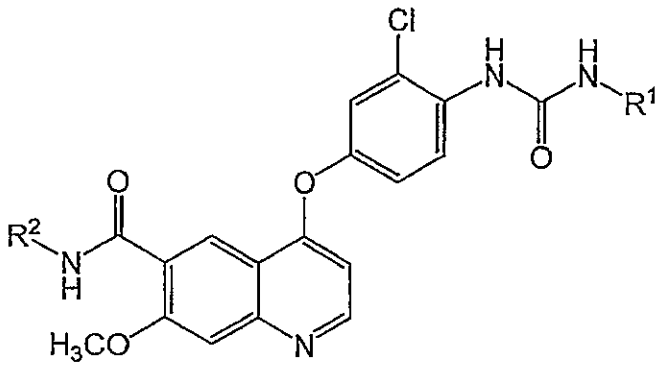
CLAIMS

1. A pharmaceutical composition comprising

(1) a compound represented by the formula (I) or pharmaceutically acceptable salt thereof or solvate thereof:



(I)

wherein R^1 is a hydrogen atom, a C_{1-6} alkyl group or a C_{3-8} cycloalkyl group; and R^2 represents a hydrogen atom or a methoxy group; and

(2) a basic substance.

2. The composition according to claim 1, wherein the basic substance is a carbonate.

3. The composition according to claim 2, wherein the salt is an alkaline earth metal salt.

4. The composition according to claim 3, wherein the alkaline earth metal salt is a magnesium salt or a calcium salt.

5. The composition according to any one of claims 1 to 4, further comprising a disintegrating agent.

6. The composition according to claim 5, wherein the disintegrating agent is carmellose sodium, carmellose calcium, carboxymethyl starch sodium, croscarmellose sodium, low-substituted hydroxypropylcellulose or crospovidone.

7. The composition according to any one of claims 1 to 6, wherein R^1 is a hydrogen atom, a methyl group, an ethyl group, an n-propyl group or a cyclopropyl group.

8. The composition according to any one of claims 1 to 7, wherein R^1 is a cyclopropyl group.

9. The composition according to any one of claims 1 to 8, wherein R^2 is a hydrogen atom, a methoxy group or an ethoxy group.

10. The composition according to any one of claims 1 to 9, wherein R^2 is a hydrogen atom.

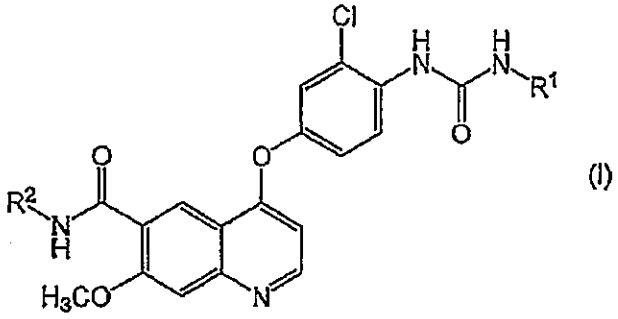
11. The composition according to any one of claims 1 to 10, wherein the pharmaceutically acceptable salt is hydrochloride, hydrobromide, p-toluenesulfonate, sulfate, methanesulfonate or ethanesulfonate.

12. The composition according to any one of claims 1 to 11, wherein the compound represented by the formula (I) is 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide methanesulfonate.

CLAIMS

1. A pharmaceutical composition comprising

(1) a compound represented by the formula (I) or pharmaceutically acceptable salt thereof or solvate thereof:



(I)

wherein R^1 is a hydrogen atom, a C_{1-6} alkyl group or a C_{3-8} cycloalkyl group; and R^2 represents a hydrogen atom or a methoxy group; and

(2) an alkaline earth metal carbonate.

2. The composition according to claim 1, wherein the alkaline earth metal carbonate is magnesium carbonate or calcium carbonate.

3. The composition according to claim 1 or 2, further comprising a disintegrating agent.

4. The composition according to claim 3, wherein the disintegrating agent is carmellose sodium, carmellose calcium, carboxymethyl starch sodium, croscarmellose sodium, low-substituted hydroxypropylcellulose or crospovidone.

5. The composition according to any one of claims 1 to 4, wherein R^1 is a hydrogen atom, a methyl group, an ethyl group, an n-propyl group or a cyclopropyl group.

6. The composition according to any one of claims 1 to 5, wherein R^1 is a cyclopropyl group.

7. The composition according to any one of claims 1 to 6, wherein R^2 is a hydrogen atom.

8. The composition according to any one of claims 1 to 7, wherein the pharmaceutically acceptable salt is hydrochloride, hydrobromide, p-toluenesulfonate, sulfate, methanesulfonate or ethanesulfonate.

9. The composition according to any one of claims 1 to 8, wherein the compound represented by the formula (I) is 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide methanesulfonate.