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(54) **ADMINISTRATION OF AN ANTITUMOR AGENT**

VERABREICHUNG EINES ANTITUMORMITTELS

ADMINISTRATION D'UN AGENT ANTI-TUMORAL

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US-A1- 2010 152 190

- **Hoffman -La Roche: "A First-In-Human Study of RO5503781 in Patients With Advanced Malignancies, Except Leukemia", INTERNET , April 2013 (2013-04), XP002696086, Retrieved from the Internet: URL:http://www.clinicaltrials.gov/ct2/show/NCT01462175?term=RO5503781&rank=2 [retrieved on 2013-04-24]**

EP 2 827 858 B1

Description

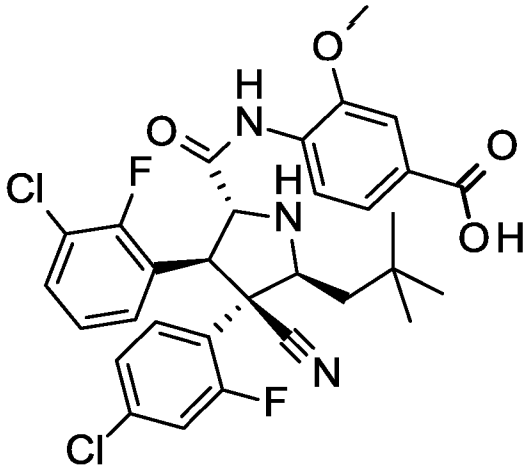
Field of the Invention

[0001] The present invention is related to improved dosage regimens of 4-{[(2R,3S,4R,5S)-4-(4-Chloro-2-fluoro-phenyl)-3-(3-chloro-2-fluoro-phenyl)-4-cyano-5-(2,2-dimethyl-propyl)-pyrrolidine-2-carbonyl]-amino}-3-methoxy-benzoic acid (referred to herein as Compound A) in the treatment of cancer. In particular, the invention relates to improved dosage regimens of Compound A that provide desirable antineoplastic effects with a tolerable level of toxicity. The regimens of the invention are characterized by administering less frequent doses comprising relatively high concentrations of Compound A. This protocol is expected to be safer and at least as effective as, possibly more effective than, administering more frequent doses at lower concentrations or larger doses at intermittent periods.

[0002] The present invention also relates to a pharmaceutical product comprising, as an active ingredient, the Compound A, characterized by administering said Compound A according to the above-mentioned improved protocol.

Background of the Invention

[0003] Compound A is an orally administered pyrrolidine that inhibits the binding of MDM2 to p53 and is thus useful in the treatment of cancer. It has the following chemical structure:



[0004] Compound A recently entered into phase I clinical trials for the treatment of solid tumors. See ClinicalTrials.gov, identifier NCT01462175. This compound is disclosed in US Pub 2010/0152190 A1. The Compound A, as well as a method for making it, is also disclosed in WO2011/098398.

[0005] Applicants have discovered that Compound A is especially effective, and best tolerated, in cancer therapy when administered in the specific doses and pursuant to the specific protocols herein described.

Summary of the Invention

[0006] The present invention relates to the compound A for treating a patient suffering with cancer, in particular colon, breast, prostate, lung or kidney cancer or osteosarcoma, wherein Compound A is administered to the patient in an amount of from 800 to 3000 mg/day, or from 1000 to 2500 mg/day, or from 1250 to 1800 mg/day, for an administration period of 5 days, on days 1-5, of a 28 day treatment cycle, followed by a rest period of 23 days.

[0007] The present invention also relates to a pharmaceutical product comprising, as an active ingredient, the Compound A for use in the treatment of cancer, characterized by administering said Compound A in the amounts and dosages indicated above.

Brief Description of the Drawings

[0008]

Figure 1 illustrates the antitumor activity, as demonstrated by the change in mean tumor volume over time, of Compound A monotherapy for a number of different dosing schedules, including a continuous 5 day dosing schedule. Figure 2 shows the increased lifespan of mice treated with Compound A for the different dosing schedules also

reflected in Figure 1.

Detailed Description of the Invention

[0009] "Tumor control" means that the perpendicular diameters of measurable lesions have not increased by 25% or more from the last measurement. *See, e.g.,* World Health Organization ("WHO") Handbook for Reporting Results of Cancer Treatment, Geneva (1979). The determination of tumor control or shrinkage (also referred to as "regression") is made by known methods. For example, by evaluation of patient symptoms, physical examination, X-ray, MRI or CAT scan or other commonly accepted evaluation modalities.

[0010] In one embodiment, the present invention relates to a pharmaceutical product comprising, as an active ingredient, the Compound A for use in the treatment of cancer, characterized by administering said Compound A in an amount of from 800 to 3000 mg/day, or from 1000 to 2500 mg/day, or from 1250 to 1800 mg/day, for an administration period of 5 days, on days 1-5, of a 28 day treatment cycle, followed by a rest period of 23 days. The course of a preferred cycle is 28 days. This treatment cycle is repeated for as long as the tumor remains under control and the regimen is clinically tolerated. The treatment cycle may, for example, be repeated up to 12 times.

[0011] In another embodiment, the present invention relates to a dosage regimen of treating a patient suffering with cancer, in particular colon, breast, prostate or kidney cancer as well as osteo or tissue sarcoma, comprising administering to the patient Compound A in an amount of from 800 to 3000 mg/day, or from 1000 to 2500 mg/day, or from 1250 to 1800 mg/day, for an administration period of 5 days, on days 1-5, of a 28 day treatment cycle, followed by a rest period of 23 days. The course of a preferred cycle is about 28 days. This treatment cycle is repeated for as long as the tumor remains under control and the regimen is clinically tolerated.

[0012] Dosages of Compound A can be applied either as a body surface area ("BSA") adapted dose (mg/m^2/day) or following flat dosing (mg/day). Compound A may be administered as a single dose daily or divided into multiple daily doses.

[0013] A patient's body measurement in square meters ("m^2") typically ranges from about 1.4 m^2 to about 2.2 m^2. Thus, the total amount of Compound A to be delivered in a treatment cycle (mg) using a BSA adapted dose would be calculated as follows:

[Dose intensity(mg/m^2/week)] x [BSA(m^2)] x [number of weeks in treatment cycle]

[0014] In an embodiment, the present product or method is characterized in that Compound A is administered daily for 5 days, on days 1-5 of a treatment cycle, followed by a rest period of 23 days ("5+/23-"). The 5+/23- treatment schedule is expected to be superior to interim schedules or to longer schedules as currently on-going Phase I studies indicate that in solid tumors, maximal apoptosis occurs only after about 48 hours of continuous exposure and longer schedules seem to present occurrence of delayed thrombocytopenia ("TCP"). Thus, a 3-5 daily treatment schedule is expected to provide the best benefit ratio taking into consideration efficacy and toxicity

[0015] In certain embodiments, the present product is characterized in that Compound A is administered daily, either once or twice (bid) daily, preferably once daily. The compound is administered to the patient in an oral unit dosage form, most preferably in tablet form.

[0016] Preferably, the 5 day treatment schedule is repeated every twenty-eight days, or as soon as permitted by recovery from toxicity, for so long as the tumor is under control or regressing and the patient tolerates the regimen. Preferably, these treatment cycles are repeated for a total of up to 12 cycles.

[0017] In an embodiment, the present product is characterized in that Compound A is administered daily in an amount from 800 to 3000 mg/day for 5 days on days 1-5 of a 28 day cycle.

[0018] In another embodiment, the present product is characterized in that Compound A is administered daily in an amount from 1000 to 2500 mg/day for 5 days on days 1-5 of a 28 day cycle.

[0019] In another embodiment, the present product is characterized in that Compound A is administered daily in an amount from 1250 to 1800 mg/day for 5 days on days 1-5 of a 28 day cycle.

[0020] In yet another embodiment there is provided a pharmaceutical product as defined above for the treatment of cancer, in particular solid tumors, more particularly colon, breast, prostate or kidney cancer as well as osteo or tissue sarcoma.

[0021] The present invention may be exemplified by controlled preclinical animal studies as shown in the Examples below, which illustrates the invention.

[0022] The superiority of the 5 day regimen of the present invention on solid tumors is demonstrated by the following experiments.

[0023] Abbreviations used herein are as follows:

x	times
po	orally

bid	twice daily
wk	week
qd	once daily
qdx5	once daily for five days
qweekly or 1 x/wk	once a week
BWL	body weight loss
SD	standard deviation

Toxicity

[0024] In the examples below, weight loss was graphically represented as percent change in mean group body weight, using the formula: $((W - W_0)/W_0) \times 100$, where 'W' represents mean body weight of the treated group at a particular day, and 'W_0' represents mean body weight of the same treated group at initiation of treatment. Maximum weight loss was also represented using the above formula, and indicated the maximum percent body weight loss that was observed at any time during the entire experiment for a particular group. Toxicity is defined as ≥20% of mice in a given group demonstrating ≥20% body weight loss and/or death.

Tumor Growth Inhibition (TGI) and Assessment of Survival/increase in Life Span (ILS)

[0025] Efficacy data was graphically represented as the mean tumor volume ± standard error of the mean (SEM). In addition, tumor volumes of treated groups were presented as percentages of tumor volumes of the control groups (%T/C), using the formula: $100 \times ((T - T_0)/(C - C_0))$, where T represented mean tumor volume of a treated group on a specific day during the experiment, T_0 represented mean tumor volume of the same treated group on the first day of treatment; C represented mean tumor volume of a control group on the specific day during the experiment, and C_0 represented mean tumor volume of the same treated group on the first day of treatment.

[0026] Tumor volume (in cubic millimeters) was calculated using the ellipsoid formula: $(D \times (d^2))/2$, where "D" represents the large diameter of the tumor and "d" represents the small diameter. In some cases, tumor regression and/or percent change in tumor volume was calculated using the formula: $((T- T_0)/ T_0) \times 100$, where 'T' represents mean tumor volume of the treated group at a particular day, and 'T_0' represents mean tumor volume of the same treated group at initiation of treatment.

[0027] Statistical analysis was determined by the rank sum test and One Way Anova and a post-hoc Bonferroni t-test (SigmaStat, version 2.0, Jandel Scientific, San Francisco, CA, USA). Differences between groups were considered to be significant when the probability value (p) was ≤0.05.

[0028] For survival assessment, the percent of increased life space (ILS) was calculated as: 100 x [(median survival day of treated group - median survival day of control group)/median survival day of control group]. Median survival was determined utilizing Kaplan Meier survival analysis. Survival in treated groups was statistically compared with the vehicle group and survival comparisons were done between groups using the log-rank test (Graph Pad Prism, La Jolla, CA, USA). Differences between groups were considered significant when the probability value (p) was ≤0.05.

Example 1

[0029] The antitumor activity of Compound A in the human osteosarcoma cancer xenograft model SJASA1 in immunocompromized mice using a variety of different schedules was assessed.

Test Compound A

[0030] Compound A was formulated as an amorphous solid dispersion micro-bulk precipitate (MBP) powder containing 30% drug substance and 70% HPMC-AS polymer was reconstituted immediately before administration as a suspension in Klucel/ Tween, and remaining suspension was discarded after dosing. All dose levels are reported as the actual dosage of Compound A rather than including drug plus polymer.

B: *In Vivo* Assays

Animals

[0031] Female athymic Crl:NU-Foxn1nu mice (10/ group), obtained from Charles River Laboratories (Wilmington, DE) were utilized when they were approximately 10-12 weeks of age and weighed 23-25 g. The health of the mice was assessed daily by gross observation and analyses of blood samples taken from sentinel animals housed on shared shelf

racks. All animals were allowed to acclimate and recover from any shipping-related stress for a minimum of 72 hours prior to experimental use. Autoclaved water and irradiated food (5058-ms Pico Lab mouse chow, Purina Mills, Richmond, IN) were provided *ad libitum,* and the animals were maintained on a 12 hour light and dark cycle. Cages, bedding and water bottles were autoclaved before use and changed weekly. All animal experiments were conducted in accordance with the Guide for the Care and Use of Laboratory Animals, local regulations, and protocols approved by the Roche Animal Care and Use Committee in an AAALAC accredited facility.

Tumors

[0032] SJSA cells (ATCC) were maintained in RPMI 1640 + 10% (v/v) heat-inactivated FBS + 1% (v/v) 200 nM L-glutamine. Each mouse received 5 x 10^6 cells in a 1:1 mixture of phosphate buffered saline and Matrigel in a total volume of 0.2 ml. Cells were implanted subcutaneously in the right flank using a 1 cc syringe and a 26 gauge needle.

Study Design:

[0033] The doses selected for Compound A and schedules utilized in this study are shown in Table 1 below.

Table 1 Study Design

Tumor Model	Treatment Groups
SJSA	1. Vehicle qd po 2. Compound A 7.5 mg/kg *qd po* 3. Compound A 15 mg/kg *qd po* 4. Compound A 30 mg/kg *qd po* 5. Compound A 20 mg/kg *20 days qd po, 8 days off* 6. Compound A 50 mg/kg *1x/week po* 7. Compound A 100 mg/kg *1x/week po* 8. Compound A 200 mg/kg *(given as two 100 mg/kg doses 8 hours apart (bid)), 1x/week po* 9. Compound A 50 mg/kg *4 days qd po, 10 days off x 2 cycles* 10. Compound A 50 mg/kg 2 *days qd po, 5 days off x 4 cycles* 11. Compound A 100 mg/kg 2 *days qd po, 5 days off x 4 cycles* 12. Compound A 80 mg/kg *5 days qd po, 23 days off* 13. Compound A 100 mg/kg 2 *days qd po, 12 days off x 2 cycles*

Treatment

[0034] Compound A was administered orally (po) using a 1 cc syringe and 18-gauge gavage needle (0.2 ml/animal). Treatment duration was 2-4 weeks. Dates of tumor implant, treatment initiation (study start date), and termination of treatment (study end date) can be found in Table 6 below. The starting tumor volume for this study was about 220 mm^3. Tumor volumes and animal body weights were measured three times per week and animals were monitored for clinical signs daily.

[0035] The results of this experiment are summarized Tables 1-3 below and Figures 1 and 2. As can be seen, the 5 day treatment schedule yielded the greatest per cent increase in life span (%ILS) as well as high per cent tumor growth inhibition (%TGI) with reasonable toxicity. Figure 1 also shows good growth inhibitory activity of the 5 day on/23 day off treatment schedule.

Table 2: Toxicity Summary

Group	Frequency	% Change in Body Weight at end of Study Day 29	Maximum % Weight loss	Maximum % Weight gain	# of animals ≥20% BWL	Mortality
Vehicle	QD	13.0	-1.2	13.0	0	0

(continued)

Group	Frequency	% Change in Body Weight at end of Study Day 29	Maximum % Weight loss	Maximum % Weight gain	# of animals ≥20% BWL	Mortality
Compound A 100 mg/kg	1 x/wk	9.1	4.2	9.1	0	0
Compound A 200 mg/kg (Two 100 mg/kg doses, 8 hr apart	1 x/wk	6.3	1.9	6.3	0	0
Compound A 50 mg/kg	2 on / 5 off x 4, QD	7.1	-0.8	7.1	0	0
Compound A 80 mg/kg	5 on / 23 off, QD	8.0	0.3	8.0	0	0
Compound A 20 mg/kg	20 on / 8 off, QD	1.2	-3.9	1.2	0	0
Compound A 100 mg/kg	2 on / 12 off x 2, QD	0.9	-0.6	1.8	0	0
Compound A 50 mg/kg	4 on / 10 off x 2, QD	1.2	-1.1	1.2	0	0
Compound A 15 mg/kg	QD	5.9	-2.2	5.9	0	0
Compound A 100 mg/kg	2 on / 5 off x 4, QD	1.3	-2.8	1.3	0	0
Compound A 30 mg/kg	QD	1.3	-0.2	1.3	0	0
Compound A 50 mg/kg	1 x/wk	6.6	-0.3	6.6	0	0
Compound A 7.5 mg/kg	QD	9.0	-0.3	9.0	0	0

Table 3: Efficacy Summary (left side)

Group Vehicle or Compound A	Frequency	Mean Tumor (mm3) Start Study Day:11	SEM	SD	Mean Tumor Volume (mm3) End Study Day:32	SD	SEM
Vehicle	QD	215.03	±19.00	±60.08	4696.49	±785.28	±296.91
50 mg/kg	1x/week	275.41	±22.66	±71.65	22.66	±1103.00	±348.80
7.5 mg/kg	QD	240.88	±18.01	±56.95	18.01	±956.45	±302.46
100 mg/kg	1 x / week	193.61	±9.67	±30.57	474.73	±273.78	±86.58
15 mg/kg	QD	232.37	±16.42	±51.93	16.42	±872.83	±276.01
50 mg/kg	2 on/5 off x 4, QD	203.43	±18.78	±59.39	257.29	±102.12	±32.29
80 mg/kg	5 on / 23 off, QD	197.38	±12.80	±40.48	128.05	±84.89	±26.84
20 mg/kg	20 on/8 off, QD	207.20	±16.97	±53.67	315.19	±277.51	±87.76
100 mg/kg	2 on/12 off x 2, QD	201.40	±9.86	±31.18	179.88	±154.02	±48.71
50 mg/kg	4 on / 10 off x 2, QD	213.61	±12.09	±38.23	244.70	±240.07	±75.92
100 mg/kg	2 on / 5 off x 4, QD	190.78	±25.68	±81.22	25.68	±15.82	±5.00
30 mg/kg	QD	250.86	±19.35	±61.19	19.35	±159.01	±50.28
100 mg/kg	200 mg/kg (Two 100 mg/kg doses, 8 hr apart) x 1x	224.88	±12.02	±38.02	158.95	±68.86	±21.78

Table 3: Efficacy Summary Continued (right side)

% T/C End of Study Day: 32	% Inhibition end of study Day:32	p value end of study Day: 32	Average % Regression per Group	Partial Regression	Full Regression	Animals per Group	% Increased Life Span	p Value versus Vehicle
-	-	-	-	0	0	7	-	-
43	57	<0.001	-	0	0	10	23	0.0036
34	66	<0.001	-	0	0	10	23	0.0012
6	94	<0.001	-	1	0	10	77	<0.0001
21	79	<0.001	-	0	0	10	62	<0.0001
1	99	<0.001	-	3	0	10	119	<0.0001
-2	regression	<0.001	35	6	2	10	127	<0.0001
2	98	<0.001	-	5	0	10	77	<0.0001
0	regression	<0.001	11	7	0	10	119	<0.0001
1	99	<0.001	-	6	0	10	112	<0.0001
-2	regression	<0.001	47	9	0	10	188	<0.0001
-1	regression	<0.001	13	7	0	10	127	<0.0001
-1	regression	<0.001	29	7	0	10	162	<0.0001

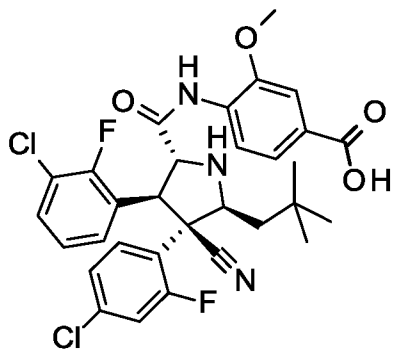
Table 4: Survival Summary

Group		50% Treatment Days	50% Vehicle days	% ILS	p value
Vehicle	QD	-	-	-	-
Compound A 100 mg/kg	1 x/wk	46	26	77	<0.0001
Compound A 200 mg/kg	Two 100 mg/kg doses, 8 hr apart 1 x / wk	68	26	162	<0.0001
Compound A 50 mg/kg	2 on / 5 off x 4, QD	57	26	119	<0.0001
Compound A 80 mg/kg	5 on / 23 off, QD	59	26	127	<0.0001
Compound A 20 mg/kg	20 on / 8 off, QD	46	26	77	<0.0001
Compound A 100 mg/kg	2 on/ 12 off x 2, QD	57	26	119	<0.0001
Compound A 50 mg/kg	4 on / 10 off x 2, QD	55	26	112	<0.0001
Compound A 15 mg/kg	QD	42	26	62	<0.0001
Compound A 100 mg/kg	2 on / 5 off x 4, QD	75	26	188	<0.0001
Compound A 30 mg/kg	QD	59	26	127	<0.0001
Compound A 50 mg/kg	1 x/wk	32	26	23	0.0036
Compound A 7.5 mg/kg	QD	32	26	23	0.0012

[0036] Overall, the 5 days on and 23 days off (5+/23-) schedule is predicted to reduce MDM2 inhibitor-induced thrombocytopenia in humans undergoing treatment for solid tumors, while still maintaining antitumor efficacy, as compared to other regimens considered.

Claims

1. A pharmaceutical product comprising as an active ingredient Compound A



(A),

for use in the treatment of cancer, **characterized by**
the daily administration of said compound A for 5 days, on days 1-5 of a treatment cycle, followed by a rest period of 23 days.

2. The pharmaceutical product for use according to claim 1, **characterized by** an administration in an amount from 800 mg/day to 3000 mg/day, daily, for up to 5 days, followed by a rest period of up to 23 days, said administration starting on the first day of a 28 day treatment cycle.

3. The pharmaceutical product for use according to claim 2, wherein Compound A is administered in an amount from 1000 mg/ day to 2500 mg/day.

4. The pharmaceutical product for use according to claim 3, wherein Compound A is administered in an amount of from 1250 mg/day to 1800 mg/day.

5. The pharmaceutical product for use according to claim 1, wherein the treatment cycle being repeated every 28 days for up to 12 cycles.

6. The pharmaceutical product for use according to claim 1, wherein Compound A is administered twice daily in equal doses.

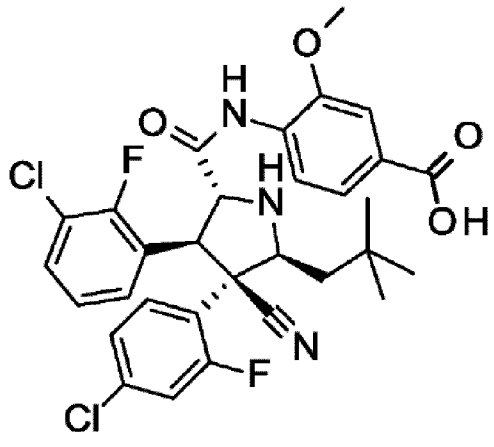
7. The pharmaceutical product for use according to claim 1 for the treatment of solid tumors.

8. The pharmaceutical product for use according to claim 1 for the treatment of colorectal cancer, prostate cancer, lung cancer, kidney cancer or breast cancer.

9. The pharmaceutical product for use according to claim 1 for the treatment of sarcoma.

Patentansprüche

1. Ein pharmazeutisches Produkt, das als einen Wirkstoff die Verbindung A umfasst



(A),

zur Verwendung bei der Behandlung von Krebs, **gekennzeichnet durch** die tägliche Verabreichung der Verbindung A für fünf Tage, an den Tagen 1 bis 5 eines Behandlungszykluses, gefolgt von einer Ruheperiode von 23 Tagen.

2. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1, **gekennzeichnet durch** eine Verabreichung in einer Menge von 800 mg/Tag bis 3000 mg/Tag, täglich, für 5 Tage, gefolgt von einer Ruheperiode von 23 Tagen, wobei die Verabreichung am ersten Tag eines 28-tägigen Behandlungszykluses beginnt.

3. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 2, wobei die Verbindung A in einer Menge von 1000 mg/Tag bis 2500 mg/Tag verabreicht wird.

4. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 3, wobei die Verbindung A in einer Menge von 1250 mg/Tag bis 1800 mg/Tag verabreicht wird.

5. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1, wobei der Behandlungszyklus alle 28 Tage für bis zu 12 Zyklen wiederholt wird.

6. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1, wobei die Verbindung A zweimal täglich in gleichen Dosen verabreicht wird.

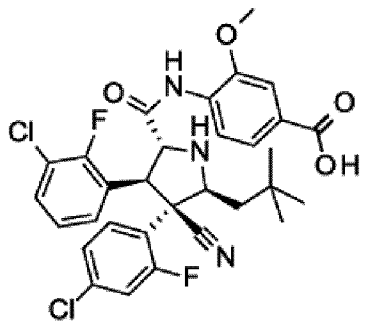
7. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1 zur Behandlung von soliden Tumoren.

8. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1 zur Behandlung von kolorektalem Krebs, Prostatakrebs, Lungenkrebs, Nierenkrebs oder Brustkrebs.

9. Das pharmazeutische Produkt zur Verwendung gemäß Anspruch 1 zur Behandlung von Sarkomen.

Revendications

1. Produit pharmaceutique comprenant comme ingrédient actif un composé A,



(A),

destiné à une utilisation dans le traitement du cancer, **caractérisé par** l'administration quotidienne dudit composé A pendant 5 jours, aux jours 1 à 5 d'un cycle de traitement, suivie d'une période d'arrêt de 23 jours.

2. Produit pharmaceutique destiné à une utilisation selon la revendication 1, **caractérisé par** une administration dans une quantité allant de 800 mg/jour à 3 000 mg/jour, chaque jour, pendant 5 jours, suivie d'une période d'arrêt de 23 jours, ladite administration commençant le premier jour d'un cycle de traitement de 28 jours.

3. Produit pharmaceutique destiné à une utilisation selon la revendication 2, dans lequel le composé A est administré dans une quantité allant de 1 000 mg/jour à 2 500 mg/jour.

4. Produit pharmaceutique destiné à une utilisation selon la revendication 3, dans lequel le composé A est administré dans une quantité allant de 1 250 mg/jour à 1 800 mg/jour.

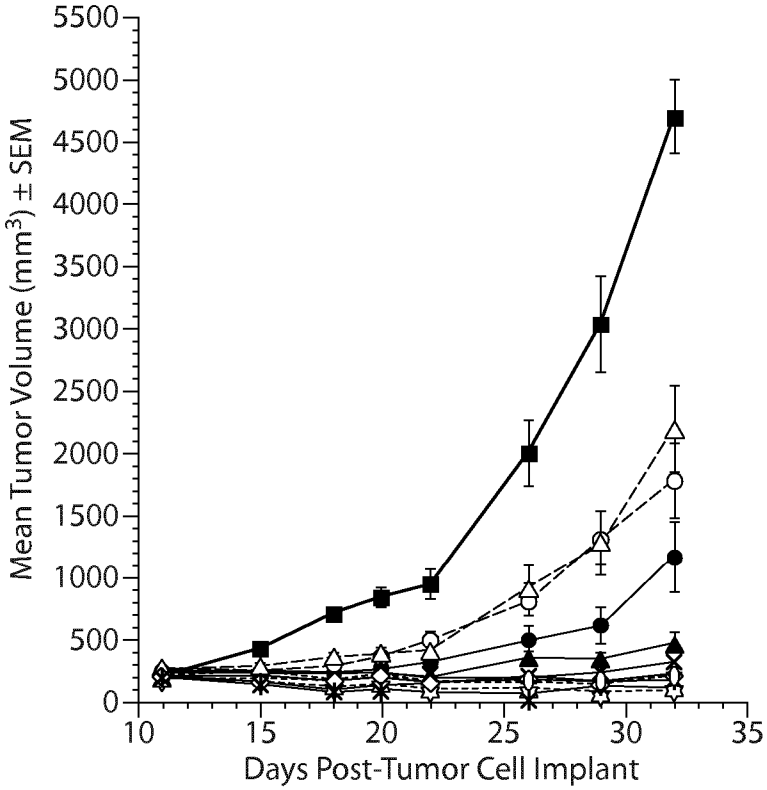
5. Produit pharmaceutique destiné à une utilisation selon la revendication 1, dans lequel le cycle de traitement est répété tous les 28 jours pendant 12 cycles au maximum.

6. Produit pharmaceutique destiné à une utilisation selon la revendication 1, dans lequel le composé A est administré deux fois par jour en des doses égales.

7. Produit pharmaceutique destiné à une utilisation selon la revendication 1, pour le traitement de tumeurs solides.

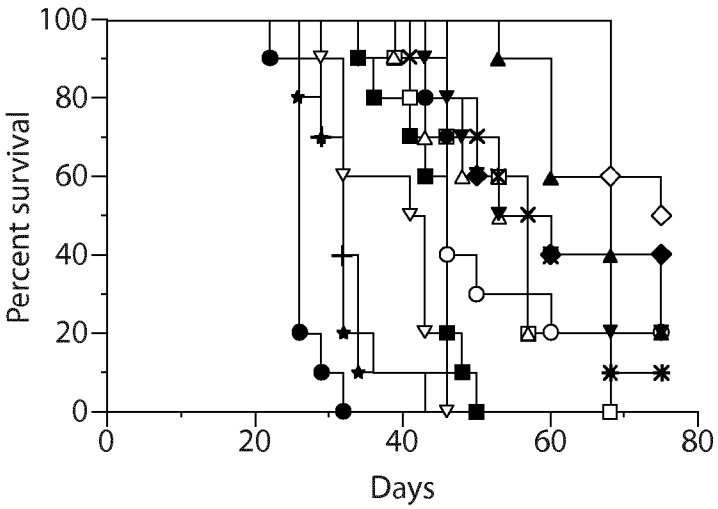
8. Produit pharmaceutique destiné à une utilisation selon la revendication 1, pour le traitement du cancer colorectal, du cancer de la prostate, du cancer du poumon, du cancer du rein ou du cancer du sein.

9. Produit pharmaceutique destiné à une utilisation selon la revendication 1, pour le traitement du sarcome.



- Vehicle QD
- Compound A 50 mg/kg 1x/wk
- Compound A 7.5 mg/kg QD
- Compound A 100 mg/kg 1x/wk
- Compound A 15 mg/kg QD
- Compound A 50 mg/kg 2 on / 5 off x 4, QD
- Compound A 80 mg/kg 5 on / 23 off, QD
- Compound A 20 mg/kg 20 on / 8 off, QD
- Compound A 100 mg/kg 2 on / 12 off x 2, QD
- Compound A 50 mg/kg 4 on / 10 off x 2, QD
- Compound A 100 mg/kg 2 on / 5 off x 4, QD
- Compound A 30 mg/kg QD
- Compound A 200 mg/kg (Two 100 mg/kg doses, 8 hr apart) 1x/wk

Fig. 1



- Vehicle, qd
- Compound A 50 mg/kg 1x/wk
- Compound A 7.5 mg/kg qd
- Compound A 100 mg/kg 1x/wk
- Compound A 15 mg/kg qd
- Compound A 50 mg/kg 2 on/5off x 4, qd
- Compound A 80 mg/kg 5 on/23 off, qd
- Compound A 20 mg/kg 20 on/8 off, qd
- Compound A 100 mg/kg 2 on/12 off x 2, qd
- Compound A 50 mg/kg 4 on/10 off x 2, qd
- Compound A 100 mg/kg 2 on/5 off x 4, qd
- Compound A 30 mg/kg qd
- Compound A 200 mg/kg (Two 100 mg/kg doses, 8 hr apart) 1x/wk

Fig. 2

REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

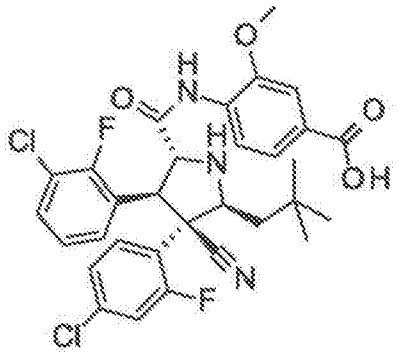
- US 20100152190 A1 **[0004]**
- WO 2011098398 A **[0004]**

Non-patent literature cited in the description

- *World Health Organization ("WHO") Handbook for Reporting Results of Cancer Treatment,* 1979 **[0009]**

Az **EP 2 827 858** lajstromszámú európai szabadalom igénypontjainak fordítása:

1. Hatóanyagként A vegyületet



(A)

tartalmazó gyógyszertermék rák kezelésében való alkalmazásra, amelyet jellemez az említett A vegyület naponta történő adagolása 5 napon át, egy kezelési ciklus 1-5 napján, amelyet egy 23 napos nyugalmi időszak követ.

2. Gyógyszertermék az 1. igénypont szerinti alkalmazásra, amelyet jellemez 800 mg/nap-tól 3000 mg/napig terjedő mennyiség naponta történő adagolása legfeljebb 5 napon át, amelyet egy legfeljebb 23 napos nyugalmi időszak követ, az említett adagolás egy 28 napos kezelési ciklus első napján kezdődik.

3. Gyógyszertermék a 2. igénypont szerinti alkalmazásra, ahol az A vegyületet 1000 mg/nap-tól 2500 mg/nap-ig terjedő mennyiségben adagoljuk.

4. Gyógyszertermék a 3. igénypont szerinti alkalmazásra, ahol az A vegyületet 1250 mg/nap-tól 1800 mg/nap-ig terjedő mennyiségben adagoljuk.

5. Gyógyszertermék az 1. igénypont szerinti alkalmazásra, ahol a kezelési ciklust minden 28 napban ismételjük legfeljebb12 cikluson át.

6. Gyógyszertermék az 1. igénypont szerinti alkalmazásra, ahol a A vegyületet naponta két alkalommal adagoljuk azonos dózisokban.

7. Gyógyszertermék az 1. igénypont szerinti alkalmazásra szilárd tumorok kezelésére.

8. Gyógyszertermék az 1. igénypont szerinti alkalmazásra vastag-végbélrák, prosztatarák, tüdőrák, veserák vagy emlőrák kezelésére.

9. Gyógyszertermék az 1. igénypont szerinti alkalmazásra szarkóma kezelésére.

A meghatalmazott:

165/1696

Dr. Miskolczi Mária