**WHAT IS CLAIMED IS:**

1. A roller compacted solid pharmaceutical composition comprising (a) the compound 4-(trifluoromethyl)-5-(2,6-dimorpholinopyrimidin-4-yl)pyridin-2-amine or a pharmaceutically acceptable salt thereof, (b) sodium stearyl fumarate, and (c) optionally at least one additional pharmaceutically acceptable carrier.

2. The roller compacted solid pharmaceutical composition according to claim 1, wherein the compound 4-(trifluoromethyl)-5-(2,6-dimorpholinopyrimidin-4-yl)pyridin-2-amine or a pharmaceutically acceptable salt thereof is present in an amount ranging from about 15% to 60% by weight of relative to the total weight of the overall composition.

3. The roller compacted solid pharmaceutical composition according to any of the preceding claims, wherein sodium stearyl fumarate is present in an amount ranging from about 3% to 8% by weight relative to the total weight of the overall composition.

4. The roller compacted solid pharmaceutical composition according to any of the preceding claims, wherein the pharmaceutical composition contains a plurality of granules having an internal phase and an external phase, and wherein said internal phase and external phase of the granules both include sodium stearyl fumarate.

5. The roller compacted solid pharmaceutical composition according to any of the preceding claims, wherein the pharmaceutical composition comprises at least one additional pharmaceutically acceptable carrier selected from a diluent, a disintegrant, a glidant, or a combination thereof.

6. The roller compacted solid pharmaceutical composition according to any of the preceding claims, wherein the pharmaceutical composition comprises at least one additional pharmaceutically acceptable carrier selected from a diluent present in an amount from about 40% to about 70% by weight of the composition, a disintegrant present in an amount from about 2% to about 15% by weight of the composition, a glidant present in an amount from about 0. 1 % to about 4% by weight of the composition, or a combination thereof.

7. The roller compacted solid pharmaceutical composition according to claim 5 or 6, wherein the diluent is selected from microcrystalline cellulose, mannitol, or a combination thereof.

8. The roller compacted solid pharmaceutical composition according to claim 5 or 6, wherein the disintegrant is crospovidone.

9. The roller compacted solid pharmaceutical composition according to claim 5 or 6, wherein the glidant is colloidal silicon dioxide.

10. The roller compacted solid pharmaceutical composition according to any of the preceding claims wherein the pharmaceutical composition shows an in vitro dissolution of the compound 4-(trifluoromethyl)-5-(2,6-dimorpholinopyrimidin-4-yl)pyridin-2-amine or a pharmaceutically acceptable salt thereof of 90% or more in 30 minutes or less.

11. The roller compacted solid pharmaceutical composition according to any of the preceding claims, wherein the solid pharmaceutical composition is in the form of a tablet.

12. A roller compacted solid pharmaceutical composition according to any of the preceding claims for use for the treatment of cancer.

13. Use of a roller compacted solid pharmaceutical composition according to any of claims 1 to 11 for the manufacture of a medicament for the treatment of cancer.

14. The roller compacted solid pharmaceutical composition for use according to claim 12, or the use according to claim 13, wherein the cancer is selected from the group consisting of lung cancer, breast cancer, prostate cancer, pancreatic cancer, colon and rectum, thyroid cancer, liver and intrahepatic bile duct cancer, hepatocellular cancer, gastric cancer, glioma / glioblastoma, endometrial cancer, melanoma, kidney and renal pelvic cancer, urinary bladder cancer, uterine corpus cancer, uterine cervix cancer, ovarian cancer, multiple myeloma, esophageal cancer, neuroendocrine tumor, leukemia, lymphoma, brain cancer, head and neck cancer, small intestinal cancer, melanoma, and villous colon adenoma.

15. A process for the manufacture of a solid pharmaceutical composition according to any of claims 1 to 11 comprising the steps of roller compacting the compound 4-(trifluoromethyl)-5-(2,6-dimorpholinopyrimidin-4-yl)pyridin-2-amine or a pharmaceutically acceptable salt thereof together with sodium stearyl fumarate and optionally at least one additional pharmaceutically acceptable carrier, milling the compacted material to form a plurality of granules, and blending the granules with sodium stearyl fumarate and optionally at least one additional pharmaceutically acceptable carrier, and optionally compressing the final blend into a tablet.