

10. ~~A method of treating breast cancer, prostate cancer or gastric cancer which comprises the steps of administering to a person in need thereof~~ Use of a therapeutically effective amount of Compound (I) in crystalline form, as claimed in any one of claims 1 to 6 in the manufacture of a medicament for treating breast cancer, prostate cancer or gastric cancer.
11. A pharmaceutical composition which comprises Compound (I) in crystalline form as claimed in any one of claims 1 to 6, in association with microcrystalline cellulose, mannitol, croscarmellose sodium and magnesium stearate.
12. A pharmaceutical tablet which comprises from 50 to 500 mg of Compound (I) in crystalline form as claimed in any one of claims 1 to 6, in association with one or more pharmaceutically acceptable excipients.
13. A pharmaceutical tablet which comprises from 0.5 to 2% by weight of magnesium stearate, from 2 to 5% by weight of croscarmellose sodium, from 15 to 60% by weight of Compound (I) in crystalline form as claimed in any one of claims 1 to 6, microcrystalline cellulose and mannitol, wherein the relative weights of microcrystalline cellulose and mannitol within the tablet are in a ratio of from 3:1 to 1:1 and wherein the amount of Compound (I) in crystalline form within the tablet is from 50 to 500 mg.