

CLAIMS

What is claimed is:

1. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of rimegepant, or a pharmaceutically acceptable salt thereof, wherein the composition is in the form of an oral solid molded fast-dispersing dosage form.
2. The pharmaceutical composition of claim 1, comprising 75 mg of rimegepant or a pharmaceutically acceptable salt thereof.
3. The pharmaceutical composition of claim 1, wherein the rimegepant is in the form of a hemisulfate sesquihydrate salt.
4. The pharmaceutical composition of claim 1 in the form of a tablet.
5. The pharmaceutical composition of claim 1, that provides an AUC_{0-t} of from 80-125% of 5000 (hr*ng/mL).
6. The pharmaceutical composition of claim 1, comprising from 70-80 weight% rimegepant hemisulfate sesquihydrate, 10-20 weight% fish gelatin, 10-20 weight% of a filler, and 0.1-5.0 weight% of a flavorant.
7. The pharmaceutical composition of claim 1, further comprising fish gelatin as a carrier.
8. The pharmaceutical composition of claim 6, wherein the filler is mannitol.
9. The pharmaceutical composition of claim 1, wherein the pharmaceutical composition is capable of disintegrating or dispersing within an interval selected from 1 to 60 seconds, 1 to 30 seconds, 1 to 10 seconds, and 2 to 8 seconds, after being placed in contact with a fluid.
- ~~10. — A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of a CGRP receptor antagonist, or a pharmaceutically acceptable salt~~

thereof, wherein the pharmaceutical composition is in a form of an oral solid molded fast-dispersing dosage form.

11. — The pharmaceutical composition of claim 10, wherein the CGRP receptor antagonist is selected from olcegepant, telcagepant, ubrogepant, atogepant, and rimegepant.

12. — The pharmaceutical composition of claim 10, further comprising gelatin as a carrier.

13. — The pharmaceutical composition of claim 10, further comprising mannitol as a filler.

14. — The pharmaceutical composition of claim 10, comprising from 70–80 weight% of the CGRP antagonist, 10–20 weight% fish gelatin, 10–20 weight% of a filler, and 0.1–5.0 weight% of a flavorant.

15. — The pharmaceutical composition of claim 10, wherein the pharmaceutical composition is capable of disintegrating or dispersing within an interval selected from 1 to 60 seconds, 1 to 30 seconds, 1 to 10 seconds, and 2 to 8 seconds, after being placed in contact with a fluid.

16. — The pharmaceutical composition of claim 15, wherein the fluid comprises saliva.