CLAIMS

1. A pharmaceutical combination preparation comprising a first layer comprising telmisartan or a pharmaceutically acceptable salt thereof, microcrystalline cellulose, calcium silicate, and an alkalizing agent and a second layer comprising (S)-amlodipine or a pharmaceutically acceptable salt thereof,

   wherein the weight ratio of telmisartan or a pharmaceutically acceptable salt thereof to calcium silicate is 10:1 to 10:5, and the alkalizing agent comprises at least one selected from the group consisting of basic amino acids, sodium carbonate, potassium carbonate, sodium hydrogen carbonate, sodium hydrogen phosphate, and mixtures thereof.

2. The pharmaceutical combination preparation of claim 1, wherein the weight ratio of telmisartan or a pharmaceutically acceptable salt thereof to microcrystalline cellulose is 10:25 to 10:35.

3. The pharmaceutical combination preparation of claim 1, wherein the weight ratio of telmisartan or a pharmaceutically acceptable salt thereof to microcrystalline cellulose is 10:25 to 10:30 and the weight ratio of telmisartan or a pharmaceutically acceptable salt thereof to calcium silicate is 10:1 to 10:5.

4. The pharmaceutical combination preparation of claim 3, wherein the microcrystalline cellulose is contained in an amount of 50 to 60 wt% with respect to the total weight of the first layer.

5. The pharmaceutical combination preparation of claim 3, wherein the weight ratio of telmisartan or a pharmaceutically acceptable salt thereof to calcium silicate is 10:2 to 10:3.

6. The pharmaceutical combination preparation of claim 1, wherein the pharmaceutically acceptable salt of (S)-amlodipine is selected from the group consisting of (S)-

amlodipine besylate, (S)-amlodipine maleate, (S)-amlodipine orotate, (S)-amlodipine camsylate, (S)-amlodipine adipate, and (S)-amlodipine mesylate.