~~[CLAIMS]~~

~~[Claim 1]~~

1. A pharmaceutical composition comprising:

fimasartan, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate thereof;

hydrochlorothiazide, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate thereof; and

a binder exhibiting a viscosity of 20 mPa·s to 1,000 mPa·s at 25 °C.

~~[Claim 2]~~

2. The pharmaceutical composition of claim 1, wherein the binder comprises at least one selected from the group consisting of hydroxypropyl cellulose, hydroxypropyl methylcellulose, and polyvinylpyrrolidone.

~~[Claim 3]~~

3. The pharmaceutical composition of claim 1, wherein the pharmaceutical composition have a relative standard deviation (RSD) of 4.0% or less in content uniformity test.

~~[Claim 4]~~

4. The pharmaceutical composition of claim 1,

wherein the pharmaceutical composition is in the form of a tablet.

~~[Claim 5]~~

5. The pharmaceutical composition of claim 4, wherein the tablet comprises an uncoated tablet and a coating layer.

~~[Claim 6]~~

6. The pharmaceutical composition of claim 5, wherein the uncoated tablet has a hardness of 7 Kp or more.

~~[Claim 7]~~

~~The pharmaceutical composition of claim 1, wherein the pharmaceutical composition is used for the treatment of hypertension.~~

~~[Claim 8]~~

7. A method for preparing a pharmaceutical composition comprising fimasartan, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate thereof and hydrochlorothiazide, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate

thereof, the method comprising the steps of:

preparing a mixture comprising fimasartan, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate thereof and hydrochlorothiazide, a pharmaceutically acceptable salt thereof, a hydrate thereof, or a solvate thereof;

preparing granules by mixing the mixture with a binding solution in which a binder is dissolved to exhibit a viscosity of 20 mPa·s to 1,000 mPa·s at 25 °C; and

preparing an uncoated tablet containing the granules.

~~[Claim 9]~~

8. The method of claim ~~8~~7, further comprising the steps of:

preparing sized materials by sizing the granules;

preparing a final mixture by adding one or more additives to the sized materials; and

preparing the uncoated tablet by compressing the final mixture.

~~[Claim 10]~~

9. The method of claim ~~8~~7, wherein the binder comprises at least one selected from the group consisting of hydroxypropyl cellulose, hydroxypropyl methylcellulose, and polyvinylpyrrolidone.

~~【Claim 11】~~

10. The method of claim ~~8~~7, wherein the pharmaceutical composition have a relative standard deviation (RSD) of 4.0% or less in content uniformity test.

~~【Claim 12】~~

11. The method of claim ~~8~~7, wherein the uncoated tablet has a hardness of 7 Kp or more.

~~【Claim 13】~~

~~The method of claim 8, wherein the pharmaceutical composition is used for the treatment of hypertension.~~

~~【Claim 14】~~

12. A tablet prepared by the method of any one of claims ~~8~~7 to ~~13~~11.