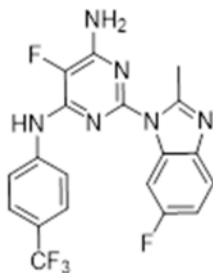


CLAIMS

We claim:

1. An amorphous solid dispersion comprising ~~An an~~ amorphous form of 5-fluoro-2-(6-fluoro-2-methyl-1H-benzimidazol-1-yl)-N-[4-(trifluoromethyl)phenyl]pyrimidine-4,6-diamine, having the formula of Compound A:



Compound A, and

- a stabilizing polymer, wherein the stabilizing polymer is hydroxypropylmethylcellulose acetate succinate.
2. The ~~form~~ dispersion of claim 1, wherein the amorphous form of Compound A is at least 90% amorphous.
 3. The ~~form~~ dispersion of claim 2, wherein the amorphous form of Compound A is a substantially pure amorphous form.
 - ~~4. An amorphous solid dispersion comprising, the amorphous form of any of claims 1, 2 or 3 and a stabilizing polymer.~~
 - ~~5. The dispersion of claim 4, wherein the stabilizing polymer is selected from polyvinylpyrrolidone or hydroxypropylmethylcellulose acetate succinate.~~
 - ~~6. The dispersion of claim 5, wherein the stabilizing polymer is hydroxypropylmethylcellulose acetate succinate.~~
 - ~~7.4.~~ The dispersion of claim ~~4~~1, wherein the amount by weight of Compound A to the amount by weight of the stabilizing polymer is in a ratio of between ~~about~~ 5:95 to ~~about~~ 60:40.
 - ~~8.5.~~ The dispersion of claim ~~7~~1, wherein the amount by weight of Compound A to the amount by weight of the stabilizing polymer is in a ratio of between ~~about~~ 10:90 to ~~about~~ 40:60.
 - ~~9.6.~~ The dispersion of claim ~~4~~1, wherein the dispersion is obtained by solvent casting of a solution comprising Compound A and the stabilizing polymer.

~~10.7.~~_____The dispersion of claim ~~4~~1, wherein the dispersion is obtained by spray drying a solution comprising Compound A and the stabilizing polymer.

~~11.8.~~_____A process for preparing the dispersion of claim ~~4~~1, comprising:

(a) dissolving Compound A and one or more stabilizing polymers in a solvent to form a solution; and,

(b) drying the solution to form the dispersion of claim ~~4~~1.

~~12.9.~~_____The process of claim ~~11~~8, wherein the solvent is selected from acetone, methyl ethyl ketone, dichloromethane, or methanol.

~~13.10.~~_____The process of claim ~~11~~8, wherein the solution is dried by solvent casting to form the dispersion of claim ~~4~~1.

~~14.11.~~_____The process of claim ~~11~~8, wherein the solution is dried by spray drying to form ~~the~~ the dispersion of claim ~~4~~1.

~~15.12.~~_____A pharmaceutical composition comprising, an admixture of the dispersion of claim ~~4~~1 and one or more pharmaceutically acceptable excipients.

~~16.13.~~_____The pharmaceutical composition of claim ~~15~~12, wherein the one or more pharmaceutically acceptable excipients are selected from a diluent, a disintegrant, a lubricant or a surfactant.

~~17.14.~~_____The pharmaceutical composition of claim ~~16~~13, wherein the surfactant is selected from sodium lauryl sulfate or Poloxamer 407.

~~18.15.~~_____A process for preparing the pharmaceutical composition of claim ~~15~~12 comprising,

(a) grinding the dispersion of claim ~~4~~1 to form a solid dispersion powder;

(b) admixing the powder with one or more pharmaceutically acceptable excipients to form an admixture;

(c) granulating the admixture to form a granulate; and,

(d) tableting or encapsulating the granulate to form tablets or capsules, respectively.