WHAT IS CLAIMED IS:

- A method for treating pulmonary hypertension, comprising administering to a subject in need thereof treprostinil or treprostinil derivative, or a pharmaceutically acceptable salt thereof by a metered dose inhaler.
- 2. The method of claim 1, wherein said metered dose inhaler is a pressured metered dose inhaler.
- 3. The method of claim 1, wherein said metered dose inhaler is a dry powder inhaler.
- 4. The method of claim 1, wherein said metered dose inhaler is a soft mist inhaler.
- 5. The method of claim 4, wherein said treprostinil is formulated in said inhaler as a solution, wherein a solvent of the solution comprises water, ethanol or a mixture thereof.
- 6. The method of claim 5, wherein a concentration of the treprostinil in the solution ranges from about 500 μ g/ml to about 2500 μ g/ml.
- 7. The method of claim 6, wherein the concentration of the treprostinil in the solution ranges from about 1000 μ g/ml to about 2000 μ g/ml.
- 8. The method of claim 1, wherein a dose of the treprostinil administered during a single event ranges from about 15 µg to about 100 µg of the treprostinil.
- 9. The method of claim 8, wherein the dose ranges from about 30 μg to about 90 μg of the treprostinil.



- 10. The method of claim 1, wherein said administering does not have a systemic side effect on said subject, wherein the systemic side effect is selected from the group consisting of headache, flush, nausea, and dizziness.
- 11. The method of claim 1, wherein said administering does not disrupt gas exchange in said subject.
- 12. The method of claim 1, wherein said administering does change heart rate of said subject.
- 13. The method of claim 1, wherein said administering does not affect systemic arterial pressure and systemic arterial resistance.
- 14. The method of claim 1, wherein said administering comprises a limited number of breaths by said subject.
- 15. The method of claim 1, wherein said administering lasts less than 5 minutes.
- 16. The method of claim 1, wherein said administering lasts less than 1 minute.
 - 17. The method of clam 1, wherein said subject is a human being.
- 18. The method of claim 1, further comprising administering to said subject at least one supplementary agent selected from the group consisting of diltiazem, amlodipine, nifedipine, sildenafil, tadalafil, vardenafil, bosentan, sitaxsentan, ambrisenatn, prostacyclin, iloprost, beraprost and pharmaceutically acceptable salts thereof.



- 19. A method of delivering to a subject in need thereof a therapeutically effective amount of treprostinil, or treprostinil derivative or a pharmaceutically acceptable salt thereof comprising administering to the subject the therapeutically effective amount of the treprostinil or treprostinil derivative or a pharmaceutically acceptable salt thereof using a metered dose inhaler.
- 20. The method of claim 19, wherein said metered dose inhaler is a pressured metered dose inhaler.
- 21. The method of claim 19, wherein said metered dose inhaler is a dry powder inhaler.
- 22. The method of claim 19, wherein said metered dose inhaler is a soft mist inhaler.
- 23. The method of claim 22, wherein said treprostinil is formulated in the metered dose inhaler as a solution, wherein a solvent of the solution comprises water, ethanol or a mixture thereof.
- 24. The method of claim 23, wherein a concentration of the treprostinil in the solution ranges from about 500 μ g/ml to about 2500 μ g/ml.
- 25. The method of claim 24, wherein the concentration of the treprostinil in the solution ranges from about 1000 μ g/ml to about 2000 μ g/ml.
- 26. The method of claim 19, wherein a dose of the treprostinil administered during a single event ranges from about 15 μ g to about 100 μ g of the treprostinil.



- 27. The method of claim 26, wherein the dose ranges from about 30 μg to about 90 μg of the treprostinil.
- 28. The method of claim 19, wherein said administering does not have a systemic side effect on said subject, wherein the systemic side effect is selected from the group consisting of headache, flush, nausea, and dizziness.
- 29. The method of claim 19, wherein said administering does not disrupt gas exchange in said subject.
- 30. The method of claim 19, wherein said administering does change heart rate of said subject.
- 31. The method of claim 19, wherein said administering does not affect systemic arterial pressure and systemic arterial resistance.
- 32. The method of claim 19, wherein said administering comprises a limited number of breaths by said subject.
- 33. The method of claim 19, wherein said administering lasts less than 5 minutes.
- 34. The method of claim 19, wherein said administering lasts less than 1 minute.
 - 35. The method of clam 19, wherein said subject is a human being.
- 36. A kit for treating pulmonary hypertension, comprising (i) a metered dose inhaler containing a pharmaceutical formulation comprising treprostinil or



treprostinil derivative, or a pharmaceutically acceptable salt thereof; and (ii) instructions for use of in treating pulmonary hypertension.

- 37. The kit of claim 36, wherein said metered dose inhaler is a pressured metered dose inhaler.
- 38. The kit of claim 36, wherein said metered dose inhaler is a dry powder inhaler.
- 39. The kit of claim 36, wherein said metered dose inhaler is a soft mist inhaler.
- 40. The kit of claim 39, wherein said formulation further comprises water, ethanol or a mixture thereof.
- 41. The kit of claim 36, wherein a concentration of the treprostinil in said formulation is from about 500 μ g/ml to about 2500 μ g/ml.
- 42. The kit of claim 41, wherein said concentration is from about 1000 $\mu g/ml$ to about 2000 $\mu g/ml$.
- 43. The kit of claim 36, further comprising an effective amount of at least one supplementary agent selected from the group consisting of diltiazem, amlodipine, nifedipine, sildenafil, tadalafil, vardenafil, bosentan, sitaxsentan, ambrisenatn, prostacyclin, iloprost, beraprost and pharmaceutically acceptable salts thereof.
- 44. A kit comprising a metered dose inhaler containing a pharmaceutical formulation comprising treprostinil or treprostinil derivative, or a pharmaceutically acceptable salt thereof.



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