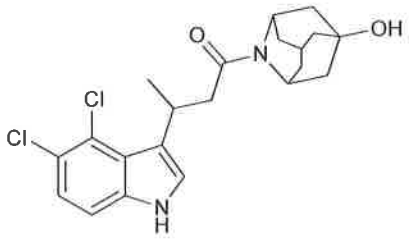
The claims defining the invention are as follows

1. Use of a compound of Formula (I), or a pharmaceutically acceptable salt thereof in the manufacture of a medicament for the treatment of a wound in a subject:

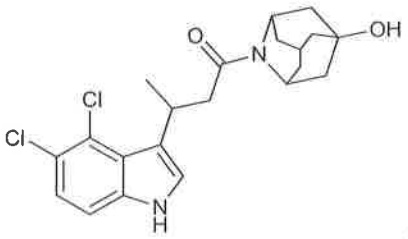


2. The use according to claim 1, wherein the wound is an incision or an excision.

3. The use according to claim 1 or claim 2, wherein the medicament is adapted to be administered topically to the site of the wound.

4. The use according to any one of claims 1 to 3, wherein the medicament is in the form of a balm, a cream, a gel, a liniment, a lotion, an ointment, a paste, a rub or a salve.

5. A compound of Formula (I), or a pharmaceutically acceptable salt thereof for use in treating a wound in a subject:

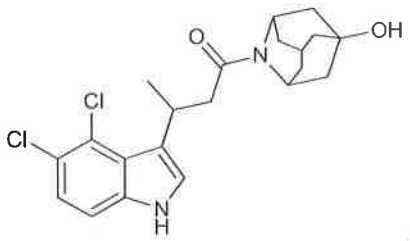


6. The compound for use according to claim 5, wherein the wound is an incision or an excision.

7. The compound for use according to claim 5 or claim 6, wherein the compound is administered topically to the site of the wound.

8. The compound for use according to any one of claims 5 to 7, wherein the compound is administered in the form of a balm, a cream, a gel, a liniment, a lotion, an ointment, a paste, a rub or a salve.

9. An agent for use in treating a wound in a subject, wherein the agent comprises a compound of Formula (I), or a pharmaceutically acceptable salt thereof:

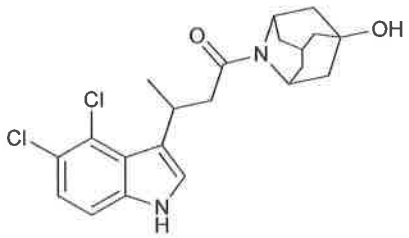


10. The agent for use according to claim 9, wherein the wound is an incision or an excision.

11. The agent for use according to claim 9 or claim 10, wherein the agent is adapted to be administered topically to the site of the wound.

12. The agent for use according to any one of claims 9 to 11, wherein the agent is a balm, a cream, a gel, a liniment, a lotion, an ointment, a paste, a rub or a salve.

13. A pharmaceutical composition for use in treating a wound in a subject, comprising a compound of Formula (I), or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable diluent, excipient or carrier:



14. The pharmaceutical composition according to claim 13, wherein the composition comprises from 0.1 to 2% (w/w) of a compound of Formula (I), or a pharmaceutically acceptable salt thereof and from 99.9 to 98% (w/w) of a carrier.

15. The pharmaceutical composition according to claim 14, wherein the compound, or a pharmaceutically acceptable salt thereof is present in the range of 0.1 to 1% (w/w), and the carrier is present in the range 99.9 to 99% (w/w).

16. The pharmaceutical composition according to claim 15, wherein the compound, or a pharmaceutically acceptable salt thereof is present at about 0.5% (w/w), and the carrier is present at about 99.5% (w/w).

17. The pharmaceutical composition according to any one of claims 14 to 16, wherein the composition comprises 0.1 to 2% (w/w) of a compound of Formula (I), or a pharmaceutically acceptable salt thereof; one or more emollients at about 5 to about 10% (w/w); one or more surfactants at about 10 to about 15% (w/w); one or more humectants at about 1 to about 10% (w/w); one or more preservatives at about 1 to about 5% (w/w); one or more antioxidants at about 0.1 to about 2% (w/w); and a solvent making the remaining balance up to 100% (w/w).

18. The pharmaceutical composition according to claim 17, wherein the one or more emollients is selected from the group consisting of heavy liquid paraffin, kokum butter and cetyl alcohol.

19. The pharmaceutical composition according to claim 17 or claim 18, wherein the one or more surfactants is selected from the group consisting of glyceryl monostearate, stearic acid, polyethylene glycol, cetostearyl alcohol and ethylene glycol monostearate.

20. The pharmaceutical composition according to any one of claims 17 to 19, wherein the one or more humectants is glycerine.

21. The pharmaceutical composition according to any one of claims 17 to 20, wherein the one or more preservatives is selected from the group consisting of phenoxy ethanol, potassium sorbate and sodium benzoate.

22. The pharmaceutical composition according to any one of claims 17 to 21, wherein the one or more antioxidants is vitamin E.

23. The pharmaceutical composition according to any one of claims 17 to 22, wherein the solvent is water.

24. The pharmaceutical composition according to claim 13 wherein the composition comprises 0.5% (w/w) of a compound of Formula (I), or a pharmaceutically acceptable salt thereof; heavy liquid paraffin - 2.5% (w/w); glyceryl mono stearate - 1.85% (w/w);stearic acid - 2.0% (w/w); kokum butter - 2.0% (w/w); polyethylene glycol - 5.0% (w/w); glycerine - 5.0% (w/w); cetostearyl alcohol - 2.5% (w/w); cetyl alcohol - 2.0% (w/w); ethylene glycol monostearate - 1.0% (w/w); phenoxy ethanol - 0.5% (w/w); potassium sorbate - 0.5% (w/w); sodium benzoate - 0.5% (w/w); vitamin E - 0.5% (w/w); and purified water - 73.65% (w/w).