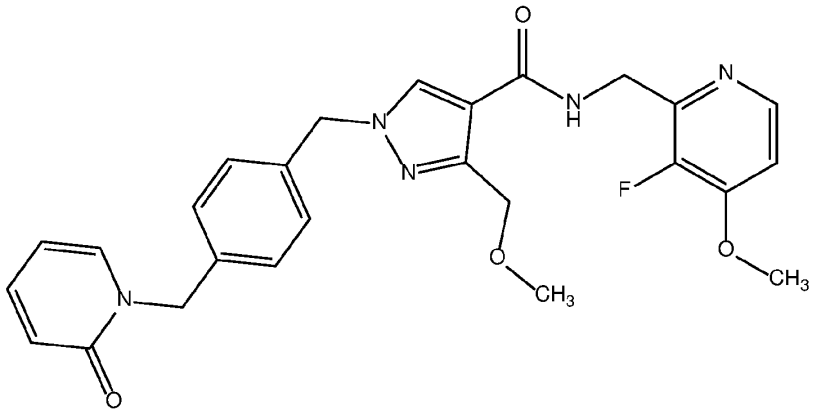
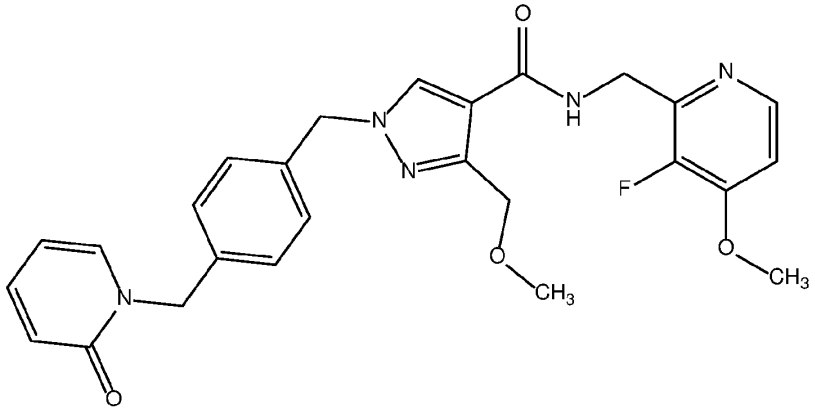
Claims

1. A method for treating bradykinin-mediated angioedema non-hereditary (BK-AEnH) on-demand comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof on-demand,



Formula A.

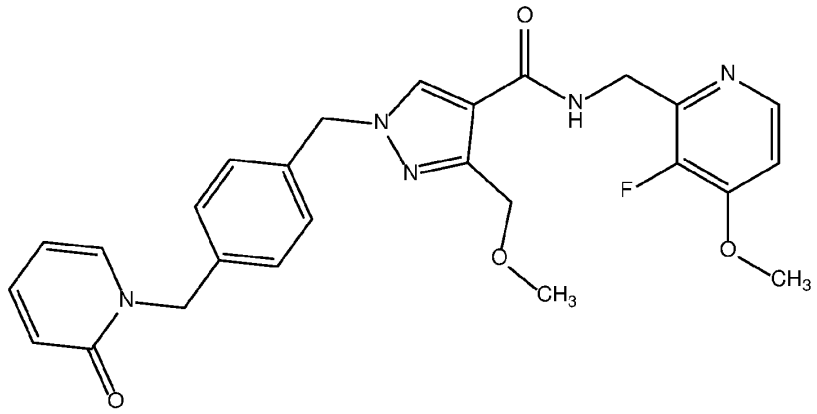
2. The compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use in treating bradykinin-mediated angioedema non-hereditary (BK-AEnH) on-demand comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof on-demand,



Formula A.

3. The use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) in the manufacture of a medicament for the treatment of bradykinin-mediated angioedema non-

hereditary (BK-AEnH) on-demand comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof on-demand,



Formula A.

4. The method according to claim 1, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 2, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 3,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is for use in treating an acute attack of bradykinin-mediated angioedema non-hereditary (BK-AEnH) on-demand and is orally administered on-demand upon recognition of a symptom of an acute BK-AEnH attack.

5. The method according to claim 4, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 4, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 4,

wherein the symptom of an acute BK-AEnH attack recognised is at least one of: swelling of tissues; fatigue; headache; muscle aches; skin tingling; abdominal pain; nausea; vomiting; diarrhoea; difficulty swallowing; hoarseness; shortness of breath; and/or mood changes.

6. The method according to any of claims 4 or 5, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 4 or 5, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 4 to 5,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand within 1 hour of the symptom of an acute BK-AEnH attack being recognised.

7. The method according to any of claims 4 to 6, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 4 to 6, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 4 to 6,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand within 30 minutes, within 20 minutes, within 10 minutes, or within 5 minutes of the symptom of an acute BK-AEnH attack being recognised.

8. The method according to any of claims 4 to 7, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 4 to 7, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 4 to 7,

wherein the compound of Formula A (or a pharmaceutically acceptable salt or solvate thereof) is orally administered on-demand in the prodromal phase of an acute BK-AEnH attack.

9. The method according to claim 8, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 8, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 8,

wherein the symptom recognised is at least one of: a slight swelling, abdominal pain or reddening of the skin.

10. The method according to claim 9, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 9, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 9,

wherein the symptom recognised is erythema marginatum.

11. The method according to any of claims 1 and 4 to 10, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 2 and 4 to 10, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 3 to 10,

wherein the treatment shortens the duration of the acute BK-AEnH attack.

12. The method according to claim 8, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 8, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 8,

wherein the treatment prevents the acute BK-AEnH attack from progressing to the swelling stage of an acute BK-AEnH attack.

13. The method according to claim 1, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 2, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 3,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand to prophylactically reduce the likelihood of an acute BK-AEnH attack.

14. The method according to claim 13, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 13, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 13,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand when it is anticipated that an acute BK-AEnH attack will be induced.

15. The method according to any of claims 13 to 14, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 13 to 14, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 13 to 14,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand to prevent an acute BK-AEnH attack.

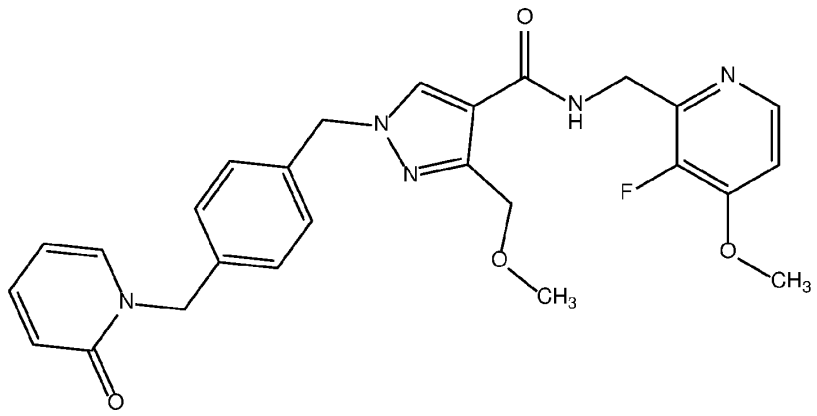
16. The method according to any of claims 14 to 15, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 14 to 15, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 14 to 15,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered on-demand when it is anticipated that an acute BK-AEnH attack will be induced by physical traumata and/or stress.

17. The method according to claim 16, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 16, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 16,

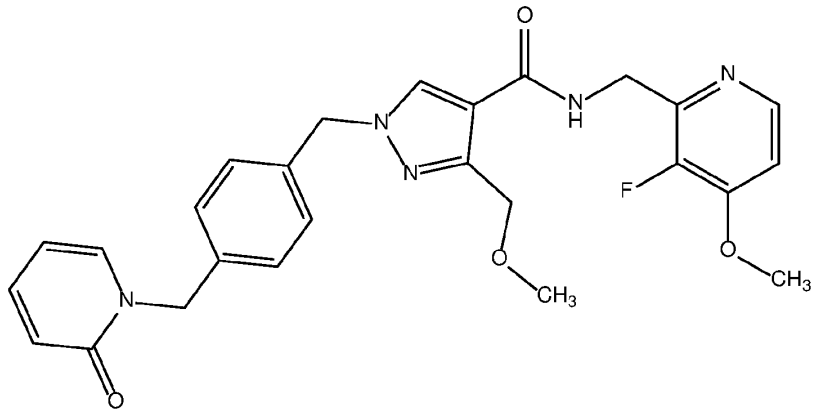
wherein it is anticipated that an acute BK-AEnH attack will be induced by the physical traumata of a dental procedure and/or the mental stress associated with a dental procedure.

18. A method for treating bradykinin-mediated angioedema non-hereditary (BK-AEnH) comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered prophylactically to reduce the likelihood of an acute BK-AEnH attack, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered regularly to the patient,



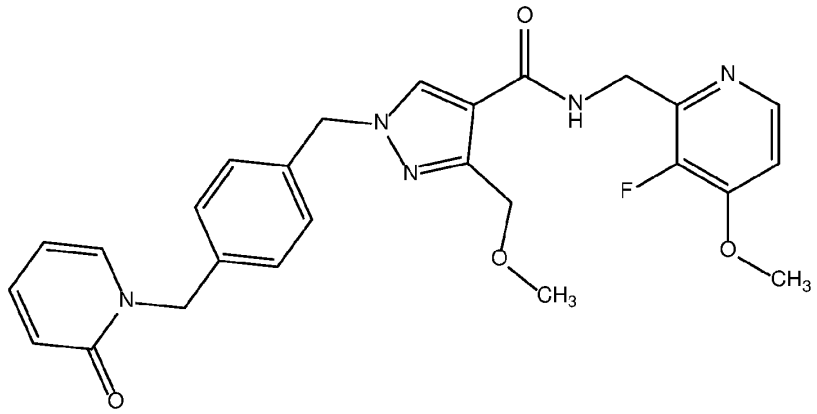
Formula A.

19. The compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use in treating bradykinin-mediated angioedema non-hereditary (BK-AEnH) comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered prophylactically to reduce the likelihood of an acute BK-AEnH attack, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered regularly to the patient,



Formula A.

20. The use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) in the manufacture of a medicament for the treatment of bradykinin-mediated angioedema non-hereditary (BK-AEnH) comprising: orally administering the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) to a patient in need thereof, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered prophylactically to reduce the likelihood of an acute BK-AEnH attack, wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered regularly to the patient,



Formula A.

21. The method according to claim 18, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 19, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 20,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered to prevent an acute BK-AEnH attack.

22. The method according to any of claims 18 or 21, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 19 or 21, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 20 to 21,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered once daily.

23. The method according to any of claims 18 or 21, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 19 or 21, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 20 to 21,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered twice daily.

24. The method according to any of claims 18 or 21, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 19 or 21, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 20 to 21,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is orally administered three times daily.

25. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the compound (or a pharmaceutically acceptable salt and/or solvate thereof) is administered as an oral dosage form comprising: (i) the compound (or a pharmaceutically acceptable salt and/or solvate thereof), and (ii) pharmaceutically acceptable excipients.

26. The method according to claim 25, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 25, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 25,

wherein the oral dosage form is a tablet comprising microcrystalline cellulose as a diluent, croscarmellose sodium as a disintegrant, polyvinyl pyrrolidone as a binder, and optionally magnesium stearate as a lubricant.

27. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the compound (or a pharmaceutically acceptable salt and/or solvate thereof) (i) inhibits plasma kallikrein, (ii) reduces cleavage of plasma prekallikrein, and/or (iii) reduces the generation of Factor XIIa from Factor XII.

28. The method according to claim 27, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 27, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 27,

wherein the patient is administered a dose of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) such that the patient's plasma has a concentration of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) of at least 500 ng/mL.

29. The method according to claim 28, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 28, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 28,

wherein the patient is administered at least 60 mg of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof).

30. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) blocks contact system activation for up to six hours.

31. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound

of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered at a daily dosage amount of between 5 mg and 2000mg.

32. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the compound of Formula A is administered at a daily dosage amount of between 100 mg and 1500 mg, between 300 mg to 1800 mg, between 100 mg and 1400 mg per day, between 200 mg and 1200 mg, between 300 mg and 1200 mg, between 600 mg and 1200 mg, between 450 mg and 900 mg, between 500 mg and 1000 mg, between 450 mg and 600 mg, between 500 mg and 700 mg, between 800 mg and 1000 mg per day, between 900 mg and 1400 mg, or between 900 mg and 1200 mg.

33. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the patient is administered the daily dosage amount in two dosage amounts within a 24 hour period starting from the time of taking the first dosage amount.

34. The method according to claim 33, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 33, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 33,

wherein the two dosage amounts are administered simultaneously, separately or sequentially.

35. The method according to any of claims 35 or 36, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 35 or 36, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 35 or 36,

wherein the second dosage amount is administered between 2 and 6 hours of the first, preferably between about 3 and 6 hours of the first dosage amount.

36. The method according to any of claims 33 or 34, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 33 or 34, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 33 or 34,

wherein the second dosage amount can be administered at least about 6 hours after the first dosage amount.

37. The method according to any of claims 1, 4 to 18, or 21 to 32, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 2, 4 to 17, 19, or 21 to 32, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 3 to 17, or 20 to 32,

wherein the patient is administered the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) as three dosage amounts per day.

38. The method according to claim 37, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 37, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 36,

wherein the three dosage amounts are administered simultaneously, separately or sequentially.

39. The method according to any of claims 37 or 38, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 37 or 38, or the use of the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 37 or 38,

wherein the second and third dosage amounts can be administered at least about 6 hours after the preceding dosage amount.

40. The method according to any of claims 30 to 39, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 30 to 39, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 30 to 39,

wherein each dosage amount comprises about 600 mg of the compound of formula A.

41. The method according to claim 40, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 40, or the use according to any claim 40,

wherein each dosage amount is administered as two tablets each comprising about 300 mg of the compound of formula A.

42. The method according to any preceding claim, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any preceding claim, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any preceding claim,

wherein the bradykinin-mediated angioedema non-hereditary (BK-AEnH) is not caused by an inherited genetic dysfunction, fault, or mutation.

43. The method according to claim 42, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 42, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 42,

wherein the BK-AEnH is selected from: non-hereditary angioedema with normal C1 Inhibitor (AE-nC1 Inh), which can be environmental, hormonal, or drug-induced; acquired angioedema; anaphylaxis associated angioedema; angiotensin converting enzyme (ACE) inhibitor-induced angioedema; dipeptidyl peptidase-4 inhibitor-induced angioedema; and tPA-induced angioedema (tissue plasminogen activator-induced angioedema).

44. The method according to claim 42, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to claim 42, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to claim 42,

wherein, the AE-nC1 Inh is environmentally-induced by air pollution and/or silver nanoparticles.

45. The method according to any of claims 13 to 15, the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) for use according to any of claims 13 to 15, or the use of a compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) according to any of claims 13 to 15,

wherein the BK-AEnH is tPA-induced angioedema,

wherein the patient is also being administered a tissue plasminogen activator,

wherein the compound of Formula A (or a pharmaceutically acceptable salt and/or solvate thereof) is administered prior to, during, or after administration of the tissue plasminogen activator to the patient.