## EXHIBIT 9

#### AMENDMENTS TO THE CLAIMS

### Listing of Claims:

- (Currently Amended) A pharmaceutical formulation which comprises azelastine, or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof, and fluticasone or a pharmaceutically acceptable ester thereofa steroid, or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof, which contains the fluticasone or a pharmaceutically acceptable ester thereof in an amount from about 50 micrograms/ml to about 5 mg/ml of the formulation.
- 2. A pharmaceutical formulation according to claim 1, wherein said azelastine (Original) is present as azelastine hydrochloride.
- 3. (Canceled)
- (Currently Amended) A formulation according to elaim 3claim 1, wherein the steroid pharmaceutically acceptable ester is beclomethasone propionate, mometasonefuroate, mometasone furoate monohydrate, fluticasone propionate or fluticasone valerate.
- 5. (Canceled)
- (Currently Amended) A formulation according to claim 1, wherein the formulation has a 6. particle size of less than about 10 μm.



- 7. (Currently Amended) A formulation according to claim 1, which is a suspension containing 0.0005 to 2% (weight/weight of the formulation) of azelastine or a pharmaceutically acceptable salt of azelastine, and from 0.5 to 1.5% (weight/weight of the formulation) of fluticasone or a pharmaceutically acceptable ester thereofsaid steroid.
- 8. (Currently Amended) A formulation according to claim 7, which contains from 0.001 to 1% (weight/weight of the formulation) azelastine, or salt thereof, and from 0.5% to 1.5% (weight/weight of the formulation) <u>fluticasone</u> or a <u>pharmaceutically acceptable ester</u> thereofsteroid.
- 9. (Previously Presented) A formulation according to claim 1, which also contains a surfactant.
- 10. (Original) A formulation according to claim 9, wherein the surfactant comprises a polysorbate or poloxamer surfactant.
- 11. (Previously Presented) A formulation according to claim 9, which contains from about 50 micrograms to about 1 milligram of surfactant per ml of the formulation.
- 12. (Previously Presented) A formulation according to claim 1, which also contains an isotonic agent.



- 13. (Original) A formulation according to claim 12, wherein the isotonic agent comprises sodium chloride, saccharose, glucose, glycerine, sorbitol or 1,2-propylene glycol.
- 14. (Previously Presented) A formulation according to claim 1, which also contains at least one additive selected from the group consisting of a buffer, a preservative, a suspending agent and a thickening agent.
- 15. (Original) A formulation according to claim 14, wherein said preservative is selected from edetic acid and its alkali salts, lower alkyl p-hydroxybenzoates, chlorhexidine, phenyl mercury borate, or benzoic acid or a salt, a quaternary ammonium compound, or sorbic acid or a salt thereof.
- 16. A formulation according to claim 14, wherein the (Previously Presented) suspending agent or thickening agent is selected from cellulose derivatives, gelatin, polyvinylpyrrolidone, tragacanth, ethoxose (water soluble binding and thickening agents on the basis of ethyl cellulose), alginic acid, polyvinyl alcohol, polyacrylic acid, or pectin.
- 17. (Previously Presented) A formulation according to claim 14, wherein the buffer comprises a citric acid-citrate buffer.
- (Currently Amended) A formulation according to claim 14, wherein the buffer maintains 18. the pH of the aqueous phase at from 3 to 7, preferably 4.5 to about 6.5.

- 19. (Previously Presented) A formulation according to claim 1, which is an aqueous suspension or solution.
- 20. (Previously Presented) A formulation according to claim 1, which is in the form of an aerosol, an ointment, eye drops, nasal drops, a nasal spray, an inhalation solution and other forms suitable for nasal or ocular administration.
- 21. (Original) A formulation according to claim 20, which is in the form of nasal drops or nasal spray.
- 22. (Original) A formulation according to claim 20, which is in the form of an aerosol.
- 23-24. (Canceled)
- 25. (Previously Presented) A formulation according to claim 1, which is in the form of an insufflation powder.
- 26. (Currently Amended) A pharmaceutical product—according to claim—1, comprising (i) azelastine, or a pharmaceutically acceptable salt, solvate or physiologically functional derivative thereof, provided in an aerosol formulation preferably together with a propellant typically suitable for MDI delivery, and (ii) <u>fluticasone</u> or a <u>pharmaceutically acceptable ester thereofat least one steroid</u>, or a <u>pharmaceutically acceptable salt</u>, solvate or <u>physiologically functional derivative</u>



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