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DESCRIPTION

Technical Field

[0001] The present invention relates to an antitumor agent containing a combination of an azabicyclo compound or a salt thereof and other antitumor agent(s), and an antitumor effect potentiator for other antitumor agent(s).

Background Art

[0002] A group of proteins called molecular chaperons is a multifunctional protein, which promotes the formation of the functional structures of other proteins or maintains these structures, promotes correct association, inhibits unnecessary aggregation, protects other proteins from degradation, and promotes secretion (Non Patent Literature 1). HSP90 is a molecular chaperon as abundant as approximately 1 to 2% of all intracellular soluble proteins and is however unnecessary for the biosynthesis of the majority of polypeptides, unlike other chaperon proteins (Non Patent Literature 1). Signaling-related factors (for example, ERBB1/EGFR, ERBB2/HER2, MET, IGF1R, KDR/VEGFR, FLT3, ZAP70, KIT, CHUK/IKK, BRAF, RAF1, SRC, and AKT), cell cycle regulators (for example, CDK4, CDK6, Cyclin D, PLK1, and BIRC5), and transcriptional regulators (for example, HIF-1 α , p53, androgen receptor, estrogen receptor, and progesterone receptor) are known as the main client proteins whose structure formation or stability is regulated by HSP90 through the interaction therebetween (Non Patent Literatures 2 and 3). HSP90 is deeply involved in cell proliferation or survival by maintaining the normal functions of these proteins. Furthermore, HSP90 is required for the normal functions of mutated or chimeric factors (for example, BCR-ABL and NPM-ALK) which cause carcinogenesis or exacerbation of cancer. This indicates the importance of HSP90 particularly for processes such as carcinogenesis, cancer survival, growth, exacerbation, and metastasis (Non Patent Literature 2).

[0003] The inhibition of the chaperon functions of HSP90 by specific inhibitors such as geldanamycin causes the inactivation, destabilization, and degradation of the client proteins, resulting in induction of a halt in cell proliferation or apoptosis (Non Patent Literature 4). In terms of the physiological functions of HSP90, HSP90 inhibitors are characterized in that they can simultaneously inhibit multiple signaling pathways involved in cancer survival and growth. Thus, the HSP90 inhibitors can serve as pharmaceuticals having extensive and effective anticancer activity. Moreover, from the findings that cancer cell-derived HSP90 has higher activity and higher affinity for ATP or inhibitors than those of normal cell-derived HSP90, it has been expected that the HSP90 inhibitors would serve as pharmaceuticals having high cancer selectivity (Non Patent Literature 5). Currently, the clinical development of multiple HSP90 inhibitors as anticancer agents is ongoing. The most advancing Ganetespib is under development as single agents as well as under test on the combined use with other antitumor

agents such as docetaxel (Non Patent Literature 6).

[0004] Further, a new type of HSP90 inhibitor has been reported (Patent Literature 1), and HSP90 inhibitors with higher antitumor effects and reduced side effects have been desired. Patent Literature 2 describes an azabicyclo compound and its cell proliferation inhibitory effect.

Citation List

Patent Literature

[0005]

Patent Literature 1: WO 2011/004610 A

Patent Literature 2: WO 2006/066937 A2

Non Patent Literatures

[0006]

Non Patent Literature 1: Nature Reviews Cancer 5, 761-772(2005)

Non Patent Literature 2: TRENDS in Molecular Medicine 10(6), 283-290 (2004)

Non Patent Literature 3: Clin Can Res 15, 9-14(2009)

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Non Patent Literature 5: Drug Resistance Updates 12, 17-27(2009)

Non Patent Literature 6: Invest New Drugs. 30(6):2201-9 (2012)

Summary of Invention

Technical Problem

[0007] An object of the invention is to provide a method of using HSP90 inhibitors with high

antitumor effects and reduced side effects.

Solution to Problem

[0008] The present inventors examined combined use of various HSP90 inhibitors and other antitumor agent(s) under the circumstances. As a result, they found that an azabicyclo compound having an HSP90 inhibitory agent remarkably potentiates the antitumor effect of extremely various ranges of antitumor agents having different action mechanisms.

[0009] That is, the invention provides the following [1] to [12].

[0010]

1. [1] The compound 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamide or a salt thereof for use in treating a tumor by administration in combination with another antitumor agent.
2. [2] The compound according to item [1] or a salt thereof for use according to item [1], wherein the other antitumor agent is one or more of kind selected from the group consisting of an antitumor antibiotic substance, a platinum preparation, a pyrimidine-based antimetabolite agent, a purine-based antimetabolite agent, a folic acid antimetabolite agent, a plant alkaloid-based antitumor agent, an immunomodulating drug, and a low molecular weight molecular targeted drug.
3. [3] The compound according to item [1] or [2] or a salt thereof for use according to item [1] or [2], wherein the other antitumor agent is one or more of kind selected from the group consisting of amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabine, cytarabine, pemetrexed, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, etoposide, lenalidomide, imatinib, gefitinib, dasatinib, erlotinib, lapatinib, and crizotinib.
4. [4] The compound according to any one of items [1] to [3] or a salt thereof for use according to any one of items [1] to [3], wherein said compound or a salt thereof and the other antitumor agent are administered to a cancer patient simultaneously or separately at an interval.
5. [5] 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamide or a salt thereof for use in potentiating the antitumor effect of another antitumor agent.
6. [6] The compound according to item [5] or a salt thereof for use according to item [5], wherein the other antitumor agent is one or more of kind selected from the group consisting of an antitumor antibiotic substance, a platinum preparation, a pyrimidine-based antimetabolite agent, a purine-based antimetabolite agent, a folic acid antimetabolite agent, a plant alkaloid-based antitumor agent, an immunomodulating drug, and a low molecular weight molecular targeted drug.
7. [7] The compound according to item [5] or [6] or a salt thereof for use according to item [5] or [6], wherein the other antitumor agent is one or more of kind selected from the group consisting of amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabine, cytarabine,

pemetrexed, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, etoposide, lenalidomide, imatinib, gefitinib, dasatinib, erlotinib, lapatinib, and crizotinib.

8. [8] The compound according to any one of items [5] to [7] or a salt thereof for use according to any one of items [5] to [7], wherein said compound or a salt thereof and the other antitumor agent are administered to a cancer patient simultaneously or separately at an interval.
9. [9] A combination of 3-ethyl-4-(3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl)benzamide or a salt thereof and another antitumor agent for use in treating a tumor.
10. [10] The combination according to item [9] or a salt thereof for use according to item [9], wherein the other antitumor agent is one or more of kind selected from the group consisting of an antitumor antibiotic substance, a platinum preparation, a pyrimidine-based antimetabolite agent, a purine-based antimetabolite agent, a folic acid antimetabolite agent, a plant alkaloid-based antitumor agent, an immunomodulating drug, and a low molecular weight molecular targeted drug.
11. [11] The combination according to item [9] or [10] or a salt thereof for use according to item [9] or [10], wherein the other antitumor agent is one or more of kind selected from the group consisting of amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabine, cytarabine, pemetrexed, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, etoposide, lenalidomide, imatinib, gefitinib, dasatinib, erlotinib, lapatinib, and crizotinib.
12. [12] The combination according to any one of items [9] to [11] or a salt thereof for use according to any one of items [9] to [11], wherein said compound or a salt thereof and the other antitumor agent are administered to a cancer patient simultaneously or separately at an interval.

[0011] Also described is an antitumor agent, wherein the azabicyclo compound 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamide or a salt thereof is administered in combination with other antitumor agent(s).

Advantageous Effects of Invention

[0012] The antitumor agent of the invention serves to perform cancer therapy exhibiting high antitumor effect while suppressing side effects, and, thus, the survival of a cancer patient can be prolonged.

Brief Description of Drawings

[0013]

Fig. 1 shows an effect of the combination of Compound 1 and docetaxel with respect to the

rate of tumor growth of a human non-small cell lung cancer line NCI-H2170.

Fig. 2 shows an effect of the combination of Compound 1 and paclitaxel with respect to the rate of tumor growth of a human non-small cell lung cancer line NCI-H441.

Fig. 3 shows an effect of the combination of Compound 1 and cisplatin with respect to the rate of tumor growth of a human stomach cancer line NCI-N87.

Fig. 4 shows an effect of the combination of Compound 1 and amrubicin with respect to the rate of tumor growth of a human small cell lung cancer cell line SBC-1.

Description of Embodiments

[0014] The HSP90 inhibitor in the invention, which provides an excellent synergic effect with other antitumor agent(s), is an azabicyclo compound. The azabicyclo compound is 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamide (hereinafter, referred to as Compound 1).

[0015] The salt of the azabicyclo compound of the invention is not particularly limited as long as it is a pharmaceutically acceptable salt, and examples thereof include acid addition salts of inorganic acids (for example, hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, nitric acid, and phosphoric acid) and organic acids (for example, formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, succinic acid, fumaric acid, maleic acid, lactic acid, malic acid, citric acid, tartaric acid, carbonic acid, picric acid, methanesulfonic acid, p-toluenesulfonic acid, and glutamic acid); salts of inorganic bases (for example, sodium, potassium, magnesium, calcium, and aluminum), organic bases (for example, methylamine, ethylamine, meglumine, and ethanolamine), or a basic amino acids (for example, lysine, arginine, and ornithine); and ammonium salts.

[0016] Incidentally, the azabicyclo compound of the invention or a salt thereof can be synthesized according to the method described in WO 2011/004610 A, for example.

[0017] As described in Examples below, when the azabicyclo compound of the invention or a salt thereof is administered in combination with extremely various ranges of antitumor agents having different action mechanisms, the antitumor effect is synergistically potentiated. The "other antitumor agent" described in the invention is not particularly limited as long as it is a pharmaceutical having antitumor activity, and as the form thereof, any of a low molecular compound, an antibody, and a nucleic acid may be employed. Specific examples of antitumor agents exhibiting the synergic effect with the azabicyclo compound of the invention or a salt thereof include antitumor antibiotic substances such as amrubicin, doxorubicin, doxyl, epirubicin, and mitomycin C; alkylating agents such as cyclophosphamide and nimustine; platinum-based agents such as cisplatin, carboplatin, and oxaliplatin; pyrimidine-based

antimetabolite agents such as 5-fluorouracil (5-FU), tegafur/gimeracil/oteracil potassium (TS-1, generic name "tegafur/gimeracil/oteracil potassium compounding agent" (trade name: "TS-1")), tegafur/uracil (UFT, generic name "tegafur/uracil compounding agent" (trade name: "UFT")), capecitabine, doxifluridine, 5-fluoro-2'-deoxyuridine (FdUrd), gemcitabine, and cytarabine; purine-based antimetabolite agents such as fludarabine, bendamustine, cladribine, and nelarabine; folic acid antimetabolite agents such as pemetrexed and methotrexate; plant alkaloid-based antitumor agents such as paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, irinotecan, and etoposide; immunomodulating drugs such as lenalidomide; low molecular weight molecular targeted drugs such as imatinib, gefitinib, dasatinib, erlotinib, lapatinib, everolimus, temsirolimus, bortezomib, crizotinib, vemurafenib, and ZD6244; and antibody molecular targeted drugs such as bevacizumab, trastuzumab, cetuximab, and rituximab. Among these, in terms of the synergic effect of antitumor effect in the case of concurrently using the azabicyclo compound of the invention or a salt thereof, one or more of agent selected from the group consisting of antitumor antibiotic substances, platinum-based agents, pyrimidine-based antimetabolite agents, purine-based antimetabolite agents, folic acid antimetabolite agents, plant alkaloid-based antitumor agents, immunomodulating drugs, and low molecular weight molecular targeted drugs are preferable.

[0018] As a specific antitumor agent, in terms of the synergic effect of antitumor effect in the case of concurrently using the azabicyclo compound of the invention or a salt thereof, one or more of kind(s) selected from the group consisting of amrubicin, doxorubicin, mitomycin C, cisplatin, oxaliplatin, 5-FU, TS-1, UFT, gemcitabine, cytarabine, pemetrexed, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, irinotecan, etoposide, lenalidomide, imatinib, gefitinib, dasatinib, erlotinib, lapatinib, bortezomib, crizotinib, vemurafenib, and ZD6244 are preferable, one or more of kind(s) selected from the group consisting of amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabine, cytarabine, pemetrexed, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, etoposide, lenalidomide, imatinib, gefitinib, dasatinib, erlotinib, lapatinib, and crizotinib are more preferable, and one or more of kind(s) selected from the group consisting of amrubicin, cisplatin, etoposide, gemcitabine, paclitaxel (for example, TAXOL or ABRAXANE), docetaxel, and crizotinib are particularly preferable.

[0019] Specific examples of a cancer to be treated by the antitumor agent of the invention include head and neck cancer, digestive organ cancer (for example, esophageal cancer, stomach cancer, duodenal cancer, liver cancer, biliary tract cancer (for example, gallbladder/bile duct cancer), pancreatic cancer, small intestinal cancer, large intestine cancer (for example, colorectal cancer, colon cancer, or rectal cancer), lung cancer (for example, non-small cell lung cancer or small cell lung cancer), breast cancer, ovarian cancer, uterus cancer (for example, cervical cancer or uterine corpus cancer), kidney cancer, bladder cancer, prostate cancer, skin cancer (for example, malignant melanoma or epidermal cancer), and blood cancer (for example, multiple myeloma or acute myelocytic leukemia). Among these, in terms of the synergic effect of the antitumor effect when the azabicyclo compound of the invention or a salt thereof is used in combination with the antitumor agent, the cancer is preferably digestive organ cancer, lung cancer, breast cancer, skin cancer, or blood cancer, and more preferably colorectal cancer, lung cancer, breast cancer, gallbladder cancer,

pancreatic cancer, stomach cancer, skin cancer, or blood cancer. Incidentally, herein, the cancer includes not only primary tumor but also cancer metastasizing to other organ(s) (for example, liver). In addition, the antitumor agent of the invention may be used for postoperative adjuvant chemotherapy which is performed for prevention of recurrence after a tumor is removed surgically or may be used for preoperative adjuvant chemotherapy which is preliminarily performed for removing surgically a tumor.

[0020] There is no particular limitation on the administration form of the antitumor agent of the invention, and appropriate form can be selected in accordance with the therapeutic purpose. Specific examples of the form of the antitumor agent include oral agents (for examples, tablets, coated tablets, powder, granules, capsules, and liquid), injections, suppositories, cataplasms, and ointments.

[0021] The administration schedule of the antitumor agent of the invention is appropriately selected in the range in which each active ingredient exerts the antitumor effect, and each active ingredient is administered simultaneously or separately at an interval.

[0022] The antitumor agent of the invention may be prepared in such a manner that respective active ingredients are separated into multiple dosage forms or are collectively prepared in one dosage form, on the basis of the administration forms or the administration schedules of the active ingredients. In addition, each drug formulation may be produced and sold in one package suitable for use at each event of combined administration or may be produced and sold in separate packages.

[0023] The antitumor agent in the invention can be prepared through a generally known method by use of a pharmacologically acceptable carrier, according to the administration form. Such a carrier can be selected from the group consisting of a variety of carriers generally employed in pharmaceuticals, and examples thereof can include an excipient, a binder, a disintegrant, a lubricant, a diluent, a solubilizing agent, a suspending agent, a tonicity agent, a pH-adjusting agent, a buffer, a stabilizer, a coloring agent, a flavoring agent, and a deodorant.

[0024] The invention also relates to an antitumor effect potentiator containing the azabicyclo compound of the invention or a salt thereof, which is used for potentiating the antitumor effect of other antitumor agent(s) with respect to a cancer patient. The antitumor effect potentiator has the formulation aspect of the antitumor agent described above.

[0025] The present description also relates to an antitumor agent containing the azabicyclo compound of the invention or a salt thereof, which is used for treating a cancer patient who has been administered other antitumor agent(s). The antitumor agent has the above-described formulation aspect.

[0026] The present description also relates to a kit preparation including directions for use in which the azabicyclo compound of the invention or a salt thereof and administration of the azabicyclo compound of the invention or a salt thereof in combination with other antitumor

agent(s) to a cancer patient are described. Herein, the "directions for use" may be directions for use in which the above-described dosage is described. Although it does not matter whether the directions for use are legally bound or not, directions for use in which the above-described dosage is recommended are preferable. Specifically, a package insert or a pamphlet is exemplified. In addition, the kit preparation including directions for use may be a kit preparation in which directions for use are printed on or attached to the package of the kit preparation or a kit preparation in which directions for use are enclosed in the package of the kit preparation together with an antitumor agent.

Examples

[0027] Hereinafter, the invention will be described in more detail by means of Examples and Reference Examples.

[0028] Example 1: A combination of Compound 1 and docetaxel exhibits the potentiated antitumor activity with respect to a human tumor cell of nude mouse xenograft model having a non-small cell lung cancer (NSCLC) cell NCI-H2170.

[0029] NCI-H2170 (ATCC #CRL-5928) as a human non-small cell lung cancer (NSCLC) cell line was obtained from American Type Culture Collection (ATCC; Manassas, Virginia, USA). In RPMI-1640 (containing 4.5 g/L of glucose, 10 mM of HEPES, and 1 mM of sodium pyruvate) (Wako Pure Chemical Industries, Ltd.) medium supplemented with 10% fetal bovine serum (FBS), cell lines were cultured. The cells cryopreserved in liquid nitrogen were quickly thawed at 37°C, transferred to a tissue culture flask containing a medium, and allowed to grow in a 5% CO₂ incubator at 37°C. The NCI-H2170 cell lines were sub-cultured once to twice a week at a dilution ratio of 1 : 5 to 1 : 10. The cells which were grown to 80 to 90% confluence in a 75 cm² flask were washed with 10 mL of phosphate buffer saline (PBS), and 10 mL of 0.25% trypsin-EDTA was then added thereto, followed by incubation until the cells were separated from the surface of the flask. 10 mL of medium for inactivation of trypsin was added thereto, a cell suspension was collected, and after centrifugal separation, the cell pellet was re-suspended in 10 mL of growth medium and then seeded in a 175 cm² flask into which 30 mL of medium was put, followed by incubation in a 5% CO₂ incubator at 37°C. When the flask reached 80 to 90% confluence, the above-described sub-culturing was repeated until cells sufficient for transplantation into mice were obtained.

[0030] Five-week old BALB/cAJcl-nu/nu mice (nude mice) were obtained from CREA Japan, Inc. Animals were housed in 5 or 6 microisolator cages under a 12-hour light/12-hour dark cycle, acclimated at least one week before use, and appropriately fed with a usual feed. The animals at the time of transplantation were six- to eight-week old. In order to transplant NCI-H2170 cells to the nude mice, cells were collected as described above, washed in PBS, and re-suspended at a concentration of 5×10^7 cell/mL in 50% PBS and 50% Matrigel basement

membrane matrix (#356237; BD Biosciences; Bedford, Massachusetts, USA). A 5×10^6 cell/0.1 mL of cell suspension was transplanted subcutaneously to the right dorsal region of the nude mouse by using a 1 mL tuberculin syringe and a 25 G needle.

[0031] Subsequently, the cell was allowed to grow for one to two weeks after transplantation until the tumor volume (TV) reached 100 to 300 mm³. Digimatic Caliper was used in measurement of the tumor diameter, the major axis and the minor axis of the tumor were measured, and TV was calculated by the following formula.

$TV \text{ (mm}^3\text{)} = (\text{Major axis} \times \text{minor axis}^2)/2$, units of the major axis and the minor axis: mm

[0032] Animals having an extremely small or large tumor volume were excluded and the remaining animals were allocated to each group by a stratified randomization allocation method using TV as an index. A relative tumor volume (RTV) was calculated from TV as an endpoint. In addition, a treatment/control (T/C) value (%) that is an RTV ratio of the RTV of the pharmaceutical administered group to the RTV of the control group on the end date of test period was calculated as an evaluation index. RTV and T/C (%) were calculated by the following formula.

$RTV = TV_n / TV_1$

$T/C \text{ (\%)} = (\text{Average RTV of each pharmaceutical administered group on test end date}) / (\text{Average RTV of control group on test end date}) \times 100$

[0033] TV₁ represents the tumor volume of Day 1.

[0034] In order to administer Compound 1, a 0.5% hypromellose aqueous solution was prepared. Hypromellose was weighed in a beaker, and distilled water (Otsuka Pharmaceutical Factory, Inc.) was added thereto in an amount of about 80% of the preparation amount. The mixture was completely dissolved by being stirred overnight using a stirrer at a low-temperature chamber, the resultant solution was transferred to a measuring cylinder, and distilled water was added to dilute to the preparation amount. Compound 1 was weighed in a necessary amount, pulverized with an agate mortar, suspended with a 0.5% hypromellose aqueous solution to have a predetermined concentration, and then subjected to ultrasonic treatment, thereby obtaining a homogeneous suspension. 10 mL of this suspension was administered orally per kg of body weight. Incidentally, the suspension was stored in a refrigerator at times other than administration. This suspension is refrigerator-stable for two weeks.

[0035] As docetaxel, 80 mg of Taxotere (registered trademark) for intravenous drip infusion (Sanofi-Aventis SA) was dissolved according to the package insert, and was diluted with physiological saline so as to have a predetermined concentration immediately before administration. On the first day of administration, 5 mL of the prepared docetaxel was

administered intravenously per kg of body weight.

[0036] Both the treatment with 5 mg/kg of body weight of Compound 1 and the treatment with 15 mg/kg of body weight of docetaxel slightly inhibited the growth of NCI-H2170 tumor in the nude mice, and the T/C values were 72 and 86, respectively. In contrast, the concomitant treatment with a combination of 5 mg/kg of body weight of Compound 1 and 15 mg/kg of body weight of docetaxel further inhibited the growth of NCI-H2170 tumor in the nude mice, and the T/C value was 43. Further, the treatment with one of 10 mg/kg of body weight of Compound 1 and 30 mg/kg of body weight of docetaxel or a combination thereof were carried out. However, the effect observed in each concomitant treatment group was potentiated more significantly ($P < 0.05$; Intersection-Union test) compared to the effect observed in the group treated only with any one of pharmaceuticals. The results thereof are shown in Fig. 1. Regarding the average body weight change on the end date of test period, all of the groups treated with the combination of Compound 1 and docetaxel were accompanied with no toxicity with respect to the control group.

[0037] Example 2: A combination of Compound 1 and paclitaxel exhibits the potentiated antitumor activity with respect to a human tumor cell of nude mouse xenograft model having NSCLC NCI-H441 cells.

[0038] An NCI-H441 cell (ATCC #HTB-174) was transplanted to nude mice according to the description in Example 1. The cell was allowed to grow for one week after transplantation until the large majority of tumors reached 100 to 300 mm³.

[0039] In order to administer paclitaxel, paclitaxel was weighed in a necessary amount, ethanol (NACALAI TESQUE, INC.) was added thereto in an amount of 10% of the preparation amount, and the mixture was dissolved by ultrasonic treatment. Next, Cremophor EL (NACALAI TESQUE, INC.) was added thereto in the same amount as ethanol, and the mixture was dissolved by ultrasonic treatment. Physiological saline was added and mixed in an amount of 10% of the preparation amount immediately before administration, thereby preparing a paclitaxel administration solution. On the first day of administration, 5 mL of the paclitaxel administration solution was administered intravenously per kg of body weight.

[0040] As shown in Fig. 2, both the treatment with 5 mg/kg of body weight of Compound 1 and the treatment with 30 mg/kg of body weight of paclitaxel inhibited the growth of NCI-H441 tumor in the nude mice, and the T/C values were 63 and 41, respectively. In contrast, in the case of concomitant treatment with these pharmaceuticals, the inhibitory effect of NCI-H441 tumor growth in the nude mice was potentiated and the T/C value was 28. Further, the treatment with 15 mg/kg of body weight or 60 mg/kg of body weight of docetaxel alone and the concomitant treatment with docetaxel and 5 mg/kg of body weight of Compound 1 were carried out. However, the effect observed in each concomitant treatment group was potentiated more significantly ($P < 0.05$; Intersection-Union test) compared to the effect observed in the group treated only with any one of pharmaceuticals. Regarding the toxicity using the body weight change on the end date of test period as an index, the group treated with the combination of

Compound 1 and paclitaxel was well in the tolerable range with respect to the control group. In addition, the same result was also obtained in the stomach cancer line.

[0041] Example 3: A combination of Compound 1 and cisplatin exhibits the potentiated antitumor activity with respect to a human tumor cell of nude mouse xenograft model having a stomach cancer line NCI-N87 cell.

[0042] An NCI-N87 cell (ATCC #CRL-5822) was transplanted to nude mice according to the description in Example 1. The cell was allowed to grow for one week after transplantation until the large majority of tumors reached 100 to 300 mm³.

[0043] As cisplatin, 25 mg of Briplatin for injection (Bristol-Myers Squibb Company) was dissolved according to the package insert, and was diluted with physiological saline so as to have a predetermined concentration immediately before administration. On the first day of administration, 14.0 mL of the prepared cisplatin was administered intravenously per kg of body weight.

[0044] As shown in Fig. 3, both the treatment with 5 mg/kg of body weight of Compound 1 and the treatment with 7 mg/kg of body weight of cisplatin inhibited the growth of NCI-N87 tumor in the nude mice, and the T/C values were 51 and 65, respectively. In contrast, in the case of concomitant treatment with these pharmaceuticals, the inhibitory effect of NCI-N87 tumor growth in the nude mice was potentiated and the T/C value was 26. Further, the treatment with 10 mg/kg of body weight or 20 mg/kg of body weight of Compound 1 alone and the concomitant treatment with Compound 1 and 7 mg/kg of body weight of cisplatin were carried out. However, the effect observed in each concomitant treatment group was potentiated more significantly ($P < 0.05$; Intersection-Union test) compared to the effect observed in the group treated only with any one of pharmaceuticals. Regarding the toxicity using the body weight change on the end date of test period as an index, the group treated with the combination of Compound 1 and cisplatin was well in the tolerable range with respect to the control group.

[0045] Example 4: A combination of Compound 1 and amrubicin exhibits the potentiated antitumor activity with respect to a human tumor cell of nude mouse xenograft model having a small cell lung cancer line SBC-1 cell.

[0046] An SBC-1 cell (purchased from Health Science Research Resources Bank, #JCRB0816) was transplanted to nude mice according to the description in Example 1. The cell was allowed to grow for one week after transplantation until the large majority of tumors reached 100 to 300 mm³.

[0047] As amrubicin, 20 mg of Calsed (registered trademark) for injection (Sumitomo Dainippon Pharma Co., Ltd.) was dissolved according to the package insert, and was diluted with physiological saline so as to have a predetermined concentration immediately before administration. On the first day of administration, 10.0 mL of the prepared amrubicin was administered intravenously per kg of body weight.

[0048] On Days 1, 3, 5, 8, 10, and 12, 10.0 mL of Compound 1 was administered orally per kg of body weight.

[0049] As shown in Fig. 4, both the treatment with 28 mg/kg of body weight of Compound 1 and the treatment with 12.5 mg/kg of body weight of amrubicin inhibited the growth of SBC-1 tumor in the nude mice, and the T/C values were 49 and 40, respectively. In contrast, in the case of concomitant treatment with these pharmaceuticals, the inhibitory effect of SBC-1 tumor growth in the nude mice was potentiated and the T/C value was 17. Further, the treatment with 20 mg/kg of body weight of Compound 1 alone and the concomitant treatment with Compound 1 and 12.5 mg/kg of body weight of amrubicin were carried out. However, the effect observed in each concomitant treatment group was potentiated more significantly ($P < 0.05$; Intersection-Union test) compared to the effect observed in the group treated only with any one of pharmaceuticals. Regarding the toxicity using the body weight change on the end date of test period as an index, the group treated with the combination of Compound 1 and amrubicin was well in the tolerable range with respect to the control group.

Example 5: In vitro combination analysis of Compound 1 and crizotinib

A. Material and Method

[0050] In RPMI-1640 medium (Wako Pure Chemical Industries, Ltd.) supplemented with 10% fetal bovine serum (Thermo Scientific), human stomach cancer cell lines NUGC-4 and MKN45 (Health Science Research Resources Bank, HSRRB) were allowed to grow. All of the cells were maintained at 37°C and 5% CO₂, and sub-cultured once to twice a week at a dilution ratio of 1 : 5 to 1 : 20.

Cell Survival Assay

[0051] The cell survival rate was measured using CellTiter-Glo. The cells were recovered by a general method, suspended in RPMI-1640 medium supplemented with 10% fetal bovine serum and antibiotic substances (penicillin and streptomycin), and seeded in a 384-well plate. The number of cells to be seeded was set to 500 cell/20 µL per well. The seeded cells were incubated at 37°C and 5% CO₂ for 24 hours, and then 5 µL of a medium supplemented with crizotinib and Compound 1 or Vehicle (DMSO) was added thereto. As crizotinib, nine of a 3-fold dilution series from 10 µM and zero concentration (DMSO) were used, and as Compound 1, seven of 3-fold dilution series from 10 µM and zero concentration (DMSO) were used. All of 80 combinations thereof were examined. With respect to each combination, four wells were allocated. Further, incubation was carried out at 37°C and 5% CO₂ for 72 hours. 25 µL of

CellTiter-Glo liquid per well was added, and then incubation was carried out at room temperature for 10 minutes. The chemiluminescence was measured by Envision as a plate reader. The average value of four wells of each combination was calculated from the obtained data, and the standardized cell survival rate was calculated with respect to the control added with a medium supplemented with Vehicle. The Fa (Fraction of Affect) was calculated by subtracting the cell survival rate from 1.

[0052] The half inhibition concentration (IC₅₀) of each pharmaceutical was determined using median effect analysis software CalcuSyn 2.0 (CalcuSyn, Inc.). Subsequently, the combination index (CI) of each combination concentration of pharmaceuticals was determined. The CI values of more than 1, equal to 1, and less than 1 each indicate antagonistic effect, additive effect, and synergic effect (Table 1) (Pharmacol Rev. 2006; 58(3):621-81, BMC Complement Altern Med. 2013; 13:212., Anticancer Res. 2005; 25(3B) : 1909-17).

[Table 1]

(Explanation of combination index value)	
Range of CI (upper limit)	Explanation
0.1	Extremely strong synergic effect
0.3	Strong synergic effect
0.7	Synergic effect
0.85	Moderate synergic effect
0.9	Slight synergic effect
1	Substantially additive
1.2	Modest antagonistic effect
1.45	Moderate antagonistic effect
3.3	Antagonistic effect
10	Strong antagonistic effect
>10	Extremely strong antagonistic effect

[0053] Further, the Fa value close to 1 is considered to be a concentration range in which the effect of one of pharmaceuticals is too strong, and the Fa value close to 0 is considered to be a concentration range in which the effect of one of pharmaceuticals is too weak. These values are not suitable for discussion on the synergistic effect. Therefore, the concentration combination of both the pharmaceuticals satisfying $0.2 \leq Fa \leq 0.8$ was extracted from the Fa values calculated by 80 concentration combinations of Compound 1 and crizotinib in total in the NUGC-4 cell, and the extracted concentration combination is used in the linear curve fitting by CalcuSyn to obtain a CI.

B. Result

[0054] The obtained CI and the concentrations of both pharmaceuticals applied therewith were examined and the concentration range of each of both the pharmaceuticals in which the CI became moderate degree or higher synergic effect (less than 0.85) was found (Table 2).

[Table 2]

Compound 1 (nM)	Crizotinib (nM)	Fa	CI		Concomitant ratio
13.7174	4.57247	0.34	0.291	Strong synergic effect	1 : 0.33333
13.7174	13.7174	0.527	0.139	Strong synergic effect	1 : 1
13.7174	41.1523	0.53	0.383	Synergic effect	1 : 3.00001
41.1523	4.57247	0.339	0.368	Synergic effect	1 : 0.11111
41.1523	13.7174	0.579	0.103	Strong synergic effect	1 : 0.33333
41.1523	41.1523	0.534	0.391	Synergic effect	1 : 1
41.1523	123.457	0.567	0.816	Moderate synergic effect	1 : 3
123.457	1.52416	0.289	0.622	Synergic effect	1 : 0.01235
123.457	4.57247	0.417	0.319	Synergic effect	1 : 0.03704
123.457	13.7174	0.581	0.149	Strong synergic effect	1 : 0.11111
123.457	41.1523	0.582	0.3	Synergic effect	1 : 0.33333
370.37	1.52416	0.645	0.148	Strong synergic effect	1 : 0.00412
370.37	4.57247	0.624	0.182	Strong synergic effect	1 : 0.01235
370.37	13.7174	0.631	0.205	Strong synergic effect	1 : 0.03704
370.37	41.1523	0.64	0.278	Strong synergic effect	1 : 0.11111
370.37	123.457	0.66	0.446	Synergic effect	1 : 0.33333
1111.11	1.52416	0.75	0.201	Strong synergic effect	1 : 0.00137
1111.11	4.57247	0.74	0.221	Strong synergic effect	1 : 0.00412
1111.11	13.7174	0.742	0.227	Strong synergic effect	1 : 0.01235
1111.11	41.1523	0.739	0.262	Strong synergic effect	1 : 0.03704
1111.11	123.457	0.722	0.41	Synergic effect	1 : 0.11111

Compound 1 (nM)	Crizotinib (nM)	Fa	CI		Concomitant ratio
3333.33	41.1523	0.717	0.833	Moderate synergic effect	1 : 0.01235

[0055] In the concentration range from 13.7174 nM to 1111.11 nM of Compound 1 and the concentration range from 4.57247 nM to 41.1523 nM of crizotinib in the NUGC-4 cell, a large number of combinations exhibiting a strong synergic effect were found. This means that in these concentration ranges, cell death was induced greater than expected from the additive effect of dosage of respective pharmaceuticals.

[0056] In addition, similarly to the result of the NUGC-4 cell described above, regarding the MKN45 cell, when the incubation was carried out for 72 hours while Compound 1 and crizotinib were simultaneously added, a large number of combinations exhibiting a strong synergic effect also were found in the concentration range from 13.7174 nM to 370.37 nM of Compound 1 and the concentration range from 13.7174 nM to 123.457 nM of crizotinib (Table 3).

[Table 3]

Compound 1 (nM)	Crizotinib (nM)	Fa	CI		Concomitant ratio
13.7174	41.1523	0.796	0.048	Extremely strong synergic effect	1 : 3.00001
13.7174	370.37	0.795	0.418	Synergic effect	1 : 27.00001
41.1523	13.7174	0.346	0.772	Moderate synergic effect	1 : 0.33333
41.1523	41.1523	0.792	0.056	Extremely strong synergic effect	1 : 1
123.457	13.7174	0.395	0.709	Moderate synergic effect	1 : 0.11111
370.37	1.52416	0.562	0.34	Synergic effect	1 : 0.00412
370.37	4.57247	0.562	0.367	Synergic effect	1 : 0.01235
370.37	13.7174	0.727	0.153	Strong synergic effect	1 : 0.03704
370.37	41.1523	0.792	0.123	Strong synergic effect	1 : 0.11111
370.37	123.457	0.796	0.21	Strong synergic effect	1 : 0.33333
370.37	370.37	0.78	0.575	Synergic effect	1 : 1
1111.11	1.52416	0.733	0.354	Synergic effect	1 : 0.00137
1111.11	4.57247	0.74	0.344	Synergic effect	1 : 0.00412
1111.11	13.7174	0.723	0.408	Synergic effect	1 : 0.01235
1111.11	41.1523	0.769	0.332	Synergic effect	1 : 0.03704
1111.11	123.457	0.753	0.523	Synergic effect	1 : 0.11111

Compound 1 (nM)	Crizotinib (nM)	Fa	CI		Concomitant ratio
3333.33	1.52416	0.764	0.846	Moderate synergic effect	1 : 0.00046
3333.33	4.57247	0.77	0.813	Moderate synergic effect	1 : 0.00137
3333.33	41.1523	0.776	0.827	Moderate synergic effect	1 : 0.01235

Example 6: In vitro combination analysis of Compound 1 and other antitumor agents

[0057] The same in vitro combination analysis was also performed on a combination of Compound 1 and other antitumor agents with respect to a cell line other than those described above. All of the antitumor agents presented in Table 4 to Table 39 exhibited moderate degree or higher synergic effect with Compound 1 ($CI < 0.85$). Particularly, in imatinib, etoposide, erlotinib, oxaliplatin, gefitinib, gemcitabine, cisplatin, cytarabine, dasatinib, doxorubicin, docetaxel, paclitaxel, pemetrexed, lapatinib, and lenalidomide, a strong synergic effect ($CI < 0.30$) exhibited in one or more of concentration combination.

[Table 4]

<Antitumor agent:5-FU,Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	5555.56	0.563	0.475	Synergic effect	1 : 405.00095
41.1523	5555.56	0.537	0.613	Synergic effect	1 : 134.99999
3333.33	7.62079	0.676	0.804	Moderate synergic effect	1 : 0.00229
3333.33	22.8624	0.678	0.797	Moderate synergic effect	1 : 0.00686
3333.33	68.5871	0.698	0.726	Moderate synergic effect	1 : 0.02058
3333.33	205.761	0.688	0.765	Moderate synergic effect	1 : 0.06173
3333.33	617.284	0.701	0.728	Moderate synergic effect	1 : 0.18519
3333.33	1851.85	0.716	0.698	Synergic effect	1 : 0.55556
3333.33	5555.56	0.712	0.786	Moderate synergic effect	1 : 1.66667

[Table 5]

<Antitumor agent:ZD6244, Cancer type: malignant melanoma, Cell line:A375>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	41.1523	0.252	0.55	Synergic effect	1 : 3.00001
13.7174	123.457	0.479	0.314	Synergic effect	1 : 9.00003
13.7174	370.37	0.59	0.442	Synergic effect	1 : 27.00001
13.7174	1111.11	0.68	0.692	Synergic effect	1 : 81.00004
41.1523	123.457	0.447	0.415	Synergic effect	1 : 3
41.1523	370.37	0.577	0.501	Synergic effect	1 : 8.99998
41.1523	1111.11	0.685	0.679	Synergic effect	1 : 26.99995
123.457	123.457	0.432	0.546	Synergic effect	1 : 1
123.457	370.37	0.572	0.576	Synergic effect	1 : 2.99999
123.457	1111.11	0.681	0.741	Moderate synergic effect	1 : 8.99998
370.37	41.1523	0.352	0.758	Moderate synergic effect	1 : 0.11111
370.37	123.457	0.443	0.772	Moderate synergic effect	1 : 0.33333
370.37	370.37	0.606	0.624	Synergic effect	1 : 1
370.37	1111.11	0.69	0.813	Moderate synergic effect	1 : 3
1111.11	41.1523	0.61	0.741	Moderate synergic effect	1 : 0.03704
1111.11	123.457	0.68	0.63	Synergic effect	1 : 0.11111
1111.11	370.37	0.753	0.547	Synergic effect	1 : 0.33333
1111.11	1111.11	0.769	0.722	Moderate synergic effect	1 : 1

[Table 6]

<Antitumor agent:SN-38, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	13.7174	0.584	0.437	Synergic effect	1 : 1
41.1523	13.7174	0.514	0.652	Synergic effect	1 : 0.33333
41.1523	41.1523	0.773	0.384	Synergic effect	1 : 1
123.457	41.1523	0.719	0.578	Synergic effect	1 : 0.33333
3333.33	1.52416	0.557	0.584	Synergic effect	1 : 0.00046
3333.33	4.57247	0.558	0.694	Synergic effect	1 : 0.00137

<Antitumor agent:SN-38, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
3333.33	13.7174	0.611	0.807	Moderate synergic effect	1 : 0.00412
3333.33	41.1523	0.785	0.556	Synergic effect	1 : 0.01235

[Table 7]

<Antitumor agent: imatinib, Cancer type: acute myelocytic leukemia, Cell line: Kasumi-1>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	370.37	0.327	0.566	Synergic effect	1 : 27.00001
13.7174	1111.11	0.488	0.272	Strong synergic effect	1 : 81.00004
13.7174	3333.33	0.51	0.595	Synergic effect	1 : 243.00013
41.1523	370.37	0.369	0.446	Synergic effect	1 : 8.99998
41.1523	1111.11	0.52	0.256	Strong synergic effect	1 : 26.99995
41.1523	3333.33	0.55	0.443	Synergic effect	1 : 80.99985
123.457	370.37	0.409	0.561	Synergic effect	1 : 2.99999
123.457	1111.11	0.564	0.335	Synergic effect	1 : 8.99998
123.457	3333.33	0.556	0.576	Synergic effect	1 : 26.99993
370.37	4.57247	0.616	0.592	Synergic effect	1 : 0.01235
370.37	13.7174	0.621	0.583	Synergic effect	1 : 0.03704
370.37	41.1523	0.576	0.673	Synergic effect	1 : 0.11111
370.37	123.457	0.599	0.631	Synergic effect	1 : 0.33333
370.37	370.37	0.632	0.578	Synergic effect	1 : 1
370.37	1111.11	0.694	0.48	Synergic effect	1 : 3
370.37	3333.33	0.697	0.516	Synergic effect	1 : 9
370.37	30000	0.737	0.722	Moderate synergic effect	1 : 81.00008

[Table 8]

<Antitumor agent: etoposide, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	1111.11	0.421	0.772	Moderate synergic effect	1 : 81.00004

<Antitumor agent: etoposide, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
41.1523	3333.33	0.638	0.741	Moderate synergic effect	1 : 80.99985
123.457	1111.11	0.495	0.638	Synergic effect	1 : 8.99998
123.457	3333.33	0.655	0.723	Moderate synergic effect	1 : 26.99993
370.37	3333.33	0.71	0.648	Synergic effect	1 : 9
1111.11	1.52416	0.637	0.716	Moderate synergic effect	1 : 0.00137
1111.11	4.57247	0.64	0.709	Moderate synergic effect	1 : 0.00412
1111.11	13.7174	0.623	0.754	Moderate synergic effect	1 : 0.01235
1111.11	41.1523	0.638	0.722	Moderate synergic effect	1 : 0.03704
1111.11	123.457	0.671	0.655	Synergic effect	1 : 0.11111
1111.11	370.37	0.685	0.662	Synergic effect	1 : 0.33333
1111.11	1111.11	0.714	0.689	Synergic effect	1 : 1
1111.11	3333.33	0.753	0.808	Moderate synergic effect	1 : 3

[Table 9]

<Antitumor agent: etoposide, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	3333.33	0.279	0.29	Strong synergic effect	1 : 243.00013
13.7174	10000	0.624	0.037	Extremely strong synergic effect	1 : 729.00112
41.1523	10000	0.672	0.028	Extremely strong synergic effect	1 : 242.99978
123.457	3333.33	0.23	0.649	Synergic effect	1 : 26.99993
123.457	10000	0.616	0.061	Extremely strong synergic effect	1 : 80.99986
370.37	10000	0.587	0.132	Strong synergic effect	1 : 27.00003
1111.11	10000	0.472	0.555	Synergic effect	1 : 9.00001
10000	3333.33	0.782	0.79	Moderate synergic effect	1 : 0.33333

<Antitumor agent: etoposide, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
10000	10000	0.783	0.789	Moderate synergic effect	1 : 1

[Table 10]

<Antitumor agent: erlotinib, Cancer type: non-small cell lung cancer, Cell line: A549>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	370.37	0.275	0.572	Synergic effect	1 : 27
13.7174	1111.11	0.377	0.715	Moderate synergic effect	1 : 81
13.7174	10000	0.695	0.611	Synergic effect	1 : 729
41.1523	123.457	0.275	0.36	Synergic effect	1 : 3
41.1523	370.37	0.345	0.426	Synergic effect	1 : 9
41.1523	1111.11	0.386	0.749	Moderate synergic effect	1 : 27
41.1523	3333.33	0.516	0.825	Moderate synergic effect	1 : 81
41.1523	10000	0.729	0.479	Synergic effect	1 : 243
123.457	123.457	0.281	0.716	Moderate synergic effect	1 : 1
123.457	370.37	0.359	0.653	Synergic effect	1 : 3
123.457	10000	0.72	0.58	Synergic effect	1 : 81
370.37	1.52416	0.558	0.554	Synergic effect	1 : 0
370.37	4.57247	0.581	0.508	Synergic effect	1 : 0.01
370.37	13.7174	0.549	0.576	Synergic effect	1 : 0.04
370.37	41.1523	0.579	0.517	Synergic effect	1 : 0.11
370.37	123.457	0.683	0.344	Synergic effect	1 : 0.33
370.37	370.37	0.718	0.305	Synergic effect	1 : 1
370.37	1111.11	0.772	0.253	Strong synergic effect	1 : 3
1111.11	13.7174	0.795	0.58	Synergic effect	1 : 0.01

[Table 11]

<Antitumor agent: oxaliplatin, Cancer type: large intestine cancer, Cell line :HCT-116>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	3333.33	0.489	0.562	Synergic effect	1 : 243.00013
13.7174	10000	0.784	0.104	Strong synergic effect	1 : 729.00112
41.1523	3333.33	0.488	0.604	Synergic effect	1 : 80.99985
41.1523	10000	0.777	0.129	Strong synergic effect	1 : 242.99978
123.457	3333.33	0.498	0.666	Synergic effect	1 : 26.99993
123.457	10000	0.79	0.158	Strong synergic effect	1 : 80.99986
370.37	1.52416	0.461	0.539	Synergic effect	1 : 0.00412
370.37	4.57247	0.484	0.508	Synergic effect	1 : 0.01235
370.37	13.7174	0.463	0.538	Synergic effect	1 : 0.03704
370.37	41.1523	0.483	0.516	Synergic effect	1 : 0.11111
370.37	123.457	0.509	0.493	Synergic effect	1 : 0.33333
370.37	370.37	0.558	0.453	Synergic effect	1 : 1
370.37	1111.11	0.604	0.439	Synergic effect	1 : 3
370.37	3333.33	0.771	0.261	Strong synergic effect	1 : 9

[Table 12]

<Antitumor agent: gefitinib, Cancer type: non-small cell lung cancer, Cell line:HCC827>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	2.28624	0.215	0.66	Synergic effect	1 : 0.16667
13.7174	6.85871	0.477	0.262	Strong synergic effect	1 : 0.5
13.7174	20.5761	0.739	0.125	Strong synergic effect	1 : 1.5
41.1523	6.85871	0.442	0.382	Synergic effect	1 : 0.16667
41.1523	20.5761	0.721	0.154	Strong synergic effect	1 : 0.5
41.1523	61.7284	0.798	0.223	Strong synergic effect	1 : 1.5
123.457	20.5761	0.68	0.248	Strong synergic effect	1 : 0.16667
123.457	61.7284	0.751	0.364	Synergic effect	1 : 0.5

<Antitumor agent: gefitinib, Cancer type: non-small cell lung cancer, Cell line:HCC827>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
123.457	185.185	0.797	0.676	Synergic effect	1 : 1.5
370.37	20.5761	0.598	0.63	Synergic effect	1 : 0.05556
370.37	61.7284	0.662	0.846	Moderate synergic effect	1 : 0.16667
1111.11	0.762079	0.75	0.321	Synergic effect	1 : 0.00069
1111.11	2.28624	0.749	0.332	Synergic effect	1 : 0.00206
1111.11	6.85871	0.753	0.346	Synergic effect	1 : 0.00617
1111.11	20.5761	0.776	0.345	Synergic effect	1 : 0.01852
1111.11	61.7284	0.774	0.53	Synergic effect	1 : 0.05556
3333.33	2.28624	0.796	0.654	Synergic effect	1 : 0.00069
3333.33	6.85871	0.787	0.726	Moderate synergic effect	1 : 0.00206
3333.33	20.5761	0.799	0.699	Synergic effect	1 : 0.00617

[Table 13]

<Antitumor agent: gemcitabine, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	13.7174	0.767	0.038	Extremely strong synergic effect	1 : 1
41.1523	13.7174	0.697	0.111	Strong synergic effect	1 : 0.33333
123.457	13.7174	0.736	0.103	Strong synergic effect	1 : 0.11111
370.37	13.7174	0.598	0.585	Synergic effect	1 : 0.03704
370.37	123.457	0.75	0.528	Synergic effect	1 : 0.33333
1111.11	4.57247	0.675	0.641	Synergic effect	1 : 0.00412
1111.11	13.7174	0.783	0.369	Synergic effect	1 : 0.01235
1111.11	41.1523	0.773	0.455	Synergic effect	1 : 0.03704
1111.11	123.457	0.762	0.713	Moderate synergic effect	1 : 0.11111

[Table 14]

<Antitumor agent: gemcitabine, Cancer type: gallbladder cancer, Cell line:TGBC2TKB>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
500.0	62.5	0.660	0.371	Synergic effect	1 : 0.13
500.0	125.0	0.708	0.365	Synergic effect	1 : 0.25
500.0	250.0	0.721	0.450	Synergic effect	1 : 0.5
500.0	500.0	0.715	0.682	Synergic effect	1 : 1
1000.0	62.5	0.652	0.686	Synergic effect	1 : 0.06
1000.0	125.0	0.695	0.649	Synergic effect	1 : 0.13
1000.0	250.0	0.700	0.756	Moderate synergic effect	1 : 0.25

[Table 15]

<Antitumor agent: gemcitabine, Cancer type: non-small cell lung cancer, Cell line:NCI-H522>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
1250	200	0.391	0.723	Moderate synergic effect	1 : 0.16
1250	400	0.470	0.595	Synergic effect	1 : 0.32
1250	800	0.567	0.490	Synergic effect	1 : 0.64
2500	200	0.527	0.435	Synergic effect	1 : 0.08
2500	400	0.602	0.329	Synergic effect	1 : 0.16
2500	800	0.594	0.502	Synergic effect	1 : 0.32
5000	200	0.578	0.522	Synergic effect	1 : 0.04
5000	400	0.644	0.376	Synergic effect	1 : 0.08
5000	800	0.627	0.548	Synergic effect	1 : 0.16
10000	200	0.618	0.718	Moderate synergic effect	1 : 0.02
10000	400	0.638	0.675	Synergic effect	1 : 0.04

[Table 16]

<Antitumor agent: gemcitabine, Cancer type: epidermal cancer, Cell line:A431>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	123.457	0.649	0.429	Synergic effect	1 : 9.00003
41.1523	370.37	0.8	0.469	Synergic effect	1 : 8.99998
123.457	123.457	0.59	0.62	Synergic effect	1 : 1

<Antitumor agent: gemcitabine, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
370.37	123.457	0.566	0.767	Moderate synergic effect	1 : 0.33333
1111.11	123.457	0.606	0.781	Moderate synergic effect	1 : 0.11111

[Table 17]

<Antitumor agent: cisplatin, Cancer type: small cell lung cancer, Cell line:NCI-H69>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
1000	625	0.515	0.787	Moderate synergic effect	1 : 0.63
1000	5000	0.798	0.791	Moderate synergic effect	1 : 5
1000	10000	0.868	0.847	Moderate synergic effect	1 : 10
2000	625	0.717	0.582	Synergic effect	1 : 0.31
2000	1250	0.694	0.783	Moderate synergic effect	1 : 0.63
2000	2500	0.785	0.679	Synergic effect	1 : 1.25
2000	5000	0.847	0.681	Synergic effect	1 : 2.5
2000	10000	0.891	0.763	Moderate synergic effect	1 : 5

[Table 18]

<Antitumor agent: cisplatin, Cancer type: non-small cell lung cancer, Cell line:A549>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	22.8624	0.337	0.149	Strong synergic effect	1 : 1.66667
13.7174	68.5871	0.321	0.193	Strong synergic effect	1 : 5.00001
13.7174	50000	0.752	0.465	Synergic effect	1 : 3645.00561
41.1523	22.8624	0.302	0.519	Synergic effect	1 : 0.55556
41.1523	68.5871	0.324	0.484	Synergic effect	1 : 1.66667
41.1523	50000	0.705	0.837	Moderate synergic effect	1 : 1214.99892
123.457	22.8624	0.507	0.516	Synergic effect	1 : 0.18519

<Antitumor agent: cisplatin, Cancer type: non-small cell lung cancer, Cell line:A549>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
123.457	68.5871	0.5	0.54	Synergic effect	1 : 0.55555
123.457	50000	0.763	0.517	Synergic effect	1 : 404.99931
370.37	22.8624	0.797	0.293	Strong synergic effect	1 : 0.06173
370.37	68.5871	0.763	0.375	Synergic effect	1 : 0.18519
370.37	1851.85	0.666	0.722	Moderate synergic effect	1 : 5
370.37	16666.7	0.751	0.558	Synergic effect	1 : 45.00014

[Table 19]

<Antitumor agent: cisplatin, Cancer type: epidermal cancer, Cell line:KB>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
250	313	0.743	0.816	Moderate synergic effect	1 : 1.25
250	625	0.871	0.758	Moderate synergic effect	1 : 2.5
500	156	0.887	0.608	Synergic effect	1 : 0.31
500	313	0.932	0.546	Synergic effect	1 : 0.63
500	625	0.953	0.568	Synergic effect	1 : 1.25

[Table 20]

<Antitumor agent: cytarabine, Cancer type: large intestine cancer, Cell line:HCT-116>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	41.1523	0.357	0.412	Synergic effect	1 : 3.00001
13.7174	123.457	0.574	0.346	Synergic effect	1 : 9.00003
13.7174	370.37	0.722	0.408	Synergic effect	1 : 27.00001
41.1523	41.1523	0.369	0.447	Synergic effect	1 : 1
41.1523	123.457	0.565	0.393	Synergic effect	1 : 3
41.1523	370.37	0.695	0.506	Synergic effect	1 : 8.99998
123.457	41.1523	0.409	0.518	Synergic effect	1 : 0.33333
123.457	123.457	0.533	0.566	Synergic effect	1 : 1
123.457	370.37	0.755	0.373	Synergic effect	1 : 2.99999
370.37	1.52416	0.393	0.774	Moderate synergic effect	1 : 0.00412

<Antitumor agent: cytarabine, Cancer type: large intestine cancer, Cell line:HCT-116>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
370.37	13.7174	0.418	0.778	Moderate synergic effect	1 : 0.03704
370.37	41.1523	0.65	0.347	Synergic effect	1 : 0.11111
370.37	123.457	0.78	0.234	Strong synergic effect	1 : 0.33333
1111.11	1.52416	0.725	0.592	Synergic effect	1 : 0.00137
1111.11	4.57247	0.739	0.555	Synergic effect	1 : 0.00412
1111.11	13.7174	0.698	0.687	Synergic effect	1 : 0.01235
1111.11	41.1523	0.77	0.5	Synergic effect	1 : 0.03704

[Table 21]

<Antitumor agent: cytarabine, Cancer type: epidermal cancer, Cell line:A431>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	370.37	0.557	0.831	Moderate synergic effect	1 : 27.00001
13.7174	1111.11	0.769	0.55	Synergic effect	1 : 81.00004
41.1523	370.37	0.56	0.82	Moderate synergic effect	1 : 8.99998
41.1523	1111.11	0.769	0.552	Synergic effect	1 : 26.99995
123.457	1111.11	0.74	0.712	Moderate synergic effect	1 : 8.99998
370.37	370.37	0.594	0.706	Moderate synergic effect	1 : 1
1111.11	41.1523	0.325	0.696	Synergic effect	1 : 0.03704
3333.33	1.52416	0.552	0.512	Synergic effect	1 : 0.00046
3333.33	4.57247	0.527	0.55	Synergic effect	1 : 0.00137
3333.33	13.7174	0.596	0.483	Synergic effect	1 : 0.00412
3333.33	41.1523	0.672	0.425	Synergic effect	1 : 0.01235
3333.33	123.457	0.688	0.481	Synergic effect	1 : 0.03704
3333.33	370.37	0.711	0.636	Synergic effect	1 : 0.11111

[Table 22]

<Antitumor agent: dasatinib, Cancer type: acute myelocytic leukemia, Cell line:Kasumi-1>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	13.7174	0.527	0.265	Strong synergic effect	1 : 1
13.7174	41.1523	0.628	0.195	Strong synergic effect	1 : 3.00001
13.7174	123.457	0.666	0.298	Strong synergic effect	1 : 9.00003
13.7174	370.37	0.668	0.793	Moderate synergic effect	1 : 27.00001
41.1523	13.7174	0.534	0.385	Synergic effect	1 : 0.33333
41.1523	41.1523	0.602	0.368	Synergic effect	1 : 1
41.1523	123.457	0.648	0.462	Synergic effect	1 : 3
123.457	13.7174	0.556	0.701	Moderate synergic effect	1 : 0.11111
123.457	41.1523	0.683	0.383	Synergic effect	1 : 0.33333
123.457	123.457	0.65	0.699	Synergic effect	1 : 1
123.457	370.37	0.711	0.665	Synergic effect	1 : 2.99999
370.37	41.1523	0.779	0.57	Synergic effect	1 : 0.11111
370.37	123.457	0.794	0.536	Synergic effect	1 : 0.33333
370.37	370.37	0.8	0.566	Synergic effect	1 : 1

[Table 23]

<Antitumor agent: doxorubicin, Cancer type: small cell lung cancer, Cell line:SBC-1>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
500.0	62.5	0.789	0.542	Synergic effect	1 : 0.13
500.0	125.0	0.886	0.509	Synergic effect	1 : 0.25
1000.0	15.6	0.647	0.584	Synergic effect	1 : 0.02
1000.0	31.3	0.826	0.300	Synergic effect	1 : 0.03
1000.0	62.5	0.906	0.244	Strong synergic effect	1 : 0.06
1000.0	125.0	0.927	0.339	Synergic effect	1 : 0.13
2000.0	7.8	0.710	0.592	Synergic effect	1 : 0.0039
2000.0	15.6	0.770	0.474	Synergic effect	1 : 0.01
2000.0	31.3	0.826	0.412	Synergic effect	1 : 0.02
2000.0	62.5	0.866	0.437	Synergic effect	1 : 0.03

<Antitumor agent: doxorubicin, Cancer type: small cell lung cancer, Cell line:SBC-1>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
2000.0	125.0	0.913	0.449	Synergic effect	1 : 0.06

[Table 24]

<Antitumor agent: doxorubicin, Cancer type: large intestine cancer, Cell line:HCT-116>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	41.1523	0.389	0.356	Synergic effect	1 : 3.00001
13.7174	123.457	0.581	0.502	Synergic effect	1 : 9.00003
13.7174	370.37	0.763	0.69	Synergic effect	1 : 27.00001
41.1523	41.1523	0.433	0.348	Synergic effect	1 : 1
41.1523	123.457	0.525	0.65	Synergic effect	1 : 3
123.457	41.1523	0.408	0.534	Synergic effect	1 : 0.33333
123.457	123.457	0.537	0.719	Moderate synergic effect	1 : 1
370.37	13.7174	0.451	0.674	Synergic effect	1 : 0.03704
370.37	41.1523	0.605	0.493	Synergic effect	1 : 0.11111
370.37	123.457	0.618	0.753	Moderate synergic effect	1 : 0.33333
1111.11	1.52416	0.737	0.618	Synergic effect	1 : 0.00137
1111.11	4.57247	0.706	0.713	Moderate synergic effect	1 : 0.00412
1111.11	13.7174	0.712	0.717	Moderate synergic effect	1 : 0.01235
1111.11	41.1523	0.705	0.804	Moderate synergic effect	1 : 0.03704

[Table 25]

<Antitumor agent: docetaxel, Cancer type: stomach cancer, Cell line:NCI-N87>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
100	4	0.568	0.807	Moderate synergic effect	1 : 0.04
100	8	0.707	0.716	Moderate synergic effect	1 : 0.08
200	4	0.731	0.697	Synergic effect	1 : 0.02
200	8	0.792	0.683	Synergic effect	1 : 0.04

<Antitumor agent: docetaxel, Cancer type: stomach cancer, Cell line:NCI-N87>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
200	16	0.848	0.709	Moderate synergic effect	1 : 0.08
400	4	0.812	0.845	Moderate synergic effect	1 : 0.01
400	8	0.85	0.792	Moderate synergic effect	1 : 0.02
400	16	0.879	0.816	Moderate synergic effect	1 : 0.04

[Table 26]

<Antitumor agent: docetaxel, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	0.457247	0.345	0.761	Moderate synergic effect	1 : 0.03333
13.7174	1.37174	0.691	0.21	Strong synergic effect	1 : 0.1
13.7174	4.11523	0.791	0.257	Strong synergic effect	1 : 0.3
41.1523	1.37174	0.71	0.206	Strong synergic effect	1 : 0.03333
41.1523	4.11523	0.789	0.279	Strong synergic effect	1 : 0.1
123.457	1.37174	0.651	0.407	Synergic effect	1 : 0.01111
370.37	1.37174	0.758	0.374	Synergic effect	1 : 0.0037
1111.11	0.152416	0.787	0.679	Synergic effect	1 : 0.00014

[Table 27]

<Antitumor agent: docetaxel, Cancer type: non-small cell lung cancer, Cell line:NCI-H226>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
500	1.25	0.640	0.265	Strong synergic effect	1 : 0.0025
500	2.50	0.723	0.195	Strong synergic effect	1 : 0.01
500	5.00	0.803	0.135	Strong synergic effect	1 : 0.01

<Antitumor agent: docetaxel, Cancer type: non-small cell lung cancer, Cell line:NCI-H226>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
500	10.00	0.852	0.110	Strong synergic effect	1 : 0.02
500	20.00	0.855	0.157	Strong synergic effect	1 : 0.04
1000	1.25	0.538	0.771	Moderate synergic effect	1 : 0.0013
2000	1.25	0.673	0.710	Moderate synergic effect	1 : 0.00063

[Table 28]

<Antitumor agent: docetaxel, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	0.457247	0.281	0.681	Synergic effect	1 : 0.03333
13.7174	1.37174	0.618	0.129	Strong synergic effect	1 : 0.1
13.7174	4.11523	0.766	0.097	Extremely strong synergic effect	1 : 0.3
41.1523	0.457247	0.281	0.705	Moderate synergic effect	1 : 0.01111
41.1523	1.37174	0.645	0.108	Strong synergic effect	1 : 0.03333
41.1523	4.11523	0.751	0.116	Strong synergic effect	1 : 0.1
123.457	1.37174	0.604	0.167	Strong synergic effect	1 : 0.01111
123.457	4.11523	0.724	0.162	Strong synergic effect	1 : 0.03333
370.37	1.37174	0.435	0.693	Synergic effect	1 : 0.0037
370.37	4.11523	0.589	0.56	Synergic effect	1 : 0.01111
3333.33	0.152416	0.653	0.562	Synergic effect	1 : 0.00005
3333.33	0.457247	0.663	0.555	Synergic effect	1 : 0.00014
3333.33	1.37174	0.672	0.584	Synergic effect	1 : 0.00041
3333.33	4.11523	0.709	0.59	Synergic effect	1 : 0.00123

[Table 29]

<Antitumor agent: paclitaxel, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	4.11523	0.651	0.313	Synergic effect	1 : 0.3
41.1523	4.11523	0.625	0.38	Synergic effect	1 : 0.1

<Antitumor agent: paclitaxel, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
123.457	4.11523	0.615	0.454	Synergic effect	1 : 0.03333
370.37	4.11523	0.598	0.667	Synergic effect	1 : 0.01111
1111.11	0.152416	0.601	0.777	Moderate synergic effect	1 : 0.00014
1111.11	1.37174	0.637	0.778	Moderate synergic effect	1 : 0.00123
1111.11	4.11523	0.79	0.469	Synergic effect	1 : 0.0037

[Table 30]

<Antitumor agent: paclitaxel, Cancer type: non-small cell lung cancer, Cell line:NCI-H2170>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
100	2.00	0.574	0.669	Synergic effect	1 : 0.02
100	4.00	0.746	0.528	Synergic effect	1 : 0.04
100	8.00	0.821	0.579	Synergic effect	1 : 0.08
200	2.00	0.753	0.576	Synergic effect	1 : 0.01
200	4.00	0.832	0.507	Synergic effect	1 : 0.02
200	8.00	0.891	0.475	Synergic effect	1 : 0.04
400	2.00	0.871	0.590	Synergic effect	1 : 0.01
400	4.00	0.892	0.577	Synergic effect	1 : 0.01
400	8.00	0.912	0.593	Synergic effect	1 : 0.02

[Table 31]

<Antitumor agent: paclitaxel, Cancer type: epidermal cancer, Cell line:A431>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	4.11523	0.675	0.281	Strong synergic effect	1 : 0.3
41.1523	4.11523	0.643	0.357	Synergic effect	1 : 0.1
41.1523	12.3457	0.777	0.367	Synergic effect	1 : 0.3
123.457	4.11523	0.636	0.386	Synergic effect	1 : 0.03333
123.457	12.3457	0.8	0.3	Synergic effect	1 : 0.1
370.37	4.11523	0.553	0.714	Moderate synergic effect	1 : 0.01111
370.37	12.3457	0.766	0.432	Synergic effect	1 : 0.03333

[Table 32]

<Antitumor agent: vemurafenib, Cancer type: malignant melanoma, Cell line:A375>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	123.457	0.367	0.651	Synergic effect	1 : 9.00003
13.7174	370.37	0.494	0.768	Moderate synergic effect	1 : 27.00001
41.1523	123.457	0.343	0.84	Moderate synergic effect	1 : 3
41.1523	370.37	0.492	0.811	Moderate synergic effect	1 : 8.99998

[Table 33]

<Antitumor agent: pemetrexed, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	123.457	0.215	0.386	Synergic effect	1 : 9.00003
13.7174	370.37	0.436	0.238	Strong synergic effect	1 : 27.00001
13.7174	1111.11	0.534	0.379	Synergic effect	1 : 81.00004
13.7174	3333.33	0.623	0.643	Synergic effect	1 : 243.00013
41.1523	370.37	0.388	0.358	Synergic effect	1 : 8.99998
41.1523	1111.11	0.526	0.424	Synergic effect	1 : 26.99995
41.1523	3333.33	0.621	0.672	Synergic effect	1 : 80.99985
123.457	1111.11	0.484	0.633	Synergic effect	1 : 8.99998
370.37	1.52416	0.337	0.662	Synergic effect	1 : 0.00412
370.37	4.57247	0.387	0.565	Synergic effect	1 : 0.01235
370.37	13.7174	0.278	0.835	Moderate synergic effect	1 : 0.03704
370.37	41.1523	0.33	0.724	Moderate synergic effect	1 : 0.11111
370.37	123.457	0.324	0.84	Moderate synergic effect	1 : 0.33333
370.37	370.37	0.499	0.55	Synergic effect	1 : 1
1111.11	1.52416	0.726	0.578	Synergic effect	1 : 0.00137
1111.11	4.57247	0.714	0.605	Synergic effect	1 : 0.00412
1111.11	13.7174	0.729	0.573	Synergic effect	1 : 0.01235
1111.11	41.1523	0.745	0.541	Synergic effect	1 : 0.03704
1111.11	123.457	0.75	0.537	Synergic effect	1 : 0.11111

<Antitumor agent: pemetrexed, Cancer type: large intestine cancer, Cell line:HCT-116>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
1111.11	370.37	0.764	0.524	Synergic effect	1 : 0.33333
1111.11	1111.11	0.721	0.697	Synergic effect	1 : 1
1111.11	3333.33	0.749	0.79	Moderate synergic effect	1 : 3

[Table 34]

<Antitumor agent: pemetrexed, Cancer type: epidermal cancer, cell line:A431>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	123.457	0.554	0.273	Strong synergic effect	1 : 9.00003
13.7174	370.37	0.638	0.38	Synergic effect	1 : 27.00001
13.7174	1111.11	0.697	0.636	Synergic effect	1 : 81.00004
41.1523	123.457	0.509	0.409	Synergic effect	1 : 3
41.1523	370.37	0.617	0.466	Synergic effect	1 : 8.99998
3333.33	1.52416	0.536	0.551	Synergic effect	1 : 0.00046
3333.33	4.57247	0.554	0.537	Synergic effect	1 : 0.00137
3333.33	13.7174	0.54	0.576	Synergic effect	1 : 0.00412
3333.33	41.1523	0.534	0.657	Synergic effect	1 : 0.01235
3333.33	123.457	0.565	0.761	Moderate synergic effect	1 : 0.03704
3333.33	370.37	0.656	0.744	Moderate synergic effect	1 : 0.11111

[Table 35]

<Antitumor agent: bortezomib, Cancer type: multiple myeloma, Cell line:RPMI-8226>

Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
400	0.5	0.4	0.832	Moderate synergic effect	1 : 0.00125
800	0.5	0.75	0.662	Synergic effect	1 : 0.00063
800	0.8	0.75	0.721	Moderate synergic effect	1 : 0.001
800	1	0.77	0.719	Moderate synergic effect	1 : 0.00125

[Table 36]

<Antitumor agent: mitomycin C, Cancer type: large intestine cancer, Cell line:HCT-116>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
500.0	12.5	0.879	0.757	Moderate synergic effect	1 : 0.025

[Table 37]

<Antitumor agent: lapatinib, Cancer type: stomach cancer, Cell line:NCI-N87>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
250	3.90625	0.728	0.658	Synergic effect	1 : 0.016
250	7.8125	0.746	0.692	Synergic effect	1 : 0.031
250	15.625	0.784	0.707	Moderate synergic effect	1 : 0.063
250	31.25	0.810	0.818	Moderate synergic effect	1 : 0.125

[Table 38]

<Antitumor agent: lapatinib, Cancer type: breast cancer, Cell line:UACC-893>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
1000	62.5	0.590	0.511	Synergic effect	1 : 0.06
1000	125	0.643	0.362	Synergic effect	1 : 0.13
1000	250	0.629	0.613	Synergic effect	1 : 0.25
1000	500	0.715	0.494	Synergic effect	1 : 0.5
1000	1000	0.726	0.848	Moderate synergic effect	1 : 1
2000	62.5	0.665	0.344	Synergic effect	1 : 0.03
2000	125	0.710	0.246	Strong synergic effect	1 : 0.06
2000	250	0.737	0.269	Strong synergic effect	1 : 0.13
2000	500	0.749	0.403	Synergic effect	1 : 0.25
2000	1000	0.761	0.661	Synergic effect	1 : 0.5
4000	62.5	0.708	0.333	Synergic effect	1 : 0.02
4000	125	0.709	0.384	Synergic effect	1 : 0.03
4000	250	0.712	0.480	Synergic effect	1 : 0.06
4000	500	0.746	0.488	Synergic effect	1 : 0.13
4000	1000	0.783	0.576	Synergic effect	1 : 0.25
8000	62.5	0.702	0.666	Synergic effect	1 : 0.01

<Antitumor agent: lapatinib, Cancer type: breast cancer, Cell line:UACC-893>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
8000	125	0.722	0.542	Synergic effect	1 : 0.02
8000	250	0.726	0.611	Synergic effect	1 : 0.03
8000	500	0.754	0.575	Synergic effect	1 : 0.06
8000	1000	0.775	0.711	Moderate synergic effect	1 : 0.13

[Table 39]

<Antitumor agent: lenalidomide, Cancer type: multiple myeloma, Cell line:MM.1S>					
Compound 1 (nM)	Antitumor agent (nM)	Fa	CI	CI_COMMENT	Concomitant ratio
13.7174	15.2416	0.227	0.657	Synergic effect	1 : 1.11111
13.7174	45.7247	0.323	0.551	Synergic effect	1 : 3.33334
13.7174	137.174	0.493	0.284	Strong synergic effect	1 : 10
13.7174	411.523	0.618	0.225	Strong synergic effect	1 : 30.00007
13.7174	1234.57	0.735	0.168	Strong synergic effect	1 : 90.00029
13.7174	3703.7	0.765	0.293	Strong synergic effect	1 : 270.00015
13.7174	11111.1	0.776	0.693	Synergic effect	1 : 810.00044
41.1523	15.2416	0.244	0.806	Moderate synergic effect	1 : 0.37037
41.1523	45.7247	0.376	0.523	Synergic effect	1 : 1.11111
41.1523	137.174	0.536	0.329	Synergic effect	1 : 3.33333
41.1523	411.523	0.652	0.263	Strong synergic effect	1 : 10
41.1523	1234.57	0.756	0.206	Strong synergic effect	1 : 30.00002
41.1523	3703.7	0.777	0.317	Synergic effect	1 : 89.99983
123.457	137.174	0.552	0.67	Synergic effect	1 : 1.11111
123.457	411.523	0.693	0.463	Synergic effect	1 : 3.33333
123.457	1234.57	0.788	0.352	Synergic effect	1 : 10

[0058] Incidentally, gemcitabine exhibited the same synergic effect as in NCI-H522 even in other non-small cell lung cancers (NCI-H441, NCI-H520). Cisplatin exhibited the same synergic effect as in A549 even in other non-small cell lung cancers (NCI-H2170, NCI-H226, NCI-H441,

NCI-H520, NCI-H522, PC-14). Doxorubicin exhibited the same synergic effect as in SBC-1 even in other small cell lung cancers (NCI-N417). Docetaxel exhibited the same synergic effect as in NCI-H226 even in other non-small cell lung cancers (NCI-H2170). Paclitaxel exhibited the same synergic effect as in NCI-H2170 even in other non-small cell lung cancers (HCC827, NCI-H1975). Lapatinib exhibited the same synergic effect as in UACC-893 even in other breast cancers (BT-474, JIMT-1, MDA-MB-361, MDA-MB-453, SK-BR-3, UACC-812).

[0059] As described above, it was clearly found that the azabicyclo compound of the invention or a salt thereof exhibited a strong synergic effect with extremely various ranges of antitumor agents having different action mechanisms.

REFERENCES CITED IN THE DESCRIPTION

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KOMBINATIONSCANCERTERAPI VED HJÆLP AF AZABICYCLOFORBINDELSE

PATENTKRAV

1. Forbindelse 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamid eller et salt deraf til anvendelse i behandling af en tumor ved indgivelse i kombination med et andet antitumormiddel.
2. Forbindelse ifølge krav 1 eller et salt deraf til anvendelse ifølge krav 1, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af et antibiotisk antitumorstof, et platinpræparat, et pyrimidin-baseret antimetabolitmiddel, et purin-baseret antimetabolitmiddel, et folinsyreantimetabolitmiddel, der er udvalgt fra pemetrexed og methotrexat, et plantealkaloid-baseret antitumormiddel, et immunregulerende lægemiddel og et målrettet molekylært lægemiddel med lav molekylvægt.
3. Forbindelse ifølge krav 1 eller 2 eller et salt deraf til anvendelse ifølge krav 1 eller 2, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabin, cytarabin, pemetrexed, paclitaxel (for eksempel TAXOL eller ABRAXANE), docetaxel, etoposid, lenalidomid, imatinib, gefitinib, dasatinib, erlotinib, lapatinib og crizotinib.
4. Forbindelse ifølge et hvilket som helst af kravene 1 til 3 eller et salt deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 3, hvor forbindelsen eller et salt deraf og det andet antitumormiddel indgives til en cancerpatient samtidigt eller separat med et interval.
5. 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamid eller et salt deraf til anvendelse i potensering af et andet antitumormiddels antitumorstof.
6. Forbindelse ifølge krav 5 eller et salt deraf til anvendelse ifølge krav 5, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af et antibiotisk antitumorstof, et platinpræparat, et pyrimidin-baseret antimetabolitmiddel, et purin-baseret antimetabolitmiddel, et folinsyreantimetabolitmiddel, der er udvalgt fra pemetrexed og methotrexat, et plantealkaloid-baseret antitumormiddel, et immunregulerende lægemiddel og et målrettet molekylært lægemiddel med lav molekylvægt.
7. Forbindelse ifølge krav 5 eller 6 eller et salt deraf til anvendelse ifølge krav 5 eller 6, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabin, cytarabin, pemetrexed, paclitaxel (for eksempel TAXOL eller ABRAXANE), docetaxel, etoposid, lenalidomid, imatinib, gefitinib, dasatinib, erlotinib, lapatinib og crizotinib.
8. Forbindelse ifølge et hvilket som helst af kravene 5 til 7 eller et salt deraf til anvendelse ifølge et hvilket som helst af kravene 5 til 7, hvor forbindelsen eller et salt deraf og det andet antitumormiddel indgives til en cancerpatient samtidigt eller separat med et interval.

- 2 -

9. Kombination af 3-ethyl-4-{3-isopropyl-4-(4-(1-methyl-1H-pyrazol-4-yl)-1H-imidazol-1-yl)-1H-pyrazolo[3,4-b]pyridin-1-yl}benzamid eller et salt deraf og et andet antitumormiddel til anvendelse i behandling af en tumor.
10. Kombination ifølge krav 9 eller et salt deraf til anvendelse ifølge krav 9, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af et antibiotisk antitumorstof, et platinpræparat, et pyrimidin-baseret antimetabolitmiddel, et purin-baseret antimetabolitmiddel, et folinsyreantimetabolitmiddel, der er udvalgt fra pemetrexed og methotrexat, et plantealkaloid-baseret antitumormiddel, et immunregulerende lægemiddel og et målrettet molekylært lægemiddel med lav molekylvægt.
- 10 11. Kombination ifølge krav 9 eller 10 eller et salt deraf til anvendelse ifølge krav 9 eller 10, hvor det andet antitumormiddel er ét eller flere af en type, der er udvalgt fra gruppen bestående af amrubicin, doxorubicin, cisplatin, oxaliplatin, gemcitabin, cytarabin, pemetrexed, paclitaxel (for eksempel TAXOL eller ABRAXANE), docetaxel, etoposid, lenalidomid, imatinib, gefitinib, dasatinib, erlotinib, lapatinib og crizotinib.
- 15 12. Kombination ifølge et hvilket som helst af kravene 9 til 11 eller et salt deraf til anvendelse ifølge et hvilket som helst af kravene 9 til 11, hvor forbindelsen eller et salt deraf og det andet antitumormiddel indgives til en cancerpatient samtidigt eller separat med et interval.

DRAWINGS

FIG. 1

Group	Dose (mg/kg/day)	T/C (%)	BWC(%) Mean $\pm$ S.D.
Control	-	100	3.8 $\pm$ 2.7
Docetaxel	15.0	86	2.7 $\pm$ 2.0
Docetaxel	30.0	63	7.7 $\pm$ 3.0
Compound 1	5.0	72	5.0 $\pm$ 3.1
Compound 1	10.0	44	3.9 $\pm$ 2.1
Docetaxel/Compound 1	15.0 / 5.0	43	5.5 $\pm$ 2.3
Docetaxel/Compound 1	15.0 / 10.0	22	2.5 $\pm$ 9.1
Docetaxel/Compound 1	30.0 / 5.0	23	0.7 $\pm$ 5.3
Docetaxel/Compound 1	30.0 / 10.0	16	-9.6 $\pm$ 15.6

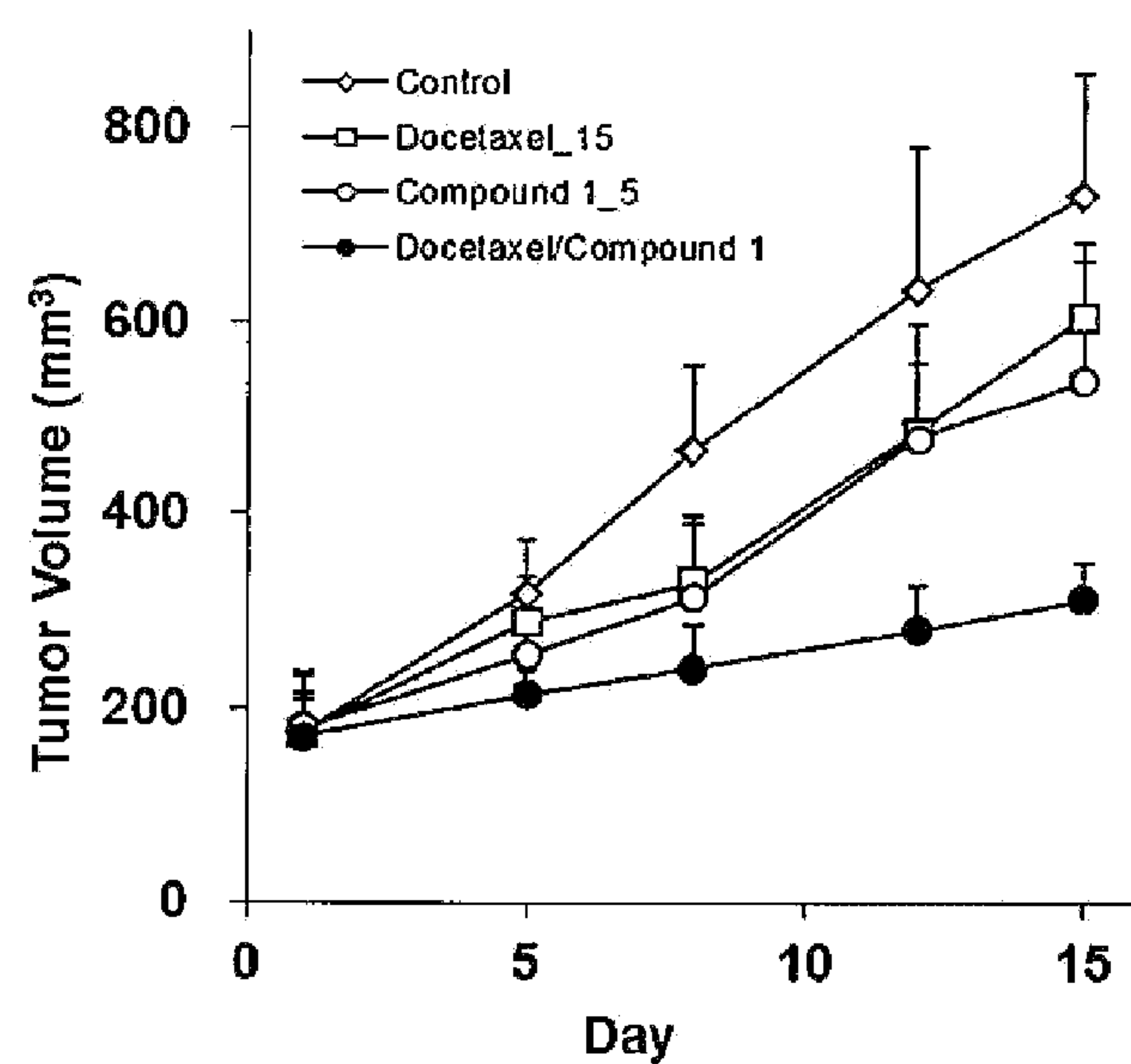


FIG. 2

Group	Dose (mg/kg/day)	T/C (%)	BWC(%) Mean $\pm$ S.D.
Control	-	100	5.8 $\pm$ 5.5
Paclitaxel	15.0	70	8.2 $\pm$ 2.8
Paclitaxel	30.0	41	8.5 $\pm$ 3.1
Paclitaxel	60.0	24	8.1 $\pm$ 3.1
Compound 1	5.0	63	4.6 $\pm$ 3.2
Paclitaxel/Compound 1	15.0/5.0	45	4.4 $\pm$ 4.6
Paclitaxel/Compound 1	30.0/5.0	28	7.0 $\pm$ 3.5
Paclitaxel/Compound 1	60.0/5.0	17	5.5 $\pm$ 3.3

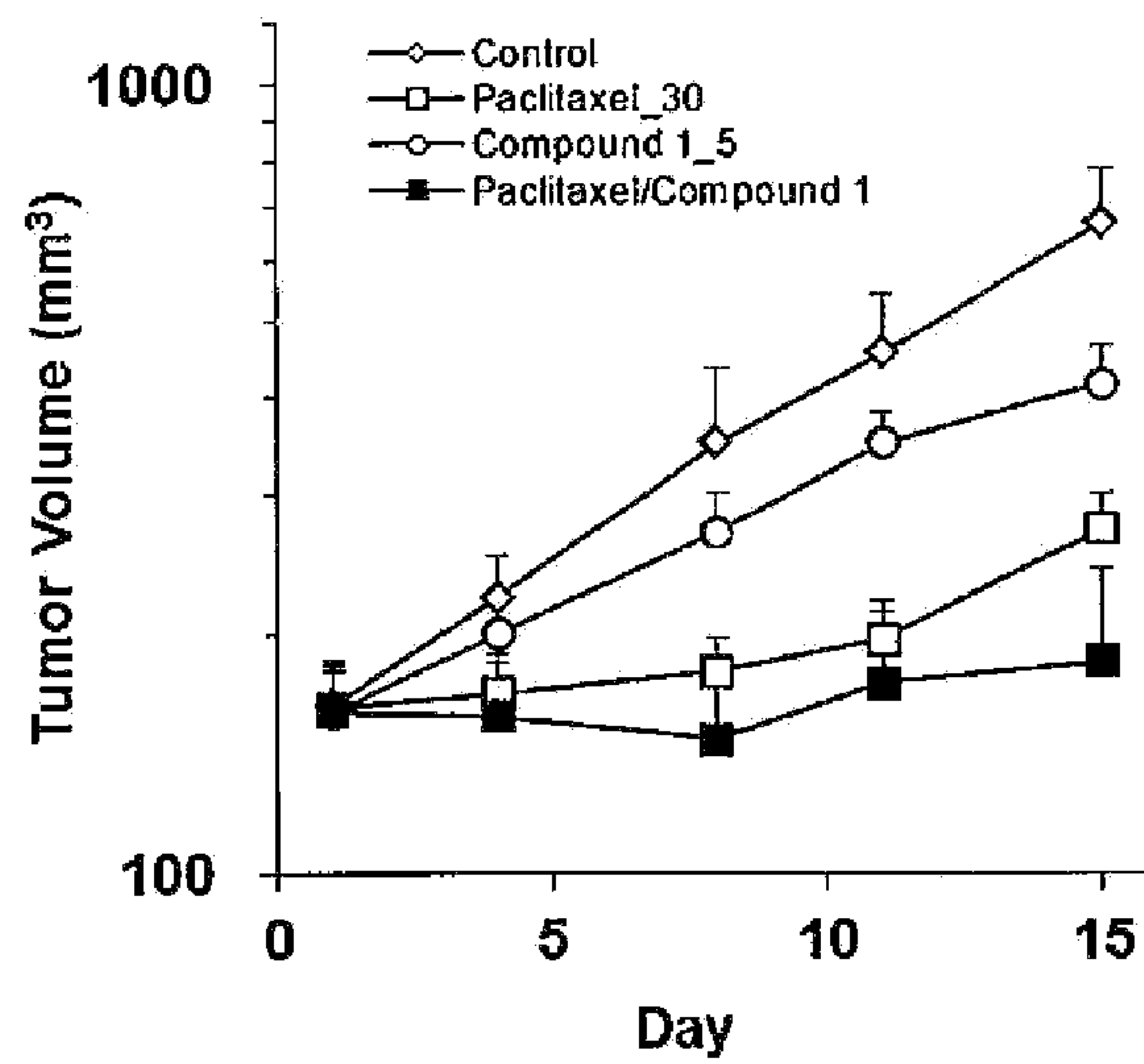


FIG. 3

Group	Dose (mg/kg/day)	Schedule (day)	T/C (%)	BWC(%) Mean $\pm$ S.D.
Control	-	-	100	6.5 $\pm$ 2.0
Cisplatin	7.0	1	65	5.6 $\pm$ 3.6
Compound 1	5.0	1-14	51	6.5 $\pm$ 6.9
Compound 1	10.0	1,3,5,8,10,12	54	3.1 $\pm$ 2.7
Compound 1	20.0	1,4,8,11	46	2.7 $\pm$ 3.2
Cisplatin/Compound 1	7.0/5.0	1/1-14	26	4.7 $\pm$ 7.5
Cisplatin/Compound 1	7.0/10.0	1/1,3,5,8,10,12	26	3.6 $\pm$ 3.0
Cisplatin/Compound 1	7.0/20.0	1/1,4,8,11	25	3.5 $\pm$ 5.3

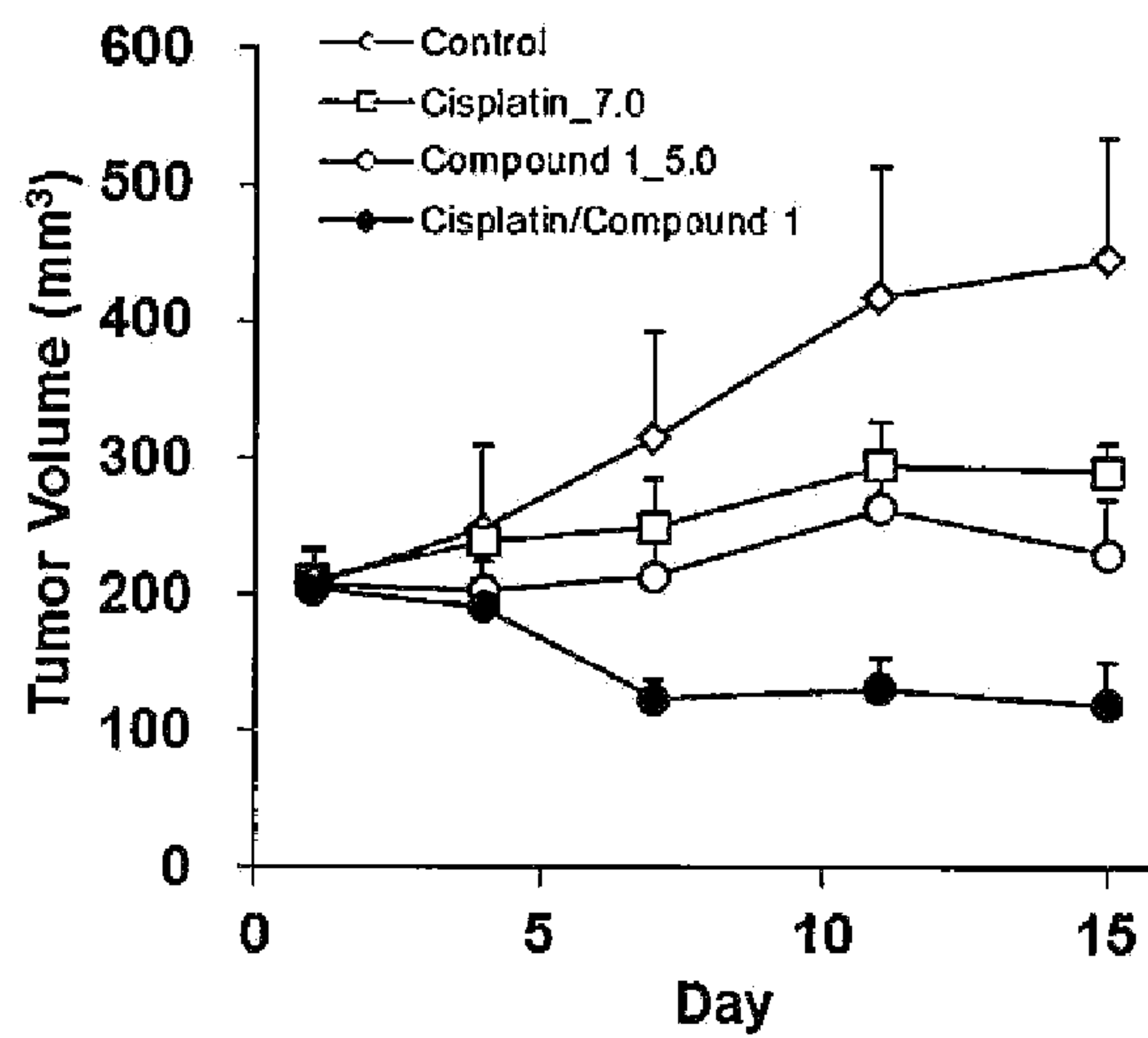


FIG. 4

Group	Dose (mg/kg/day)	T/C (%)	BWC(%) Mean $\pm$ S.D.
Control	-	100	7.8 $\pm$ 4.6
Amrubicin	12.5	40	8.0 $\pm$ 3.2
Compound 1	20.0	62	5.9 $\pm$ 3.8
Compound 1	28.0	49	5.4 $\pm$ 6.5
Amrubicin/Compound 1	12.5/20.0	21	5.3 $\pm$ 2.0
Amrubicin/Compound 1	12.5/28.0	17	-0.6 $\pm$ 15.6

