CLAIMS

1. A pharmaceutical composition comprising VS-6063, or a pharmaceutically acceptable salt thereof, admixed with ~~an excipient, wherein the excipient is~~ polyvinylpyrrolidone vinyl acetate copolymer (PVP VA) ~~or~~and a copolymer comprising polyethylene glycol, polyvinylcaprolactam, and polyvinylacetate, wherein the composition is configured in an oral dosage form.

2. The pharmaceutical composition of claim 1, wherein the oral dosage form comprises 5 to 50%, 5 to 30%, 10 to 30%, 10 to 20%, 12 to 15%, or 13% of VS-6063, or a pharmaceutically acceptable salt thereof, per weight of the oral dosage form.

3. The pharmaceutical composition of claim 1, wherein the VS-6063, or a pharmaceutically acceptable salt thereof, is present at 13% ~~weight~~ per weight of the oral dosage form.

4. The pharmaceutical composition of claim 1, wherein the oral dosage form is a tablet.

5. The pharmaceutical composition of claim 4, wherein the tablet is provided by direct compression of a physical blend or mixture.

6. The pharmaceutical composition of claim 5, wherein the physical blend or mixture is provided by a dry granulation process.

7. The pharmaceutical composition of claim 1, further comprising a disintegrant ~~(e.g., a crosslinked polymer, e.g., crosslinked polyvinylpyrrolidone or crospovidone, carboxymethyl cellulose or croscarmellose sodium; modified starch, e.g., sodium starch glycolate)~~.

8. The pharmaceutical composition of claim 7, wherein the disintegrant is crosslinked polyvinylpyrrolidone or crospovidone, carboxymethyl cellulose or croscarmellose sodium, or a modified starch.

~~8~~9. The pharmaceutical composition of claim 7, wherein the disintegrant is a polyvinylpolypyrrolidone ~~(e.g., crospovidone)~~.

~~9~~10. The pharmaceutical composition of claim 7, wherein the disintegrant is present at 0 to 30%, 5 to 25%, 5 to 20%, 5 to 15%, or 10% per weight of the oral dosage form.

~~10~~11. The pharmaceutical composition of claim 1, further comprising a lubricant.

12. The pharmaceutical composition of claim 11, wherein the lubricant is ~~(e.g.,~~ talc, silica, or fats, ~~e.g., magnesium stearate),~~ optionally wherein the lubricant is magnesium stearate, optionally wherein the lubricant is present at 0.1 to 2%, 0.2 to 1.5%, 1%, or 0.5% per weight of the oral dosage form.

~~11. The pharmaceutical composition of claim 1, comprising a mixture of two excipients.~~

~~12~~13. The pharmaceutical composition of claim 1, further comprising one or more fillers.

~~13~~14. The pharmaceutical composition of claim ~~12~~13, wherein the filler is a polyol ~~(e.g., mannitol-starch (e.g., Pearlitol Flash), mannitol, sorbitol)~~.

15. The pharmaceutical composition of claim 14, wherein the polyol is mannitol-starch, mannitol, or sorbitol.

~~14~~16. The pharmaceutical composition of claim 1, wherein the composition comprises VS-6063 free base or VS-6063 hydrochloride.

~~15~~17. An oral dosage form comprising:

VS-6063 ~~(e.g., VS-6063 free base)~~, or a pharmaceutically acceptable salt thereof, ~~(e.g., VS-6063 hydrochloride)~~ in an amount from ~~about~~ 10 to 15% by weight on a dry weight basis,

a first pharmaceutically acceptable excipient ~~(e.g., a binder (e.g., polyvinyl pyrrolidone (e.g., polyvinyl pyrrolidone vinyl acetate copolymer (PVP/VA), Copovidone))~~ in an amount from ~~about~~ 15 to 25% by weight on a dry weight basis,

a second pharmaceutically acceptable excipient ~~(e.g., a binder (e.g., polyvinyl acetate (e.g., Soluplus)))~~ in an amount from ~~about~~ 15 to 25% by weight on a dry weight basis,

a pharmaceutically acceptable filler ~~(e.g., e.g., polyol (e.g., mannitol-starch (e.g., Pearlitol Flash)))~~ in an amount from ~~about~~ 30 to 40% by weight on a dry weight basis,

a pharmaceutically acceptable disintegrant ~~(e.g., crospovidone~~ in an amount from ~~about~~ 5 to 15% by weight on a dry weight basis, and

a pharmaceutically acceptable lubricant ~~(e.g., magnesium stearate)~~ in an amount from ~~about~~ 0.2 to 1.5% by weight on a dry weight basis.

18. The oral dosage form of claim 17, wherein the first pharmaceutically acceptable excipient is polyvinyl pyrrolidone vinyl acetate copolymer (PVP VA).

19. The oral dosage form of claim 17, wherein the second pharmaceutically acceptable excipient is a copolymer comprising polyethylene glycol, polyvinylcaprolactam, and polyvinylacetate.

20. The oral dosage form of claim 17, wherein the pharmaceutically acceptable filler is mannitol-starch.

21. The oral dosage form of claim 17, wherein the pharmaceutically acceptable disintegrant is crospovidone.

22. The oral dosage form of claim 17, wherein the pharmaceutically acceptable lubricant is magnesium stearate.