® (Novo Nordisk A/S, Bagsvaerd, Denmark).

Epsilon-aminocaproic acid:

Epsilon-aminocaproic acid is described by, e.g., Vander Salm TJ, et al; J Thorac Cardiovasc Surg, 1996 Oct; Penta de Peppo A, et al; Tex Heart Inst J, 1995; and Leipzig TJ, et al; J Neurosurg, 1997
10 Feb

In the present context the three-letter or one-letter indications of the amino acids have been used in their conventional meaning as indicated in table 1. Unless indicated explicitly, the amino acids mentioned herein are L-amino acids. It is to be understood, that the first letter in,
15 for example, K337 represent the amino acid naturally present at the indicated position wild-type factor VII, and that, for example, [K337A]-FVIIa designates the FVII-variant wherein the amino acid represented by the one-letter code K naturally present in the indicated position is replaced by the amino acid represented by the one-letter code A.

Table 1: Abbreviations for amino acids:

Amino acid	Tree-letter code	One-letter code
Glycine	Gly	G
Proline	Pro	P
Alanine	Ala	A
Valine	Val	V
Leucine	Leu	L
Isoleucine	Ile	I
Methionine	Met	M
Cysteine	Cys	C
Phenylalanine	Phe	F
Tyrosine	Tyr	Y
Tryptophan	Trp	W
Histidine	His	H
Lysine	Lys	K
Arginine	Arg	R
Glutamine	Gln	Q
Asparagine	Asn	N
Glutamic Acid	Glu	E
Aspartic Acid	Asp	D

20

The term "factor VIIa" or "FVIIa" may be used interchangeably.

In this context, "subjects with an impaired thrombin generation" means subjects who cannot generate a full thrombin burst on the activated platelet surface and includes subjects having
25 a generation of thrombin less that the thrombin-generation in subjects having a fully functioning,

normal haemostatic system, including a normal amount and function of coagulation factors, platelets and fibrinogen (e.g., as in pooled, normal human plasma), and includes, without limitations, subjects lacking factor VIII; subjects having a lowered number of platelets or platelets with a defective function (e.g., thrombocytopenia or thrombasthenia Glanzmann or subjects with excessive bleeds); subjects having lowered levels of prothrombin, FX or FVII; subjects having a lowered level of several coagulation factors (e.g., due to excessive bleeding as a consequence of trauma or extensive surgery); and subjects with lowered plasma concentrations of fibrinogen (e.g., multitransfused subjects).

By "level of thrombin generation" or "normal thrombin generation" is meant the level of the patient's level of thrombin generation compared to the level in healthy subjects. The level is designated as a percentage of the normal level. The terms may, where appropriate, be used interchangeably.

The term "enhancement of the haemostatic system" means an enhancement of the ability to generate thrombin. The term "enhancing haemostasis" is intended to encompass the situations when the measured thrombin generation for a test sample containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid is prolonged relative to the individual thrombin generation of a control sample containing only the factor VII or factor VII-related polypeptide or the epsilon-aminocaproic acid, respectively, when tested in the same thrombin generation assay. The thrombin generation may be assayed as described in the thrombin generation assay of the present description (see "assay part").

"Sole" agents or factors as used herein refers to situations in which the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid, taken together, are the only haemostatic agents, or active haemostatic agents, or coagulation factors contained in the pharmaceutical composition or kit, or are the only haemostatic agents, or active haemostatic agents, or coagulation factors administered to the patient in the course of a particular treatment, such as, e.g., in the course of a particular bleeding episode. It will be understood that these situations encompass those in which other haemostatic agents or coagulation factors, as applicable, are not present in either sufficient quantity or activity so as to significantly influence one or more coagulation parameters.

Clot lysis time, clot strength, fibrin clot formation, and clotting time are clinical parameters used for assaying the status of patient's haemostatic system. Blood samples are drawn from the patient at suitable intervals and one or more of the parameters are assayed by means of, e.g., thromboelastography as described by, e.g., Meh et al., *Blood Coagulation & Fibrinolysis* 2001;12:627-637; Vig et al., *Hematology*, Vol. 6 (3) pp. 205-213 (2001); Vig et al., *Blood coagulation & fibrinolysis*, Vol. 12 (7) pp. 555-561 (2001) Oct; Glidden et al., *Clinical and applied thrombosis/hemostasis*, Vol. 6 (4) pp. 226-233 (2000) Oct; McKenzie et al., *Cardiology*, Vol. 92 (4) pp. 240-247 (1999) Apr; or Davis et al., *Journal of the American Society of Nephrology*, Vol. 6 (4) pp. 1250-1255 (1995).

The term "prolonging clot lysis time" is intended to encompass the situations when the measured clot lysis time for a test sample containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid is prolonged relative to the individual clot lysis time of a control sample containing only the factor VII or factor VII-related polypeptide or the epsilon-aminocaproic acid, respectively, when tested in the same clot lysis assay. The clot lysis time may be assayed as described above.

The term "increasing clot strength" is intended to encompass the situations when the measured clot strength, e.g., mechanical strength, for a test sample containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid is increased relative to the individual clot lysis time of a control sample containing only the factor VII or factor VII-related polypeptide or the epsilon-aminocaproic acid, respectively, when tested in the same clot strength assay. The clot strength may be assayed as described, e.g. in Carr et al, 1991. (Carr ME, Zekert SL. Measurement of platelet-mediated force development during plasma clot formation. AM J MED SCI 1991; 302: 13-8), or as described above by means of thromboelastography.

The term "enhancing fibrin clot formation" is intended to encompass the situations when the measured rate for or degree of fibrin clot formation for a test sample containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of a preparation of epsilon-aminocaproic acid is increased relative to the individual rate for or degree of fibrin clot formation of a control sample containing only the factor VII or factor VII-related polypeptide or the epsilon-aminocaproic acid, respectively, when tested in the same clotting assay. The fibrin clot formation may be assayed as described above.

The term "shortening clotting time" is intended to encompass the situations when the measured time for clot formation (clotting time) for a test sample containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of a preparation of epsilon-aminocaproic acid is increased relative to the individual clotting time of a control sample containing only the factor VII or factor VII-related polypeptide or the epsilon-aminocaproic acid respectively, when tested in the same clotting assay. The clotting time may be assayed by means of standard PT og aPTT assays, which are known to the general skilled person.

The term "lowered count or activity of platelets" refers to the number of platelets (thrombocytes) present in the subject's plasma and to the biological, coagulation-related activity of such platelets. Lowered counts may be due, e.g., to increased platelet destruction, decreased platelet production, and pooling of a larger than normal fraction of platelets in the spleen. Thrombocytopenia, for example, is defined as a platelet count less than 150,000 platelets per microliter; the upper limit of the normal platelet count is generally considered to be between 350,000 and 450,000 platelets per microliter. Platelet count may be measured by automated platelet counters; this is a well known method to the skilled worker. Syndromes due to lowered platelet count include, without limitation, thrombocytopenia, coagulopathy. "Activity" includes, without limitation, aggrega-

tion, adhesion, and coagulant activity of the platelets. Decreased activity may be due, e.g., to glycoprotein abnormalities, abnormal membrane-cytoskeleton interaction, abnormalities of platelet granules, abnormalities of platelet coagulant activity, abnormalities of signal transduction and secretion. Platelet activity, including aggregation, adhesion, and coagulant activity, are measured by standard methods known to the skilled worker, see e.g., Platelets. A Practical Approach, Ed. S.P. Watson & K.S. Authi: Clinical Aspects of Platelet Disorders (K.J. Clemetson) 15:299-318, 1996, Oxford University Press; Williams Hematology, Sixth Edition, Eds. Beutler, Lichtman, Coller, Kipps & Seligsohn, 2001, McGraw-Hill. Syndromes due to lowered platelet activity includes, without limitation, Glanzmann thrombathenesis, Bernard-Soulier syndrome, anticoagulant treatment and thrombolytic treatment. "Lowered" refers to the count or activity of a sample of the test plasma compared to the count or activity in a sample of normal pooled plasma when measured in the same assay

As used herein the term "bleeding disorder" reflects any defect, congenital, acquired or induced, of cellular or molecular origin that is manifested in bleeding episodes. Examples of bleeding disorders include, but are not limited to, clotting factor deficiencies (e.g. deficiency of coagulation factors VIII, IX, XI or VII), clotting factor inhibitors, defective platelet function (e.g., Glanzmann thrombasthenia and Bernard-Soulier syndrome), thrombocytopenia, von Willebrand's disease, and coagulopathy such as that caused by a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to bleedings and/or transfusions (e.g., in multi transfused subjects having been subjected to surgery or trauma).

As used herein, "coagulopathy" refers to a tendency to bleed occurring in subjects having a basically normal coagulation system but experiencing a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to bleedings and/or transfusions (e.g., in multi transfused subjects having been subjected to surgery or trauma).

Bleeding refers to extravasation of blood from any component of the circulatory system. The term "bleeding episodes" is meant to include unwanted, uncontrolled and often excessive bleeding in connection with surgery, trauma, or other forms of tissue damage, as well as unwanted bleedings in subjects having bleeding disorders. Bleeding episodes may occur in subjects having a basically normal coagulation system but experiencing a (temporary) coagulopathy, as well as in subjects having congenital or acquired coagulation or bleeding disorders. In subjects having a defective platelet function, the haemostatic system lacks or has abnormal essential clotting "compounds" (e.g., platelets). In subjects who experience extensive tissue damage, for example in association with surgery or vast trauma, the normal haemostatic mechanism may be overwhelmed by the demand of immediate haemostasis and they may develop excessive bleeding in spite of a basically (pre-trauma or pre-surgery) normal haemostatic mechanism. Such subjects, who further often are multi transfused, develop a (temporary) coagulopathy as a result of the bleeding and/or transfusions (i.e., a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to the bleeding and/or transfusions). Bleedings may also

occur in organs such as the brain, inner ear region and eyes; these are areas with limited possibilities for surgical haemostasis and thus problems with achieving satisfactory haemostasis. Similar problems may arise in the process of taking biopsies from various organs (liver, lung, tumour tissue, gastrointestinal tract) as well as in laparoscopic surgery and radical retropubic prostatectomy. Common for all these situations is the difficulty to provide haemostasis by surgical techniques (sutures, clips, etc.) which also is the case when bleeding is diffuse (e.g., haemorrhagic gastritis and profuse uterine bleeding). Bleedings may also occur in subjects on anticoagulant therapy in whom a defective haemostasis has been induced by the therapy given; these bleedings are often acute and profuse. Anticoagulant therapy is often given to prevent thromboembolic disease. Such therapy may include heparin, other forms of proteoglycans, warfarin or other forms of vitamin K-antagonists as well as aspirin and other platelet aggregation inhibitors, such as, e.g., antibodies or other inhibitors of GP IIb/IIIa activity. The bleeding may also be due to so-called thrombolytic therapy which comprises combined treatment with an antiplatelet agent (e.g., acetylsalicylic acid), an anticoagulant (e.g., heparin), and a fibrinolytic agent (e.g., tissue plasminogen activator, tPA). Bleeding episodes are also meant to include, without limitation, uncontrolled and excessive bleeding in connection with surgery or trauma in subjects having acute haemarthroses (bleedings in joints), chronic haemophilic arthropathy, haematomas, (e.g., muscular, retroperitoneal, sublingual and retropharyngeal), bleedings in other tissue, haematuria (bleeding from the renal tract), cerebral haemorrhage, surgery (e.g., hepatectomy), dental extraction, and gastrointestinal bleedings (e.g., UGI bleeds). The bleeding episodes may be associated with inhibitors against factor VIII; haemophilia A; haemophilia A with inhibitors; haemophilia B; deficiency of factor VII; deficiency of epsilon-aminocaproic acid; thrombocytopenia; deficiency of von Willebrand factor (von Willebrand's disease); severe tissue damage; severe trauma; surgery; laparoscopic surgery; haemorrhagic gastritis; taking biopsies; anticoagulant therapy; upper gastrointestinal bleedings (UGI); or stem cell transplantation. The bleeding episodes may be profuse uterine bleeding; occurring in organs with a limited possibility for mechanical haemostasis; occurring in the brain; occurring in the inner ear region; or occurring in the eyes. The terms "bleeding episodes" and "bleedings" may, where appropriate, be used interchangeably.

30 As used herein, "non-haemophilic" means not lacking blood coagulation factor VIII or factor IX, or von Willebrand's factor, and not having inhibitors to any of these factors. The term "non-haemophilic subject" refers to subjects not lacking coagulation factors VIII or IX, or von Willebrand's factor, and not having inhibitors to any of these factors. The term includes subjects having congenital or acquired haemophilia A or B, with or without inhibitors, and von Willebrand's disease.

35 may be used interchangeably with "subject not lacking coagulation factors VIII or IX, or von Willebrand's factor, and not having inhibitors to any of these factors"

As used herein, the term "non-haemophilic bleeding disorder" reflects any defect, congenital, acquired or induced, of cellular or molecular origin that is manifested in bleeding episodes. Examples of bleeding disorders include, but are not limited to, defective platelet function (e.g., Glanzmann thrombasthenia and Bernard-Soulier syndrome), thrombocytopenia, and coagulopathy such as that caused by a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to bleedings and/or transfusions (e.g., in multi transfused subjects having been subjected to surgery or trauma).

The term "bleeding episodes in non-haemophilic subjects" is meant to include unwanted, uncontrolled and often excessive bleeding in connection with surgery, trauma, or other forms of tissue damage, as well as unwanted bleedings in subjects having non-haemophilic bleeding disorders. Bleeding episodes may occur in subjects having a basically normal coagulation system but experiencing a (temporary) coagulopathy, as well as in subjects having congenital or acquired non-haemophilic bleeding disorders. In subjects having a defective platelet function, the haemostatic system lacks or has abnormal essential clotting "compounds" (e.g., platelets). In subjects who experience extensive tissue damage, for example in association with surgery or vast trauma, the normal haemostatic mechanism may be overwhelmed by the demand of immediate haemostasis and they may develop excessive bleeding in spite of a basically (pre-trauma or pre-surgery) normal haemostatic mechanism. Such subjects, who further often are multi transfused, develop a (temporary) coagulopathy as a result of the bleeding and/or transfusions (i.e., a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to the bleeding and/or transfusions). Bleedings may also occur in organs such as the brain, inner ear region and eyes; these are areas with limited possibilities for surgical haemostasis and thus problems with achieving satisfactory haemostasis. Similar problems may arise in the process of taking biopsies from various organs (liver, lung, tumour tissue, gastrointestinal tract) as well as in laparoscopic surgery and radical retropubic prostatectomy. Common for all these situations is the difficulty to provide haemostasis by surgical techniques (sutures, clips, etc.) which also is the case when bleeding is diffuse (e.g., haemorrhagic gastritis and profuse uterine bleeding). Bleedings may also occur in subjects on anticoagulant therapy in whom a defective haemostasis has been induced by the therapy given; these bleedings are often acute and profuse. Anticoagulant therapy is often given to prevent thromboembolic disease. Such therapy may include heparin, other forms of proteoglycans, warfarin or other forms of vitamin K-antagonists as well as aspirin and other platelet aggregation inhibitors, such as, e.g., antibodies or other inhibitors of GP IIb/IIIa activity. The bleeding may also be due to so-called thrombolytic therapy which comprises combined treatment with an antiplatelet agent (e.g., acetylsalicylic acid), an anticoagulant (e.g., heparin), and a fibrinolytic agent (e.g., tissue plasminogen activator, tPA). The bleeding episodes may be associated with deficiency of epsilon-aminocaproic acid; thrombocytopenia; severe tissue damage; severe trauma; surgery; laparoscopic surgery;

haemorrhagic gastritis; taking biopsies; anticoagulant therapy; upper gastrointestinal bleedings (UGI); or stem cell transplantation. The bleeding episodes may be profuse uterine bleeding; occurring in organs with a limited possibility for mechanical haemostasis; occurring in the brain; occurring in the inner ear region; or occurring in the eyes. The terms "bleeding episodes" and
5 "bleedings" may, where appropriate, be used interchangeably.

In this context, the term "treatment" is meant to include both prevention of an expected bleeding, such as, for example, in surgery, and regulation of an already occurring bleed-
ing, such as, for example, in trauma, with the purpose of inhibiting or minimising the bleeding.
10 The above-referenced "expected bleeding" may be a bleeding expected to occur in a particular tissue or organ, or it may be an unspecified bleeding. Prophylactic administration of a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid is thus included in the term "treatment".

The term "subject" as used herein is intended to mean any animal, in particular mammals,
15 such as humans, and may, where appropriate, be used interchangeably with the term "patient". The present invention also encompasses the use of factor VII or FVII-related polypeptides, and tPA inhibitors within veterinary procedures.

The factor VII or factor VII-related polypeptides and epsilon-aminocaproic acids as de-
fined in the present specification may be administered simultaneously or sequentially. The fac-
20 tors may be supplied in single-dosage form wherein the single-dosage form contains both coagu-
lation factors, or in the form of a kit-of-parts comprising a preparation of factor VII or a factor
VII-related polypeptide as a first unit dosage form and a preparation of epsilon-aminocaproic
acid as a second unit dosage form. Whenever a first or second or third, etc., unit dose is men-
tioned throughout this specification this does not indicate the preferred order of administration,
25 but is merely done for convenience purposes

By "simultaneous" dosing of a preparation of factor VII or a factor VII-related polypep-
tide and a preparation of epsilon-aminocaproic acid is meant administration of the coagulation
factor proteins in single-dosage form, or administration of a first coagulation factor protein fol-
lowed by administration of a second coagulation factor protein with a time separation of no
30 more than 15 minutes, preferably 10, more preferred 5, more preferred 2 minutes. Either factor
may be administered first.

By "sequential" dosing is meant administration of a first coagulation factor protein fol-
lowed by administration of a second coagulation factor protein with a time separation of up to
2 hours, preferably from 1 to 2 hours, more preferred up to 1 hour, more preferred from 30
35 minutes to 1 hour, more preferred up to 30 minutes, more preferred from 15 to 30 minutes.
Either of the two unit dosage form, or coagulation factor proteins, may be administered first.
Preferably, both products are injected through the same intravenous access.

By "level of factor VII" or "factor VII level" is meant the level of the patient's factor VII activity compared to the level in healthy subjects. The level is designated as a percentage of the normal level. The terms may, where appropriate, be used interchangeably.

By "reduced level of factor VII" or "reduced factor VII level" is meant a decrease in the presence or activity of factor VII in the blood stream compared to the mean factor VII level in a population of subjects having no factor VII deficiency. The level of circulating factor VII can be measured by either a coagulant or an immunologic assay. Factor VII activity is determined by the ability of the patient's plasma to correct the clotting time of factor VII-deficient plasma (e.g., an APTT assay, see below; see also "assay part" of the present description).

One unit of factor VII is defined as the amount of factor VII present in 1 ml of normal plasma, corresponding to about 0.5 µg protein. After activation 50 units correspond to about 1 µg protein.

By "deficiency" is meant a decrease in the presence or activity of, e.g., factor VII in plasma compared to that of normal healthy individuals. The term may, where appropriate, be used interchangeably with "reduced factor VII level".

By "APTT" or "aPTT" is meant the activated partial thromboplastin time (described by, e.g., Proctor RR, Rapaport SI: The partial thromboplastin time with kaolin; a simple screening test for first-stage plasma clotting factor deficiencies. Am J Clin Pathol 36:212, 1961).

By "factor VII-responsive syndrome" is meant a syndrome where exogenous factor VII, preferably factor VIIa, administered to the subject in need thereof may prevent, cure or ameliorate any symptoms, conditions or diseases, expected or present, caused by the syndrome. Included are, without limitation, syndromes caused by a reduced level of clotting factors VIII, IX, XI or VII, clotting factor inhibitors, defective platelet function (e.g., Glanzmann thrombasthenia and Bernard-Soulier syndrome), thrombocytopenia, von Willebrand's disease, and coagulopathy such as that caused by a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to bleedings and/or transfusions (e.g., in multi transfused subjects having been subjected to surgery or trauma).

"Half-life" refers to the time required for the plasma concentration of factor VII or a factor VII-related polypeptide, or epsilon-aminocaproic acid to decrease from a particular value to half of that value.

By "primary haemostasis" is meant the initial generation of thrombin by FXa and TF:factor VIIa, the subsequent activation of platelets and formation of the initial loose plug of activated, adhered platelets which has not yet been stabilized by fibrin and, finally, by cross-linked fibrin. If not stabilized by the fibrin formed during the second step of the haemostatic process (maintained haemostasis), the plug is easily dissolved by the fibrinolytic system.

By "secondary haemostasis" or "maintained haemostasis" is meant the secondary, full, and major, burst or generation of thrombin taking place on the surface of activated platelets

and catalysed by factor VIIIa and factor XIIIa, the subsequent formation of fibrin and the stabilization of the initial platelet plug. Stabilization of the plug by fibrin leads to full haemostasis.

By "full haemostasis" is meant the formation of a stable and solid fibrin clot or plug at the site of injury which effectively stops the bleeding and which is not readily dissolved by the fibrinolytic system. In this context, the term haemostasis will be used to represent full haemostasis as described above.

The total amount of protein in a preparation may be measured by generally known methods, e.g. by measuring optical density. Amounts of epsilon-aminocaproic acid- or factor VII protein ("antigen") may be measured by generally known methods such as standard Elisa immuno assays. In general terms, such assay is conducted by contacting, e.g., a solution of the epsilon-aminocaproic acid protein-containing preparation with an anti-thrombomodulin antibody immobilised onto the elisa plate, subsequently contacting the immobilised antibody-epsilon-aminocaproic acid complex with a second anti-epsilon-aminocaproic acid antibody carrying a marker, the amounts of which, in a third step, are measured. The amounts of each coagulation factor may be measured in a similar way using appropriate antibodies. The total amount of coagulation factor protein present in a preparation is determined by adding the amounts of the individual coagulation factor proteins. In one embodiment, the preparation comprises isolated coagulation factor. In another embodiment the preparation is essentially free of coagulation factor II and coagulation factor IIa (prothrombin and thrombin) and/or factor X or Xa.

As used herein, the term "isolated" refers to coagulation factors, e.g., epsilon-aminocaproic acids that have been separated from the cell in which they were synthesized or the medium in which they are found in nature (e.g., plasma or blood). Separation of polypeptides from their cell of origin may be achieved by any method known in the art, including, without limitation, removal of cell culture medium containing the desired product from an adherent cell culture; centrifugation or filtration to remove non-adherent cells; and the like. Separation of polypeptides from the medium in which they naturally occur may be achieved by any method known in the art, including, without limitation, affinity chromatography, such as, e.g., on an anti-factor VII or anti-epsilon-aminocaproic acid antibody column, respectively; hydrophobic interaction chromatography; ion-exchange chromatography; size exclusion chromatography; electrophoretic procedures (e.g., preparative isoelectric focusing (IEF)), differential solubility (e.g., ammonium sulfate precipitation), or extraction and the like.

Within the present invention an "effective amount" of factor VII or a factor VII-related polypeptide, and epsilon-aminocaproic acid is defined as the amount of factor VII or a factor VII-related polypeptide, e.g., FVIIa, and epsilon-aminocaproic acid, that together suffices to prevent or reduce bleeding or blood loss, so as to cure, alleviate or partially arrest the disease and its complications.

The term "activity of factor VIIa" or "factor VIIa-activity" includes the ability to generate thrombin; the term also includes the ability to generate thrombin on the surface of activated platelets in the absence of tissue factor.

5 Abbreviations

TF	tissue factor
FVII	factor VII in its single-chain, unactivated form
FVIIa	factor VII in its activated form
rFVIIa	recombinant factor VII in its activated form
10 TAFI	TAFI in its zymogenic, unactivated form

Preparation of compounds:

Human purified factor VIIa suitable for use in the present invention is preferably made by DNA recombinant technology, e.g. as described by Hagen et al., Proc.Natl.Acad.Sci. USA 83: 2412-
15 2416, 1986, or as described in European Patent No. 200.421 (ZymoGenetics, Inc.).

Factor VII may also be produced by the methods described by Broze and Majerus, J.Biol.Chem. 255 (4): 1242-1247, 1980 and Hedner and Kisiel, J.Clin.Invest. 71: 1836-1841, 1983. These methods yield factor VII without detectable amounts of other blood coagulation factors. An even further purified factor VII preparation may be obtained by including an additional gel filtration as
20 the final purification step. factor VII is then converted into activated factor VIIa by known means, e.g. by several different plasma proteins, such as epsilon-aminocaproic acid, IX a or Xa. Alternatively, as described by Bjoern et al. (Research Disclosure, 269 September 1986, pp. 564-565), factor VII may be activated by passing it through an ion-exchange chromatography column, such as Mono Q® (Pharmacia fine Chemicals) or the like.

25 Factor VII -related polypeptides may produced by modification of wild-type factor VII or by recombinant technology. factor VII -related polypeptides with altered amino acid sequence when compared to wild-type factor VII may be produced by modifying the nucleic acid sequence encoding wild-type factor VII either by altering the amino acid codons or by removal of some of the amino acid codons in the nucleic acid encoding the natural factor VII by known means, e.g. by site-specific
30 mutagenesis.

It will be apparent to those skilled in the art that substitutions can be made outside the regions critical to the function of the factor VIIa and still result in an active polypeptide. Amino acid residues essential to the activity of the factor VII or factor VII-related polypeptide, and therefore preferably not subject to substitution, may be identified according to procedures
35 known in the art, such as site-directed mutagenesis or alanine-scanning mutagenesis (see, e.g., Cunningham and Wells, 1989, *Science* 244: 1081-1085). In the latter technique, mutations are introduced at every positively charged residue in the molecule, and the resultant mutant mole-

cules are tested for coagulant, respectively cross-linking activity to identify amino acid residues that are critical to the activity of the molecule. Sites of substrate-enzyme interaction can also be determined by analysis of the three-dimensional structure as determined by such techniques as nuclear magnetic resonance analysis, crystallography or photoaffinity labelling (see, e.g., de Vos
5 *et al.*, 1992, *Science* 255: 306-312; Smith *et al.*, 1992, *Journal of Molecular Biology* 224: 899-904; Wlodaver *et al.*, 1992, *FEBS Letters* 309: 59-64).

The introduction of a mutation into the nucleic acid sequence to exchange one nucleotide for another nucleotide may be accomplished by site-directed mutagenesis using any of the methods known in the art. Particularly useful is the procedure that utilizes a super coiled, double
10 stranded DNA vector with an insert of interest and two synthetic primers containing the desired mutation. The oligonucleotide primers, each complementary to opposite strands of the vector, extend during temperature cycling by means of *Pfu* DNA polymerase. On incorporation of the primers, a mutated plasmid containing staggered nicks is generated. Following temperature cycling, the product is treated with *DpnI*, which is specific for methylated and hemi-
15 methylated DNA to digest the parental DNA template and to select for mutation-containing synthesized DNA. Other procedures known in the art for creating, identifying and isolating variants may also be used, such as, for example, gene shuffling or phage display techniques.

Separation of polypeptides from their cell of origin may be achieved by any method known in the art, including, without limitation, removal of cell culture medium containing the
20 desired product from an adherent cell culture; centrifugation or filtration to remove non-adherent cells; and the like.

Optionally, factor VII or factor VII-related polypeptides may be further purified. Purification may be achieved using any method known in the art, including, without limitation, affinity chromatography, such as, e.g., on an anti-factor VII antibody column (see, e.g., Wakabayashi
25 *et al.*, *J. Biol. Chem.* 261:11097, 1986; and Thim *et al.*, *Biochem.* 27:7785, 1988); hydrophobic interaction chromatography; ion-exchange chromatography; size exclusion chromatography; electrophoretic procedures (e.g., preparative isoelectric focusing (IEF), differential solubility (e.g., ammonium sulfate precipitation), or extraction and the like. See, generally, Scopes, *Protein Purification*, Springer-Verlag, New York, 1982; and *Protein Purification*, J.C. Janson and Lars Ryden,
30 editors, VCH Publishers, New York, 1989. Following purification, the preparation preferably contains less than about 10% by weight, more preferably less than about 5% and most preferably less than about 1%, of non-factor VII or factor VII-related polypeptides derived from the host cell.

Factor VII or factor VII-related polypeptides may be activated by proteolytic cleavage,
35 using factor XIa or other proteases having trypsin-like specificity, such as, e.g., factor IXa, kallikrein, factor Xa, and thrombin. See, e.g., Osterud *et al.*, *Biochem.* 11:2853 (1972); Thomas, U.S. Patent No. 4,456,591; and Hedner *et al.*, *J. Clin. Invest.* 71:1836 (1983). Alternatively, factor VII or

factor VII-related polypeptides may be activated by passing it through an ion-exchange chromatography column, such as Mono Q® (Pharmacia) or the like. The resulting activated factor VII or factor VII-related polypeptide may then be formulated and administered as described below.

Epsilon-aminocaproic acid for use within the present invention may be isolated from, e.g., placenta or lung, according to known methods; it is preferred, however, to use recombinant epsilon-aminocaproic acid so as to avoid to the use of blood- or tissue-derived products that carry a risk of disease transmission. Methods for isolating epsilon-aminocaproic acid and preparing recombinant epsilon-aminocaproic acid are known in the art; see, for example, Gomi et al., Blood 75:1396, 1990 ; Ogata et al., Appl Microbiol Biotechnol 38: 520, 1993 . Epsilon-aminocaproic acid variants may be produced by means of site-directed mutagenesis as described above.

Pharmaceutical Compositions and Methods of Use

The preparations of the present invention may be used to treat any factor VII responsive syndrome, such as, e.g., bleeding disorders, including, without limitation, syndromes caused by a reduced level of clotting factors VIII, IX, XI or VII, clotting factor inhibitors, defective platelet function (e.g., Glanzmann thrombasthenia and Bernard-Soulier syndrome), thrombocytopenia, von Willebrand's disease, and coagulopathy such as that caused by a dilution of coagulation proteins, increased fibrinolysis and lowered number of platelets due to bleedings and/or transfusions (e.g., in multi transfused subjects having been subjected to surgery or trauma). Pharmaceutical compositions comprising a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid according to the present invention are primarily intended for parenteral administration for prophylactic and/or therapeutic treatment. Preferably, the pharmaceutical compositions are administered parenterally, i.e., intravenously, subcutaneously, or intramuscularly; intravenously being most preferred. They may also be administered by continuous or pulsatile infusion.

Pharmaceutical compositions or formulations according to the invention comprise a factor VII or a factor VII-related polypeptide, and epsilon-aminocaproic acid, either formulated in a single-unit dosage form or in the form of a kit-of parts, preferably dissolved in, a pharmaceutically acceptable carrier, preferably an aqueous carrier or diluent. Briefly, pharmaceutical compositions suitable for use according to the present invention is made by mixing factor VII or a factor VII-related polypeptide, or a epsilon-aminocaproic acid, or factor VII or a factor VII-related polypeptide in combination with a epsilon-aminocaproic acid, preferably in purified form, with suitable adjuvants and a suitable carrier or diluent. A variety of aqueous carriers may be used, such as water, buffered water, 0.4% saline, 0.3% glycine and the like. The preparations of the invention can also be formulated using non-aqueous carriers, such as, e.g., in the form of a gel or as liposome

preparations for delivery or targeting to the sites of injury. Liposome preparations are generally described in, e.g., U.S. Patents Nos. 4,837,028, 4,501,728, and 4,975,282. The compositions may be sterilised by conventional, well-known sterilisation techniques. The resulting aqueous solutions may be packaged for use or filtered under aseptic conditions and lyophilised, the lyophilised preparation being combined with a sterile aqueous solution prior to administration.

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.

Formulations may further include one or more diluents, emulsifiers, preservatives, buffers, excipients, etc. and may be provided in such forms as liquids, powders, emulsions, controlled release, etc. One skilled in this art may formulate the compositions of the invention in an appropriate manner, and in accordance with accepted practices, such as those disclosed in Remington's Pharmaceutical Sciences, Gennaro, ed., Mack Publishing Co., Easton, PA, 1990.

Thus, a typical pharmaceutical composition for intravenous infusion could be made up to contain 250 ml of sterile Ringer's solution and 10 mg of the preparation.

The compositions containing the preparations of the present invention can be administered for prophylactic and/or therapeutic treatments. In therapeutic applications, compositions are administered to a subject already suffering from a disease, as described above, in an amount sufficient to cure, alleviate or partially arrest the clinical manifestations of the disease and its complications. An amount adequate to accomplish this is defined as "therapeutically effective amount". Effective amounts for each purpose will depend on the severity of the disease or injury as well as the weight and general state of the subject. It will be understood that determining an appropriate dosage may be achieved using routine experimentation, by constructing a matrix of values and testing different points in the matrix.

Local delivery of the preparations of the present invention, such as, for example, topical application, may be carried out, e.g., by means of a spray, perfusion, double balloon catheters, stent, incorporated into vascular grafts or stents, hydrogels used to coat balloon catheters, or other well established methods. In any event, the pharmaceutical compositions should provide a quantity of the preparation sufficient to effectively treat the condition.

The concentration of factor VII or factor VII-related polypeptide, epsilon-aminocaproic acid, or factor VII or factor VII-related polypeptide in combination with epsilon-aminocaproic acid in these formulations can vary widely, i.e., from less than about 0.5% by weight, usually at or at least about 1% by weight to as much as 15 or 20% by weight and will be selected primarily by fluid volumes, viscosities, etc., in accordance with the particular mode of administration selected. Administration by injection or infusion, in particular injection, is preferred. Thus, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid are prepared in a

form suitable for intravenous administration, such as a preparation that is either a dissolved lyophilized powder or a liquid formulation containing both the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid in one dosage form, or a dissolved lyophilized powder or a liquid formulation containing the factor VII or factor VII-related polypeptide in one dosage form and dissolved lyophilized powder or a liquid formulation containing the epsilon-aminocaproic acid in another dosage form.

It is to be understood that the amount of factor VII or factor VII-related polypeptide and the amount of epsilon-aminocaproic acid together comprise an aggregate effective amount for treating the bleeding episode.

It must be kept in mind that the materials of the present invention may generally be employed in serious disease or injury states, that is, life threatening or potentially life threatening situations. In such cases, in view of the minimization of extraneous substances and general lack of immunogenicity of factor VIIa and epsilon-aminocaproic acid in humans, it is possible and may be felt desirable by the treating physician to administer a substantial excess of these compositions.

In prophylactic applications, compositions containing a preparation of factor VII or a factor VII-related polypeptide and a preparation of epsilon-aminocaproic acid are administered to a subject susceptible to or otherwise at risk of a disease state or injury to enhance the subject's own coagulative capability. Such an amount is defined to be a "prophylactically effective dose." It is to be understood that the amount of factor VII or factor VII-related polypeptide and the amount of epsilon-aminocaproic acid together comprise an aggregate effective amount for preventing a bleeding episode.

Single or multiple administrations of the compositions can be carried out with dose levels and patterns being selected by the treating physician. The compositions may be administered one or more times per day or week. An effective amount of such a pharmaceutical composition is the amount that provides a clinically significant effect against bleeding episodes. Such amounts will depend, in part, on the particular condition to be treated, age, weight, and general health of the subject, and other factors evident to those skilled in the art.

The composition of the invention is generally administered in a single dose before the expected bleeding or at the start of the bleeding. It may however also be given repeatedly (in multiple doses) preferably with intervals of 2-4-6-12 hour, depending on the dose given and the condition of the subject.

For treatment in connection with deliberate interventions, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid will typically be administered within about 24 hours prior to performing the intervention, and for as much as 7 days or more thereafter. Administration as a coagulant can be by a variety of routes as described herein.

The composition may be in the form of a single preparation (single-dosage form) comprising both a preparation of a preparation of factor VII or a factor VII-related polypeptide and a preparation of a preparation of epsilon-aminocaproic acid in suitable concentrations. The composition may also be in the form of a kit-of-parts consisting of a first unit dosage form
5 comprising a preparation of a preparation of factor VII or a factor VII-related polypeptide and a second unit dosage form comprising a preparation of a preparation of epsilon-aminocaproic acid. In this case, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid should be administered one after the other, preferably within about 15 minutes of each other, for example within 10 minutes of each other or, preferably, within 5 minutes or, more preferred, within
10 2 minutes of each other. Either of the two unit dosage forms can be administered first.

The kit includes at least two separate pharmaceutical compositions. The kit includes container means for containing the separate compositions such as a divided bottle or a divided foil packet. Typically the kit includes directions for the administration of the separate components. The kit form is particularly advantageous when the separate components are preferably administered in
15 different dosage forms, are administered at different dosage intervals, or when titration of the individual components of the combination is desired by the prescribing physician.

The amount of factor VII or factor VII-related polypeptide and the amount of epsilon-aminocaproic acid administered according to the present invention may vary from a ratio of between about 1:100 to about 100:1 (w/w). The ratio of factor VII to epsilon-aminocaproic acid
20 may thus be, e.g., about 1:100, or 1:90, or 1:80, or 1:70 or 1:60, or 1:50, or 1:40, or 1:30, or 1:20, or 1:10, or 1:5, or 1:2, or 1:1, or 2:1, or 5:1, or 10:1, or 20:1, or 30:1, or 40:1, or 50:1, or 60:1, or 70:1, or 80:1, or 90:1, or 100:1; or between about 1:90 to about 1:1, or between about 1:80 to about 1:2, or between about 1:70 to about 1:5, or between about 1:60 to about 1:10, or between about 1:50 to about 1:25, or between about 1:40 to about 1:30, or between about 90:1 to about 1:1, or between
25 about 80:1 to about 2:1, or between about 70:1 to about 5:1, or between about 60:1 to about 10:1, or between about 50:1 to about 25:1, or between about 40:1 to about 30:1.

The dose of the Factor VII polypeptide ranges from about 0.05 mg to about 500 mg/day, e.g., from about 1 mg to about 200 mg/day, or, e.g., from about 5 mg to about 175 mg/day for a 70-kg subject as loading and maintenance doses, depending on the weight of the
30 subject, the condition and the severity of the condition.

The dose of epsilon-aminocaproic acid ranges from about 0.05 mg to about 6000 mg/day, e.g., from about 1 mg to about 5000 mg/day, or, e.g., from about 5 mg to about 5000 mg/day for a 70-kg subject as loading and maintenance doses, depending on the weight of the
subject, the condition and the severity of the condition.

35 The combination of factor VII or a factor VII-related polypeptide and epsilon-aminocaproic acid shows a synergistic effect in an *in vitro* clot firmness- and fibrinolysis time-assay. Moreover, the combination of factor VII or a factor VII-related polypeptide and epsilon-

aminocaproic acid shows a synergistic effect in forming stable fibrin clots, increasing the half-clot lysis time, increasing clot strength and increasing resistance to fibrinolysis.

The composition may be in the form of a single preparation comprising both factor VII or a factor VII-related polypeptide and epsilon-aminocaproic acid in suitable concentrations. The composition may also be in the form of a kit consisting of a first unit dosage form comprising factor VII or a factor VII-related polypeptide, and a second unit dosage form comprising epsilon-aminocaproic acid. In this case, the factor VII or factor VII-related polypeptide and the epsilon-aminocaproic acid should be administered sequentially, preferably within about 1-2 hours of each other, for example within 30 minutes of each other or, preferably, within 10 minutes or, more preferred, within 5 minutes of each other. Either of the two unit dosage forms can be administered first.

Since the present invention relates to the prevention or treatment of bleeding episodes or for coagulative treatment by treatment with a combination of active ingredients that may be administered separately, the invention also relates to combining separate pharmaceutical compositions in kit form. The kit includes at least two separate pharmaceutical compositions. The kit includes container means for containing the separate compositions such as a divided bottle or a divided foil packet. Typically the kit includes directions for the administration of the separate components. The kit form is particularly advantageous when the separate components are preferably administered in different dosage forms, are administered at different dosage intervals, or when titration of the individual components of the combination is desired by the prescribing physician

Assays:

25 Test for factor VIIa activity:

A suitable assay for testing for factor VIIa activity and thereby selecting suitable factor VIIa variants can be performed as a simple preliminary *in vitro* test:

In Vitro Hydrolysis Assay

Native (wild-type) factor VIIa and factor VIIa variant (both hereafter referred to as "factor VIIa") may be assayed for specific activities. They may also be assayed in parallel to directly compare their specific activities. The assay is carried out in a microtiter plate (MaxiSorp, Nunc, Denmark). The chromogenic substrate D-Ile-Pro-Arg-p-nitroanilide (S-2288, Chromogenix, Sweden), final concentration 1 mM, is added to factor VIIa (final concentration 100 nM) in 50 mM HEPES, pH 7.4, containing 0.1 M NaCl, 5 mM CaCl₂ and 1 mg/ml bovine serum albumin. The absorbance at 405 nm is measured continuously in a SpectraMax™ 340 plate reader (Molecular Devices, USA). The absorbance developed during a 20-minute incubation, after subtraction of the absorbance in a blank well containing no enzyme, is used to calculate the ratio between the activities of variant and wild-type factor VIIa:

$$\text{Ratio} = (A_{405 \text{ nm}} \text{ factor VIIa variant}) / (A_{405 \text{ nm}} \text{ factor VIIa wild-type}).$$

Based thereon, factor VIIa variants with an activity comparable to or higher than native factor VIIa may be identified, such as, for example, variants where the ratio between the activity of the variant and the activity of native factor VII (wild-type FVII) is around, versus above 1.0.

5

The activity of factor VIIa or factor VIIa variants may also be measured using a physiological substrate such as factor X, suitably at a concentration of 100-1000 nM, where the factor Xa generated is measured after the addition of a suitable chromogenic substrate (eg. S-2765). In addition, the activity assay may be run at physiological temperature.

10

In Vitro Proteolysis Assay

Native (wild-type) factor VIIa and factor VIIa variant (both hereafter referred to as "factor VIIa") are assayed in parallel to directly compare their specific activities. The assay is carried out in a microtiter plate (MaxiSorp, Nunc, Denmark). factor VIIa (10 nM) and factor X (0.8 microM) in 100 microL 50 mM Hepes, pH 7.4, containing 0.1 M NaCl, 5 mM CaCl₂ and 1 mg/ml bovine serum albumin, are incubated for 15 min. factor X cleavage is then stopped by the addition of 50 microL 50 mM Hepes, pH 7.4, containing 0.1 M NaCl, 20 mM EDTA and 1 mg/ml bovine serum albumin. The amount of factor Xa generated is measured by addition of the chromogenic substrate Z-D-Arg-Gly-Arg-p-nitroanilide (S-2765, Chromogenix, Sweden), final concentration 0.5 mM. The absorbance at 405 nm is measured continuously in a SpectraMax™ 340 plate reader (Molecular Devices, USA). The absorbance developed during 10 minutes, after subtraction of the absorbance in a blank well containing no FVIIa, is used to calculate the ratio between the proteolytic activities of variant and wild-type factor VIIa:

15
20
25

$$\text{Ratio} = (A_{405 \text{ nm}} \text{ factor VIIa variant}) / (A_{405 \text{ nm}} \text{ factor VIIa wild-type}).$$

Based thereon, factor VIIa variants with an activity comparable to or higher than native factor VIIa may be identified, such as, for example, variants where the ratio between the activity of the variant and the activity of native factor VII (wild-type FVII) is around, versus above 1.0.

Thrombin generation assay:

30

The ability of factor VII or factor VII-related polypeptides or epsilon-aminocaproic acids (e.g., variants) to generate thrombin can be measured in an assay comprising all relevant coagulation factors and inhibitors at physiological concentrations and activated platelets (as described on p. 543 in Monroe et al. (1997) Brit. J. Haematol. 99, 542-547 which is hereby incorporated as reference).

35

The present invention is further illustrated by the following examples, which, however, are not to be construed as limiting the scope of protection. The features disclosed in the fore-

going description and in the following examples may, both separately and in any combination thereof, be material for realizing the invention in diverse forms thereof.

5 EXAMPLES'

Example 1

Improving Haemostatic Clot Stability by Combining Coagulation Factor VIIa and epsilon-Aminocaproic Acid

10 METHODS:

Clot lysis assay: Normal human plasma diluted 10-fold with buffer (20 mM HEPES, 150 mM NaCl, 5 mM CaCl, pH 7.4) containing Innovin (Dade Behring, 2000-fold dilution), rFVIIa (Novo Nordisk A/S Bagsvaerd, Denmark, various concentrations) and t-PA (American Diagnostica, 8 nM) was added to 96-well ELISA plates and turbidity at 650 nm was measured over time at room temperature. Where indicated, epsilon-Aminocaproic Acid (Sigma, various concentrations) was included.

RESULTS:

Clot lysis assay: Addition of FVIIa results in a dose-dependent prolongation of the clot lysis time (Fig. 1). This effect was optimal at 10 nM FVIIa. In the presence of 10 nM FVIIa, addition of ϵ -Aminocaproic Acid resulted in a further prolongation of the clot lysis time (Fig. 2). The effect was dose-dependent and optimal at 1 μ M epsilon-Aminocaproic Acid.

CONCLUSION:

25 These results demonstrate that FVIIa and epsilon-Aminocaproic Acid addition to plasma in a synergistic fashion improve clot resistance to fibrinolysis.

CLAIMS

1. A pharmaceutical composition comprising a factor VII-related polypeptide and epsilon-aminocaproic acid.
- 5
2. A composition according to claim 1, wherein said factor VII-related polypeptide is a factor VII amino acid sequence variant.
3. A composition according to claim 1 or claim 2, wherein the ratio between the activity of said
10 factor VII-related polypeptide and the activity of native human factor VIIa (wild-type FVIIa) is at least about 1.25 when tested in the "In Vitro Hydrolysis Assay" as described in the present description.
4. A composition according to any one of claims 1 to 3, wherein said factor VII-related
15 polypeptide and epsilon-aminocaproic acid are present in a ratio by mass of between about 100:1 and about 1:100 (w/w factor VII:epsilon-aminocaproic acid)
5. A composition according to any one of claims 1 to 4, wherein the composition further comprises pharmaceutically acceptable excipients suitable for injection or infusion, in particular
20 injection.
6. A composition according to any of claims 1 to 5, which is in the form of a kit of parts, comprising
- 25
- a) An effective amount of a preparation of a factor VII-related polypeptide and a pharmaceutically acceptable carrier in a first-unit dosage form;
 - b) An effective amount of a preparation of epsilon-aminocaproic acid and a pharmaceutically acceptable carrier in a second-unit dosage form; and
 - c) Container means for containing said first and second dosage forms.
- 30
7. Use of a factor VII-related polypeptide in combination with epsilon-aminocaproic acid for the manufacture of a medicament for treating bleeding episodes.
8. Use of a composition according to any one of claims 1 to 6, for the manufacture of a medicament for treating bleeding episodes.
- 35
9. Use according to claim 7 or claim 8, wherein the medicament is for reducing clotting time.

10. Use according to claim 7 or claim 8, wherein the medicament is for prolonging the clot lysis time.
11. Use according to claim 7 or claim 8, wherein the medicament is for increasing clot strength.
- 5 12. Use according to any one of claims 7 to 11, wherein the medicament is formulated for injection or infusion, in particular injection.
- 10 13. Use according to any one of claims 7 to 12, wherein the bleeding episodes are due to trauma, surgery, or lowered count or activity of platelets.
14. Use according to any one of claims 7 to 12, wherein the bleeding episodes are due to lack of coagulation factors VIII or IX, or lack of von Willebrand factor.
- 15 15. Use according to any one of claims 7 to 12, wherein the bleeding episodes are due to anticoagulant treatment.
16. Use according to any one of claims 7 to 15, wherein the medicament is in single-dosage form.
- 20 17. Use according to any one of claims 7 to 15, wherein the medicament is prepared in the form of a first unit dosage form comprising a preparation of a factor VII-related polypeptide and a second unit dosage form comprising a preparation of epsilon-aminocaproic acid.
- 25 18. A method for treating bleeding episodes in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amount together are effective to treat bleedings.
- 30 19. A method for reducing clotting time in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to reduce clotting time.
- 35 20. A method to enhance haemostasis in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a

second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to enhance haemostasis.

21. A method for prolonging the clot lysis time in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to prolong the clot lysis time.

22. A method for increasing clot strength in a subject, the method comprising administering to a subject in need thereof a first amount of a preparation of a factor VII-related polypeptide and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to increase clot strength.

23. Method according to any one of claims 18 to 22, wherein the factor VII-related polypeptide, and the epsilon-aminocaproic acid are administered in single-dosage form.

24. Method according to any one of claims 18 to 22, wherein the factor VII-related polypeptide and the epsilon-aminocaproic acid are administered in the form of a first-unit dosage form comprising a preparation of a factor VII-related polypeptide, and a second-unit dosage form comprising a preparation of epsilon-aminocaproic acid.

25. Method according to claim 24, wherein the first dosage form and the second dosage form are administered with a time separation of no more than 15 minutes.

26. A kit containing a treatment for bleeding episodes comprising

- a) An effective amount of a factor VII-related polypeptide and an effective amount of epsilon-aminocaproic acid and a pharmaceutically acceptable carrier in a single-unit dosage form; and
- b) Container means for containing said single-unit dosage form.

27. Use of factor VII in combination with epsilon-aminocaproic acid for the manufacture of a medicament for treating bleeding episodes in non-haemophilic subjects.

28. Use according to claim 27, wherein the medicament is for reducing clotting time.

29. Use according to claim 27, wherein the medicament is for prolonging the clot lysis time.

30. Use according to claim 27, wherein the medicament is for increasing clot strength.
31. Use according to any one of claims 27 to 30, wherein the medicament is formulated for injection or infusion, in particular injection.
- 5 32. Use according to any one of claims 27 to 31, wherein the bleeding episodes are due to coagulopathy or lowered count or activity of platelets.
33. Use according to claim 32, wherein the bleeding episodes are due to coagulopathy.
- 10 34. Use according to claim 32, wherein the bleeding episodes are due to lowered count of platelets.
35. Use according to any one of claims 27 to 31, wherein the bleeding episodes are due to
- 15 anticoagulant treatment.
36. Use according to any one of claims 27 to 35, wherein the medicament is in single-dosage form.
- 20 37. Use according to any one of claims 27 to 35, wherein the medicament is prepared in the form of a first unit dosage form comprising a preparation of factor VII, and a second unit dosage form comprising a preparation of epsilon-aminocaproic acid.
38. A method for treating bleeding episodes in a non-haemophilic subject, the method comprising
- 25 administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid, wherein the first and second amount together are effective to treat bleedings.
39. A method for reducing clotting time in a non-haemophilic subject, the method comprising
- 30 administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to reduce clotting time.
40. A method to enhance haemostasis in a non-haemophilic subject, the method comprising
- 35 administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to enhance haemostasis.

41. A method for prolonging the clot lysis time in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to prolong the clot lysis time.

42. A method for increasing clot strength in a non-haemophilic subject, the method comprising administering to a subject in need thereof a first amount of a preparation of factor VII and a second amount of a preparation of epsilon-aminocaproic acid wherein the first and second amount together are effective to increase clot strength.

43. Method according to any one of claims 38 to 42, wherein factor VII and epsilon-aminocaproic acid are administered in single-dosage form.

44. Method according to any one of claims 38 to 42, wherein factor VII and epsilon-aminocaproic acid are administered in the form of a first-unit dosage form comprising a preparation of factor VII, and a second-unit dosage form comprising a preparation of epsilon-aminocaproic acid.

45. Method according to claim 44, wherein the first dosage form and the second dosage form are administered with a time separation of no more than 15 minutes.

46. A kit containing a treatment for non-haemophilic bleeding episodes comprising

- c) An effective amount of factor VII and an effective amount of epsilon-aminocaproic acid and a pharmaceutically acceptable carrier in a single-unit dosage form; and
- d) Container means for containing said single-unit dosage form.

1/2

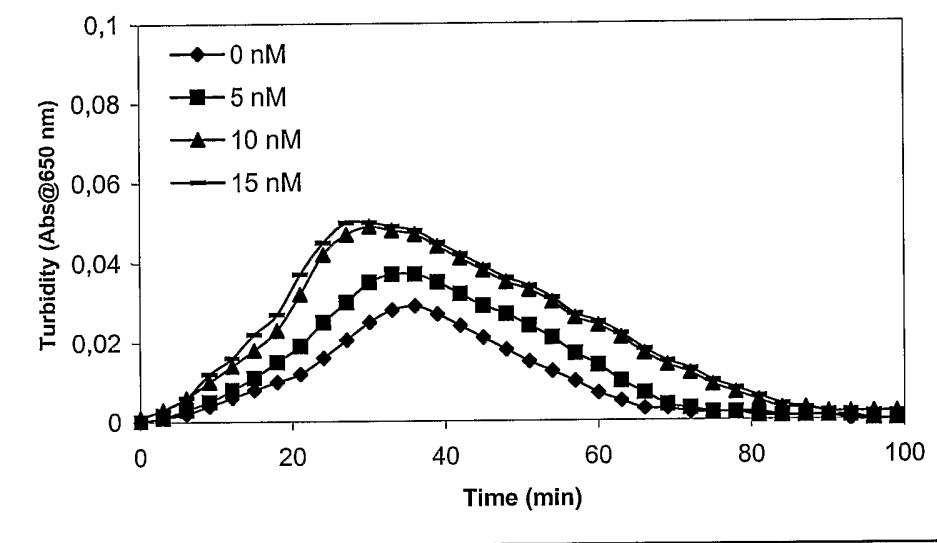
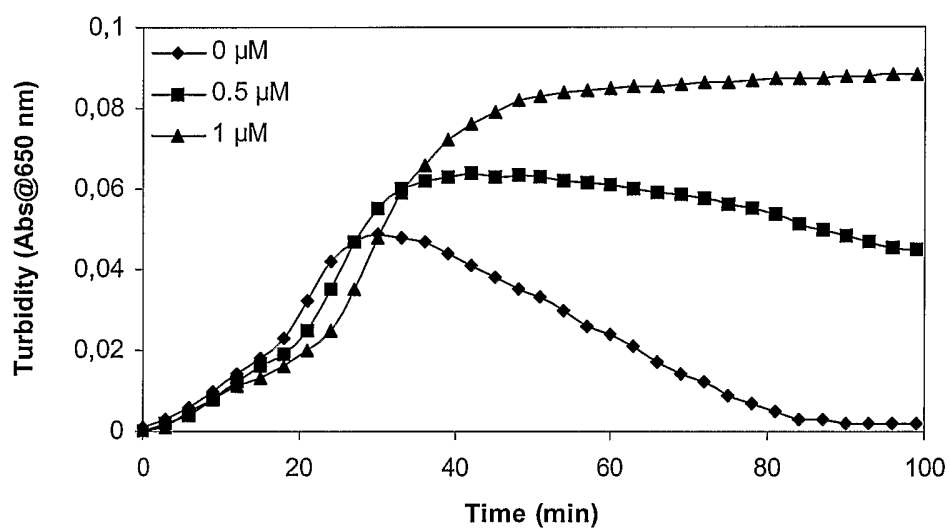


Figure 1

2/2

**Figure 2**

INTERNATIONAL SEARCH REPORT

International Application No
PCT/DK 02/00752

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K38/36 A61K38/48

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

BIOSIS, EPO-Internal, MEDLINE, WPI Data, PAJ, EMBASE, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>HIRAWAT SAMIT ET AL: "Recombinant factor VIIa prophylaxis for a patient with severe factor VII deficiency undergoing dental extraction." BLOOD, vol. 96, no. 11 Part 2, 16 November 2000 (2000-11-16), page 84b XP002231356 42nd Annual Meeting of the American Society of Hematology; San Francisco, California, USA; December 01-05, 2000 ISSN: 0006-4971 abstract</p> <p style="text-align: center;">--- -/--</p>	1-46

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

17 February 2003

Date of mailing of the international search report

10. 03. 2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Tereše Persson

INTERNATIONAL SEARCH REPORT

International Application No
PCT/DK 02/00752

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	MATHEW PRASAD ET AL: "Successful use of rFVIIa (NovoSeven) and fibrin sealant (Tisseal) during dental work and tooth extraction in a pediatric patient with Glanzmann's thrombasthenia." BLOOD, vol. 96, no. 11 Part 2, 16 November 2000 (2000-11-16), page 67b XP002231357 42nd Annual Meeting of the American Society of Hematology; San Francisco, California, USA; December 01-05, 2000 ISSN: 0006-4971 abstract	1-46
X	--- WO 93 06855 A (NOVONORDISK AS) 15 April 1993 (1993-04-15) page 9, line 22 - line 23; claims 1,14	1-46
Y	--- MUNOZ JOHN J ET AL: "Is epsilon-aminocaproic acid as effective as aprotinin in reducing bleeding with cardiac surgery? A meta-analysis." CIRCULATION, vol. 99, no. 1, pages 81-89, XP002231358 ISSN: 0009-7322 abstract	1-46
Y	--- WO 98 58661 A (NOVONORDISK AS ; HEDNER ULLA (SE)) 30 December 1998 (1998-12-30) abstract; claims	1-46
Y	--- US 5 866 122 A (EIBL JOHANN ET AL) 2 February 1999 (1999-02-02) column 2, line 7 - line 23	1-46
A	--- KONKLE BARBARA A ET AL: "Approaches to successful knee replacement in hemophilia patients with inhibitors." BLOOD, vol. 96, no. 11 Part 2, 16 November 2000 (2000-11-16), page 84b XP002231359 42nd Annual Meeting of the American Society of Hematology; San Francisco, California, USA; December 01-05, 2000 ISSN: 0006-4971 the whole document	1-46
	--- -/--	

INTERNATIONAL SEARCH REPORT

International Application No

PCT/DK 02/00752

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>SHETH S ET AL: "Heart transplant in a factor VIII-deficient patient with a high-titre inhibitor: perioperative management using high-dose continuous infusion factor VIII and recombinant factor VIIa." HAEMOPHILIA: THE OFFICIAL JOURNAL OF THE WORLD FEDERATION OF HEMOPHILIA. ENGLAND MAR 2001, vol. 7, no. 2, March 2001 (2001-03), pages 227-232, XP002231360 ISSN: 1351-8216 the whole document</p>	1-46
P,X	<p>--- RIVERA ENID ET AL: "Use of recombinant factor VII in Hermansky-Pudlak syndrome." BLOOD, vol. 98, no. 11 Part 2, 16 November 2001 (2001-11-16), page 63b XP002231361 43rd Annual Meeting of the American Society of Hematology, Part 2;Orlando, Florida, USA; December 07-11, 2001, November 16, 2001 ISSN: 0006-4971 abstract</p>	1-46
P,X	<p>--- WO 01 85198 A (PETERSEN LARS CHRISTIAN ;NOVO NORDISK AS (DK); ROEJKJAER RASMUS (D) 15 November 2001 (2001-11-15) claims page 10, line 7 - line 8 -----</p>	1-46

INTERNATIONAL SEARCH REPORT

international application No.
PCT/DK 02/00752

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: **18-25 and 38-45**
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210

2. Claims Nos.: **1-26 (all partially)**
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
- No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Claims Nos.: 18-25 and 38-45

Claims 18-25 and 38-45 relate to methods of treatment of the human or animal body by surgery or by therapy/diagnostic methods practised on the human or animal body/Rule. 39.1.(iv)). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds/compositions.

Continuation of Box I.2

Claims Nos.: 1-26 (all partially)

Present claims 1 and 3-26 relate to an extremely large number of possible polypeptides, designated "factor VII-related polypeptides". Furthermore, claim 2 relates to an extremely large number of possible variants, designated "factor VII amino acid sequence variants".

In fact, the claims contain so many possible polypeptides and variants that a lack of clarity within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible.

Consequently, the search has mainly been focused on the general aspect of combining factor VII and epsilon-aminocaproic acid, as well as on the example in the description.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/DK 02/00752

Patent document cited in search report	A	Publication date	Patent family member(s)	Publication date
WO 9306855	A	15-04-1993	AU 2793292	A 03-05-1993
			CA 2121028	A1 15-04-1993
			CZ 9400831	A3 16-11-1994
			WO 9306855	A1 15-04-1993
			EP 0613377	A1 07-09-1994
			FI 941628	A 08-06-1994
			HU 67693	A2 28-04-1995
			JP 7500095	T 05-01-1995
			NO 941285	A 07-06-1994
			US 2002192271	A1 19-12-2002
WO 9858661	A	30-12-1998	AU 7907398	A 04-01-1999
			WO 9858661	A1 30-12-1998
US 5866122	A	02-02-1999	AT 404673	B 25-01-1999
			AT 407116	B 27-12-2000
			AT 408612	B 25-01-2002
			AT 408947	B 25-04-2002
			AT 17797	A 15-09-2001
			AT 51896	A 15-06-1998
			AU 725442	B2 12-10-2000
			AU 1645197	A 25-09-1997
			CA 2200394	A1 20-09-1997
			CZ 9700819	A3 15-10-1997
			EP 0796623	A2 24-09-1997
			HU 9700603	A2 28-06-1999
			JP 10045620	A 17-02-1998
			US 6039945	A 21-03-2000
			US 6099837	A 08-08-2000
			US 6165974	A 26-12-2000
			US 6224862	B1 01-05-2001
AT 157396	A 15-05-2000			
AT 167396	A 15-06-2001			
WO 0185198	A	15-11-2001	AU 5822801	A 20-11-2001
			AU 5823001	A 20-11-2001
			BR 0110659	A 11-02-2003
			WO 0185198	A1 15-11-2001
			WO 0185199	A1 15-11-2001
			EP 1282438	A1 12-02-2003
			EP 1282439	A1 12-02-2003
			NO 20025355	A 09-01-2003