



Office de la Propriété
Intellectuelle
du Canada

Un organisme
d'Industrie Canada

Canadian
Intellectual Property
Office

An agency of
Industry Canada

CA 2694389 C 2016/01/19

(11)(21) **2 694 389**

(12) **BREVET CANADIEN
CANADIAN PATENT**

(13) **C**

(86) **Date de dépôt PCT/PCT Filing Date:** 2008/07/03
(87) **Date publication PCT/PCT Publication Date:** 2009/02/05
(45) **Date de délivrance/Issue Date:** 2016/01/19
(85) **Entrée phase nationale/National Entry:** 2010/01/22
(86) **N° demande PCT/PCT Application No.:** EP 2008/058601
(87) **N° publication PCT/PCT Publication No.:** 2009/015981
(30) **Priorité/Priority:** 2007/08/01 (EP07113603.0)

(51) **Cl.Int./Int.Cl. A61K 31/4745** (2006.01),
A61P 35/00 (2006.01)
(72) **Inventeurs/Inventors:**
PISANO, CLAUDIO, IT;
VESCI, LOREDANA, IT;
CARMINATI, PAOLO, IT
(73) **Propriétaire/Owner:**
SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE
S.P.A, IT
(74) **Agent:** FETHERSTONHAUGH & CO.

(54) **Titre : TRAITEMENT DE TUMEURS PEDIATRIQUES**
(54) **Title: TREATMENT OF PEDIATRIC TUMORS**

(57) **Abrégé/Abstract:**

A subclass of camptothecin derivatives is disclosed to be useful for the preparation of a medicament for the treatment of pediatric tumors such as for example rhabdomyosarcoma, primitive neuroectodermal tumors (PNET) and neuroblastoma.



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 February 2009 (05.02.2009)

(10) International Publication Number
WO 2009/015981 A3

(51) International Patent Classification:

A61K 31/4745 (2006.01) A61P 35/00 (2006.01)

(21) International Application Number:

PCT/EP2008/058601

(22) International Filing Date:

3 July 2008 (03.07.2008)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

07113603.0 1 August 2007 (01.08.2007) EP

(71) Applicant: SIGMA-TAU Industrie Farmaceutiche Riunite S.p.A. [IT/IT]; Viale Shakespeare, 47, I-00144 Rome (IT).

(72) Inventors; and

(75) Inventors/Applicants (for US only): PISANO, Claudio [IT/IT]; Via Cherubini, 60/B, I-04011 Aprilia (It) (IT). VESCI, Loredana [IT/IT]; Via Elio Petri, 3, I-00128 Rome (IT). CARMINATI, Paolo [IT/IT]; Via Agnello, 19, I-20121 Milan (IT).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ,

EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

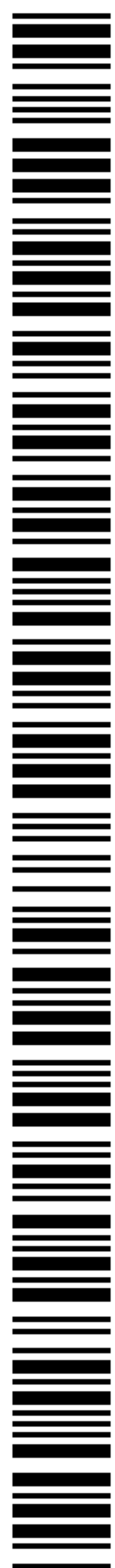
- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))

(88) Date of publication of the international search report:

3 September 2009

(54) Title: TREATMENT OF PEDIATRIC TUMORS

(57) Abstract: A subclass of camptothecin derivatives is disclosed to be useful for the preparation of a medicament for the treatment of pediatric tumors such as for examplerhabdomyosarcoma, primitive neuroectodermal tumors (PNET) and neuroblastoma.



WO 2009/015981 A3

TREATMENT OF PEDIATRIC TUMORS

FIELD OF THE INVENTION

The present invention relates to the use of a subclass of camptothecin
5 derivatives for the preparation of a medicament for the treatment of pediatric
tumors such as, for example, rhabdomyosarcoma, primitive neuroectodermal
tumors (PNET) and neuroblastoma.

BACKGROUND OF THE INVENTION

Camptothecin derivatives are DNA-topoisomerase I inhibitors that have
10 emerged as a prominent class of anticancer agents. Together with the taxanes,
the topoisomerase I inhibitors are presumably the most important new class of
anticancer drugs introduced into clinical practice. Pre-clinical studies demonstrated
significant *in vitro* and *in vivo* activity of topoisomerase I inhibitors, such as
camptothecin and its derivatives, on a broad range of tumors. The results from
15 clinical trials were promising, as shown by the registration of two topoisomerase
inhibitors, topotecan and irinotecan (also known as CPT-11), in many European
countries and in the USA, for treatment of patients with ovarian and colorectal
cancer, respectively. Other derivatives are currently at different stages of clinical
development.

20 In patent application EP1044977 and in J. Med. Chem. 2001, 44, 3264-3274,
camptothecin derivatives are described which bear an alkyloxime O-substituted at
position 7 and which are endowed with antitumor activity higher than the
compound of reference topotecan. Moreover these camptothecin derivatives
bearing an imino group on position 7, also show an improved therapeutic index.

Among these compounds one of the preferred molecules was shown to be 7-t-butoxyiminomethylcamptothecin (CPT 184, also known as ST1481 or gimatecan).

Although the annual mortality rate of pediatric cancer has decreased over the past two decades, the proportion of deaths from tumors of the central nervous
5 system in the same population has increased from 18 percent to 30 percent. The cause of childhood brain tumors is largely unknown. While radiation exposure is a recognized risk factor for brain tumors, the role of other environmental toxins is unclear in children. Less than 5 percent of pediatric brain tumors are associated with a known genetic disease, such as neurofibromatosis, a common genetic
10 condition associated with benign tumor growths on nerve tissue (Serletis D, Parkin P, Bouffet E, Shroff M, Drake JM, Rutka JT J Neurosurg. 2007 106: 363-7).

Although only 1 percent of childhood brain tumors are detected at birth or in the first few months of life, a significant number are diagnosed before age five, suggesting a developmental defect. As a matter of fact, defects in developmental
15 growth signaling pathways have recently been identified in embryonal tumors.

Historically, a diagnosis of brain cancer is provided by a pathologist, who views tissue samples under a microscope. Upon visual inspection of brain cells (histology), pathologists can then classify the tumor type (Miller CR, Perry A Arch Pathol Lab Med. 2007 131: 397-406). The limitations of this practice are that
20 many brain tumors have a similar histology when they are actually very different tumors with greatly different prognoses and responses to therapies. There are some children diagnosed with medulloblastoma who respond well to therapy while others do not. Therefore, tumor classification is moving toward the use of molecular signatures to more precisely classify and grade tumor tissues (Sardi I, Cavalieri D, Massimino M Paediatr Drugs. 2007 9:81-96).

In addition to confusing tumor classifications, in the past, pediatric tumors were considered to be similar to tumors in adults. However, recent studies have revealed that pediatric brain tumors are very different biologically than their adult counterparts. One example is a tumor called fibrillary astrocytoma, a tumor that
5 occurs both in children and adults (*Collins Nat Clin Pract Oncol. 2007 4: 362-74*).

Biologically they behave very differently even though they look the same under the microscope. This disease in children rarely will become a high-grade tumor during childhood years, but in adults it can turn into higher grade tumors.

These observations are further supported by recent studies of molecular
10 markers. Mutations in specific genes that cause disease in adults may not be the cause of disease in children. Future studies should provide fertile opportunities for drug target discoveries and related molecularly targeted therapies.

The use of surgery in treatment of pediatric brain tumors is well-established, but more effective treatments are needed. Imaging technologies have been used
15 to non-invasively assess tumor status and treatment in children, thus eliminating the need to obtain repeated biopsies of the same tumor. The gains achieved in improved surgical resection of brain tumors also can be attributed to improved imaging technologies (Khatua S, Jalali R *Pediatr Hematol Oncol.* 2005 22: 361-71). Surgeons are now better able to locate a tumor and assess the margins,
20 removing less of the normal brain tissue. This is a significant improvement because there is a direct correlation between the extent of tumor resection and survival in some types of brain tumors in pediatric patients. Improved imaging technologies have also spurred advances in radiation therapy techniques (Greco C, Wolden S. *Cancer.* 2007 109: 1227-38). In addition to providing information
25 about the size and location of a tumor, imaging techniques are also providing data

to evaluate the biochemical profile of the tumor, as well. Studies have shown that changes in the ratio of certain biochemical components of a tumor can aid an oncologist in determining if a tumor is actively growing (Chen L, Madura K. Cancer Res. 2005 65:5599-606; Chung TD, Broaddus WC Mol Interv. 2005 5:15-9). These results would support the choice of aggressive or less stringent treatment regiments.

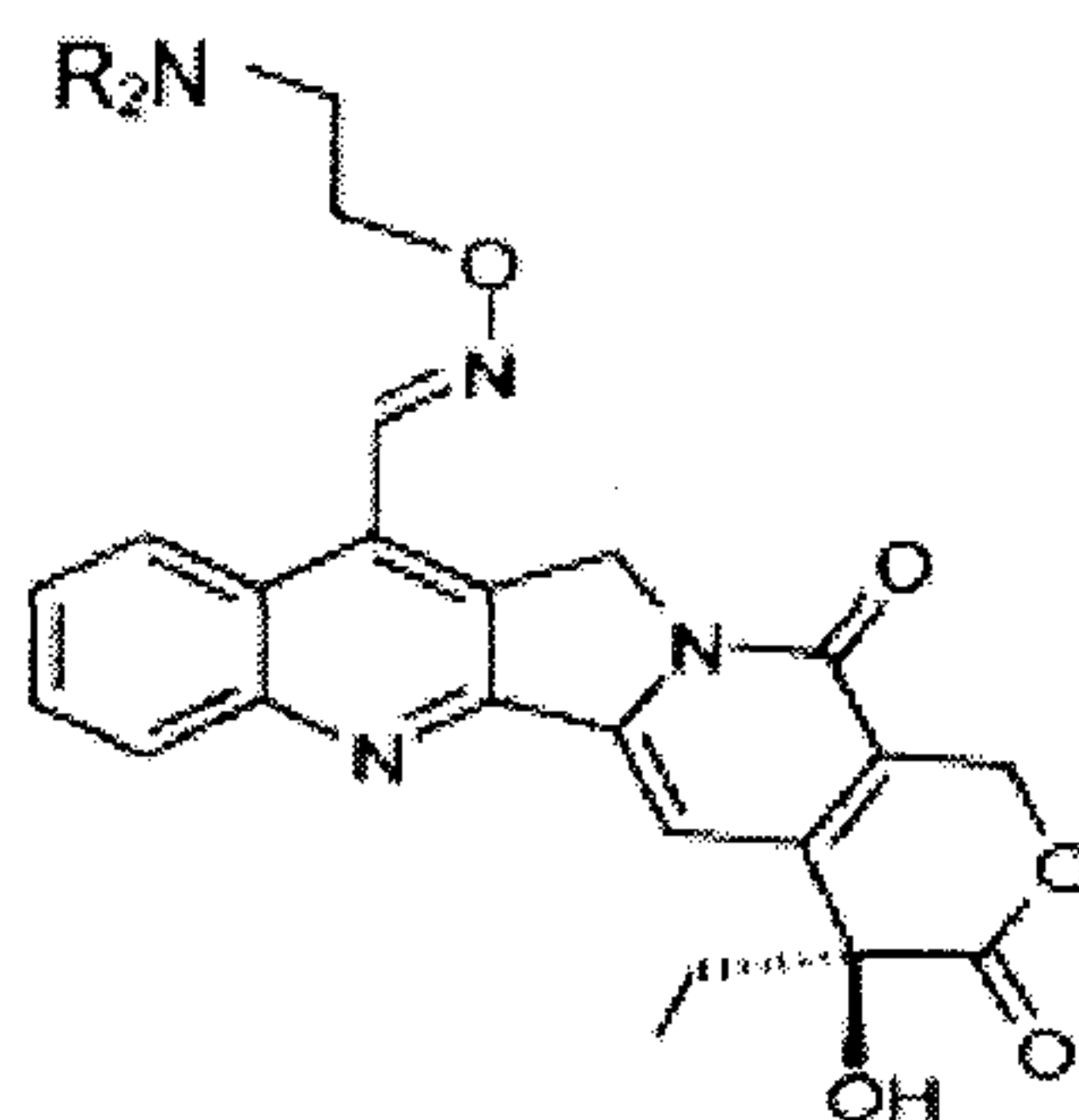
Blood flow to tumors and tissues in the brain is also being examined through imaging. Evaluation of changes in the amount of blood flowing to tissues in the brain is essential to assess the effectiveness of anti-angiogenic drugs. These drugs do not target the tumor directly, but attack the cells lining the blood vessels that support tumor growth. The tumor size may not shrink, but a decrease in the number of blood vessels surrounding the tumor is a significant advance (Kieran MW J Neurooncol. 2005 75: 327-34; Kibble A. IDrugs. 2007 10: 5-7). Anti-angiogenic drugs will most likely be combined with other agents that target the tumor directly. Currently, these agents are being tested alone for safety.

Neuroblastoma is one of the most common extracranial solid tumours in childhood with a poor prognosis in its advanced stage. Treatment failure is often associated to the occurrence of drug resistance. To date, treatment of paediatric neuroblastoma is still dismal, and therefore novel effective drugs are awaited. (Gutierrez JC, Fischer AC, Sola JE, Perez EA, Koniaris LG Pediatr Surg Int. 2007 23: 637-46). Rhabdomyosarcoma is the most common soft-tissue sarcoma of childhood, representing 5% of all childhood cancers (M. Beth McCarville, Sheri L. Spunt and Alberto S. Pappo AJR 2001; 176:1563-1569). It is thought to arise from primitive mesenchymal cells committed to skeletal muscle differentiation and can occur in a variety of organs and tissues, including those that lack striated muscle.

Primitive neuroectodermal tumors (PNETs) develop from primitive or undifferentiated neuroepithelial cells from the early development of the nervous system. PNET of the posterior fossa, or medulloblastoma, is the most common brain tumor in children. In 80% of cases, patients with PNETs develop acute hydrocephalus accompanied by severe symptoms of headache and vomiting, and they require urgent resection of the mass (*de Bont JM, den Boer ML, Kros JM, Passier MM, Reddingius RE, Smitt PA, Luider TM, Pieters R. J Neuropathol Exp Neurol. 2007 66: 505-516*).

SUMMARY OF INVENTION

The present invention provides use of a compound of Formula (1),

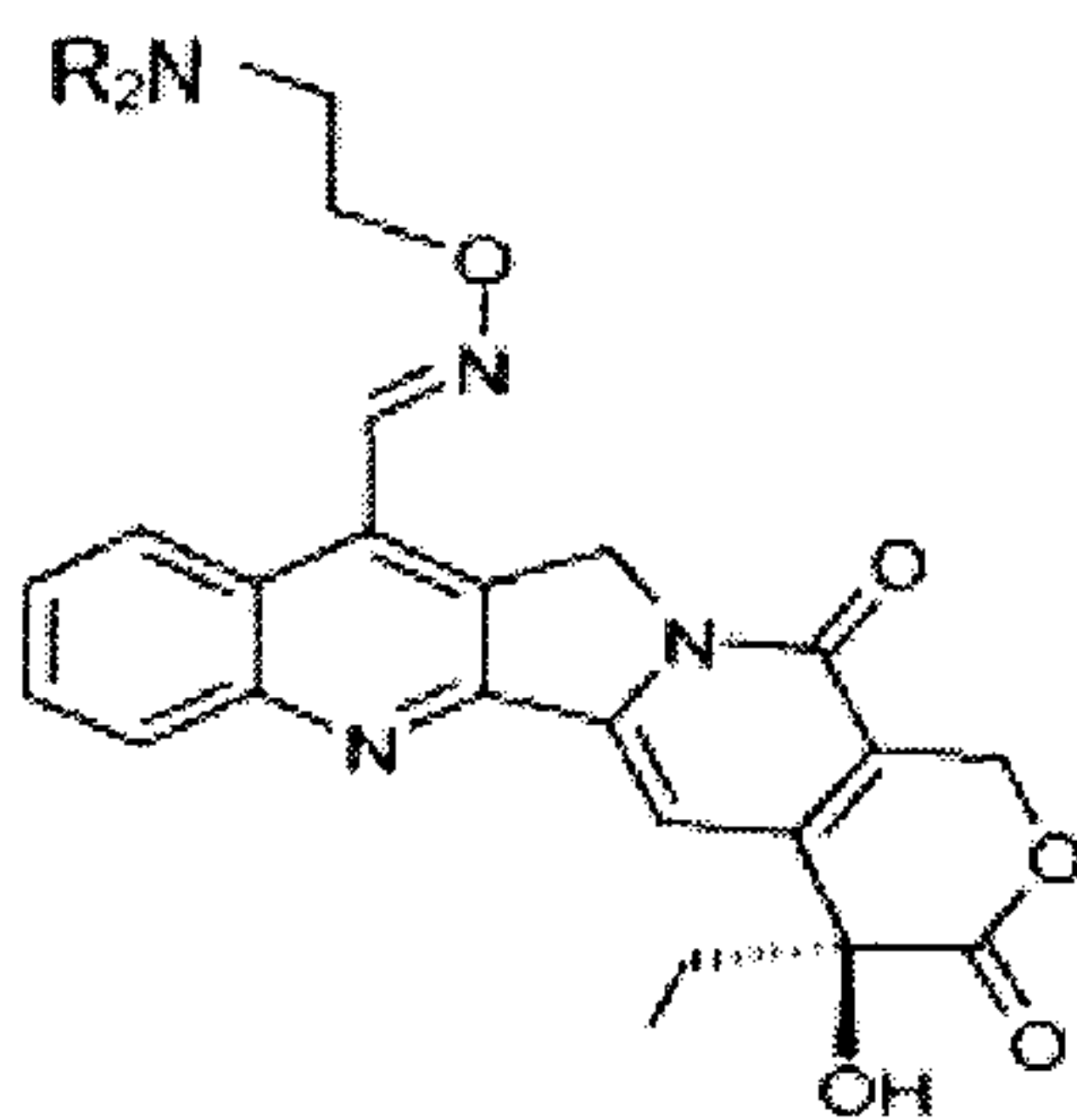


(I)

wherein R is hydrogen or C₁-C₄ alkyl, for the treatment of a pediatric tumor, or for the manufacture of a medicament for the treatment of a pediatric tumor.

5a

The present invention also provides a compound of Formula (1),



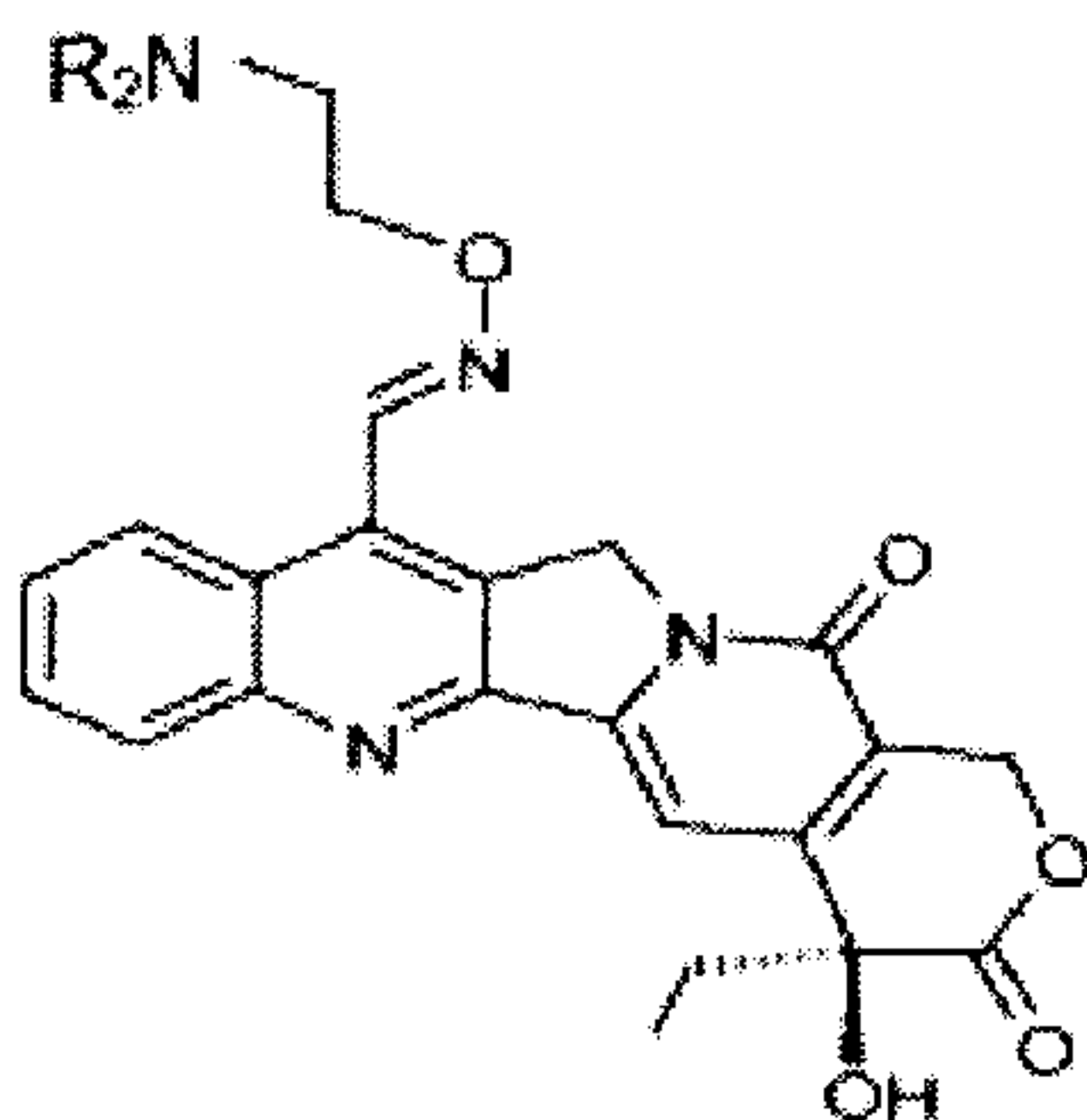
(I)

wherein R is hydrogen or C₁-C₄ alkyl, for use in the treatment of a pediatric tumor

DESCRIPTION OF THE INVENTION

It has now surprisingly found that two camptothecin derivatives have shown a superior efficacy with respect to the reference compound (irinotecan) in terms of tumor volume inhibition (TVI%), against a wide spectrum of pediatric tumors xenografts in vivo. Moreover, the treatment was generally well tolerated by the animals, with no signs of acute or delayed toxicity.

In particular the main object of the present invention is the use of a compound of Formula I,



(I)

where R is hydrogen or C₁-C₄ alkyl,

for the preparation of a medicament for the treatment of pediatric tumors such as, for example, rhabdomyosarcoma, primitive neuroectodermal tumors (PNET) and neuroblastoma.

5 Compounds of Formula (I) also comprise tautomers, geometrical isomers, optically active forms as enantiomers, diastereomers and racemate forms, as well as pharmaceutically acceptable salts of the compounds of Formula (I).

Preferred pharmaceutically acceptable salts of the Formula (I) are acid addition salts formed with pharmaceutically acceptable acids like hydrochloride, hydrobromide, sulfate or bisulfate, phosphate or hydrogen phosphate, acetate, 10 benzoate, succinate, fumarate, maleate, lactate, citrate, tartrate, gluconate, methanesulfonate, benzenesulfonate, and para-toluenesulfonate salts.

Preferably R is hydrogen or methyl.

Preferred compounds of Formula (I) are:

15 7-(2-amino)ethoxyiminomethylcamptothecin, (ST1968, also known as CPT188) and 7-(2-dimethylamino)ethoxyiminomethylcamptothecin (ST1969).

The compounds of Formula (I) may be prepared from readily available starting materials using the following general methods and procedures. It will be appreciated that where typical or preferred experimental conditions (i.e. reaction 20 temperatures, time, moles of reagents, solvents, etc.) are given, other experimental conditions can also be used, unless otherwise stated. Optimum reaction conditions may vary with the particular reactants or solvents used, but such conditions can be determined by one skilled in the art by routine optimisation procedures. Specific reference is made to the methods described in patent 25 application EP1044977 and in J. Med. Chem. 2001, 44, 3264-3274.

A method of treating a patient suffering from tumor pathology selected from the group consisting of rhabdomyosarcoma, primitive neuroectodermal tumors (PNET) and neuroblastoma comprising administering a therapeutically effective amount of a compound of Formula (I) as described above represents one of the aspects of the present invention.

The term "therapeutically effective amount" as used herein refers to an amount of a therapeutic agent needed to treat, ameliorate a targeted disease or condition, or to exhibit a detectable therapeutic effect.

For any compound, the therapeutically effective dose can be estimated initially either in cell culture assays, for example, of neoplastic cells, or in animal models, usually mice or rats.

The animal model may also be used to determine the appropriate concentration range and route of administration. Such information can then be used to determine useful doses and routes for administration in humans.

The precise effective amount for a human subject will depend upon the severity of the disease state, general health of the subject, age, weight, and gender of the subject, diet, time and frequency of administration, drug combination(s), reaction sensitivities, and tolerance/response to therapy. This amount can be determined by routine experimentation and is within the judgement of the clinician. Generally, an effective dose will be from 0.01 mg/kg to 100 mg/kg, preferably 0.05 mg/kg to 50 mg/kg. Compositions may be administered individually to a patient or may be administered in combination with other agents, drugs or hormones.

The medicament may also contain a pharmaceutically acceptable carrier, for administration of a therapeutic agent. Such carriers include antibodies and other polypeptides, genes and other therapeutic agents such as liposomes,

provided that the carrier does not itself induce the production of antibodies harmful to the individual receiving the composition, and which may be administered without undue toxicity.

Suitable carriers may be large, slowly metabolised macromolecules such as
5 proteins, polysaccharides, polylactic acids, polyglycolic acids, polymeric amino acids, amino acid copolymers and inactive virus particles.

A thorough discussion of pharmaceutically acceptable carriers is available in Remington's Pharmaceutical Sciences (Mack Pub. Co. , N. J.1991).

Pharmaceutically acceptable carriers in therapeutic compositions may
10 additionally contain liquids such as water, saline, glycerol and ethanol. Additionally, auxiliary substances, such as wetting or emulsifying agents, pH buffering substances, and the like, may be present in such compositions. Such carriers enable the pharmaceutical compositions to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions, and the like, for
15 ingestion by the patient.

Once formulated, the compositions of the invention can be administered directly to the subject. The subjects to be treated can be animals; in particular, human pediatric subjects can be treated.

The medicament of this invention may be administered by any number of
20 routes including, but not limited to, oral, intravenous, intramuscular, intra- arterial, intramedullary, intrathecal, intraventricular, transdermal or transcutaneous applications, subcutaneous, intraperitoneal, intranasal, enteral, topical, sublingual, intravaginal, rectal means or locally on the diseased tissue after surgical operation.

Dosage treatment may be a single dose schedule or a multiple dose
25 schedule.

The invention will now be illustrated in greater detail by means of non-limiting Examples.

EXAMPLES

IN VIVO ANTITUMORAL ACTIVITY ON DIFFERENT PEDIATRIC TUMOR

5 XENOGRAFT MODELS

The antitumor effect was evaluated against different human pediatric tumor models: SK-ND-AS (neuroblastoma); PSFK (primitive neuroectodermal tumors/PNET); TE-671 (rhabdomyosarcoma). ST 1968 was evaluated in comparison to CPT-11 (irinotecan) by intravenously delivery of 30 mg/10 ml/kg for
10 ST1968 and 50 mg/10 ml/kg for CPT-11 according to the schedule q4dx4.

Methods

Exponentially growing tumor cells were injected s.c. into nude athymic mice. The number of tumor cells was previously chosen by a growth curve. Mice were housed inside cages of makrolon (33.2 x 15 x 13 cm) with stainless steel cover-
15 feed and sterilized and dust-free bedding cobs. Animals were housed under a light-dark cycle, keeping temperature and humidity constant. Parameters of the animal rooms were assessed as follows: 22±2 °C temperature, 55±10% relative humidity, about 15-20 filtered air changes/hour and 12 hour circadian cycle of artificial light (7 a.m., 7 p.m.). At request, the environmental conditions were
20 monitored and the data are retained in Animal Housing Archives. Drinking water was supplied ad libitum. Each mouse was offered daily a complete pellet diet (GLP 4RF21, Mucedola) throughout the study. The analytical certificates of animal food and water are retained at Sigma-Tau premises. All animals were weighed before starting the experiment and were subdivided into the different dosage groups.

Each cage was identified by a paper tag indicating: cage number, group, date of tumor injection, starting date of treatment, name of the test item, dose and route of administration, date of sacrifice.

Tumor growth was followed by biweekly measurements of tumor diameters with a Vernier caliper. Tumor volume (TV, mm³) was calculated as: [length (mm) x width (mm)²]/2, where the width and the length are the shortest and the longest diameters of each tumor, respectively.

The efficacy of the drug treatment was assessed as tumor volume inhibition (TVI%) in treated versus control mice, calculated as: 100- [(mean tumor volume of treated animals/mean tumor volume of control animals)x100].

The toxicity of the drug treatments was determined as: body weight loss percent (% BWL max) = 100- (mean BW_{day x}/mean BW_{day 1} x 100), where BW_x is the mean BW at the day of maximal loss during the treatment and BW₁ is the mean BW on the 1st day of treatment.

Results

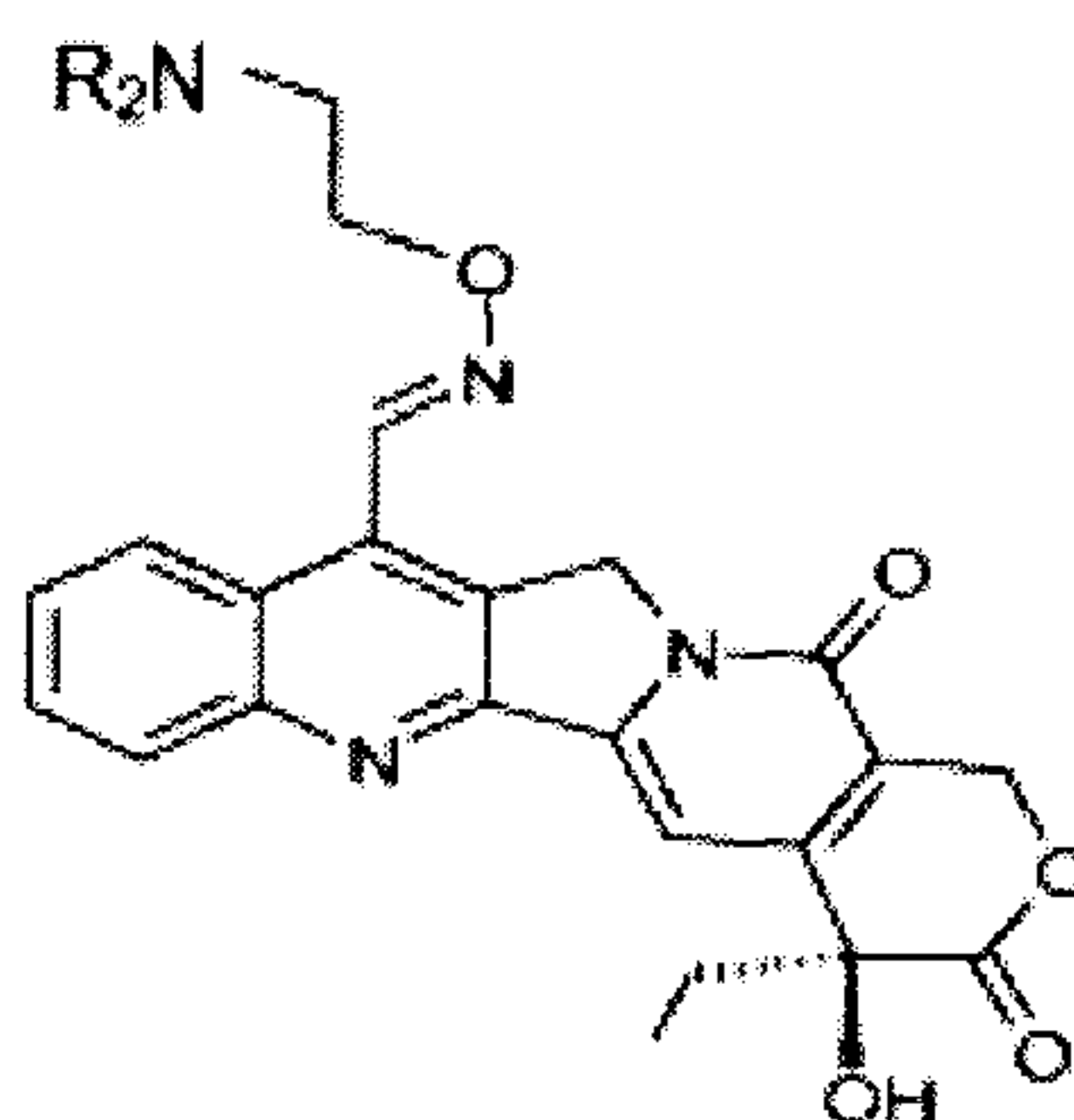
In the *in vivo* studies, ST1968 showed a remarkable antitumor activity with an appreciable rate of complete tumor regressions in all models compared with CPT-11. The most impressive antitumor effects were observed against PNET and rabdomiosarcoma models, as documented by the high rate of complete responses: the tumor volume inhibition (TWI) was 99% or 100% respectively (Table 1). Interestingly, ST1968 treatment was generally well tolerated by mice with no signs of acute or delayed toxicity.

Table 1. Antitumor activity of ST1968 on human pediatric tumor xenograft models

Tumor	Line	Compound	Dose (mg/kg)	Method of administ.	TVI%
Rhabdomyosarcoma	TE-671	ST1968	30	q4dx4	99
		irinotecan	50	q4dx4	73
Primitive neuroectodermal tumors (PNET)	PFSK	ST1968	30	q4dx4	100
		irinotecan	50	q4dx4	95
Neuroblastoma	SK-N-AS	ST1968	30	q4dx4	98
		irinotecan	50	q4dx4	91

CLAIMS

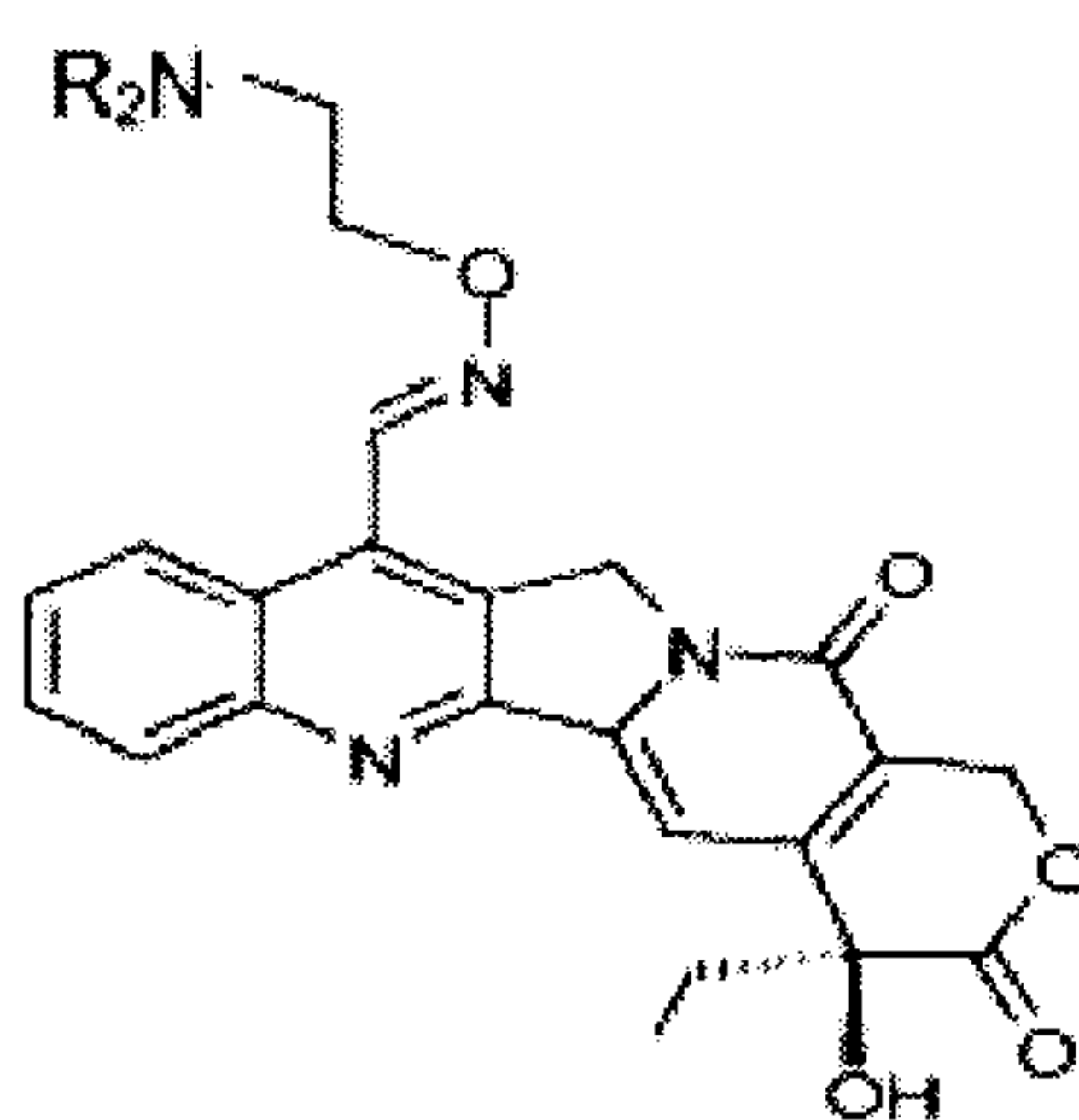
1. Use of a compound of Formula (1),



(I)

wherein R is hydrogen or C_1 - C_4 alkyl,
for the treatment of a pediatric tumor.

2. Use of a compound of Formula (1),



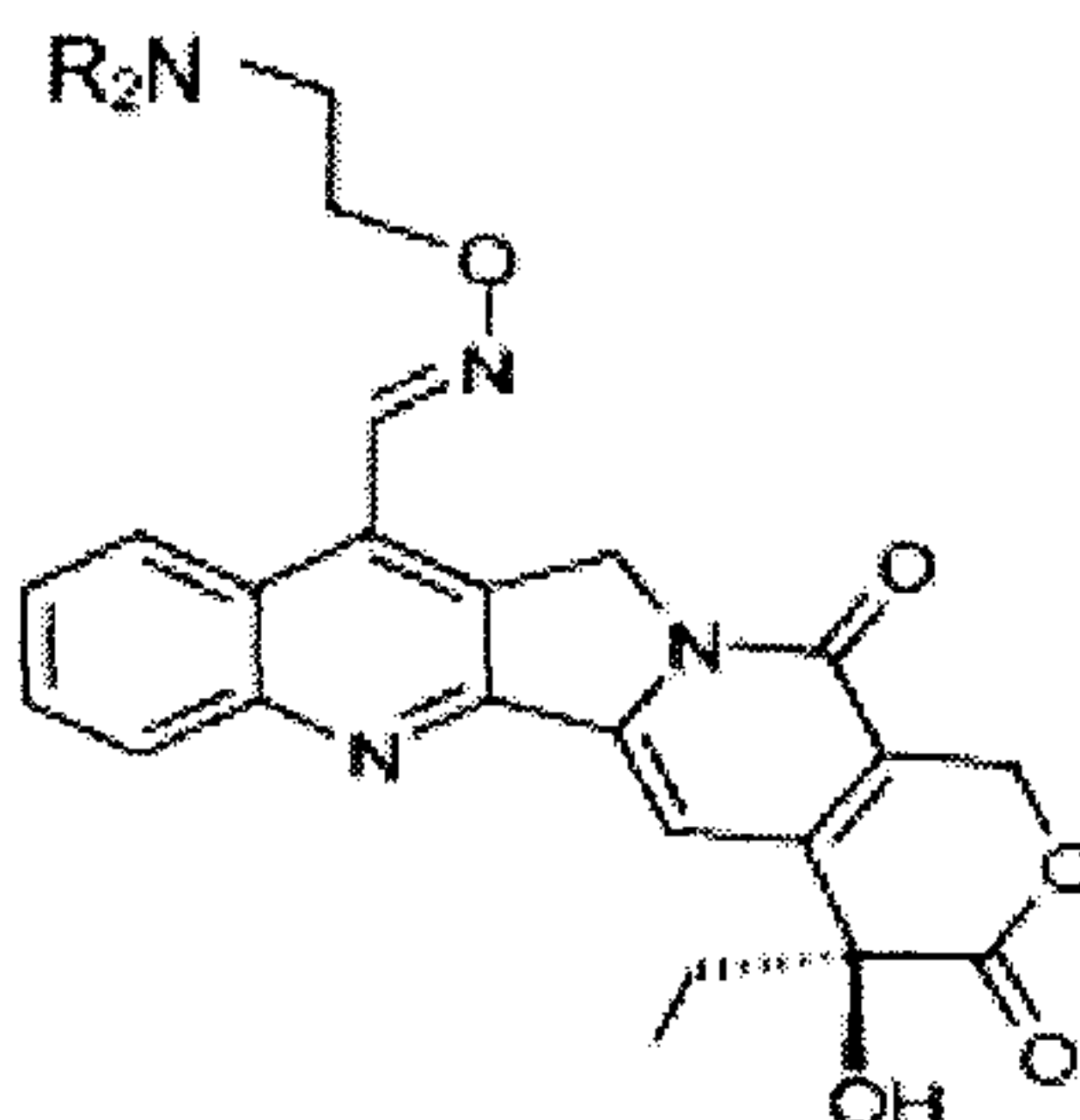
(I)

wherein R is hydrogen or C_1 - C_4 alkyl,
in the manufacture of a medicament for the treatment of a pediatric tumor.

3. The use according to claim 1 or 2, wherein R is hydrogen or methyl.

4. The use according to any one of claims 1 to 3, wherein the pediatric tumor is a head and neck carcinoma selected from the group consisting of: rhabdomyosarcoma, primitive neuroectodermal tumor (PNET) and neuroblastoma.

5. A compound of Formula (1),



(I)

wherein R is hydrogen or C₁-C₄ alkyl,
for use in the treatment of a pediatric tumor.

6. The compound according to claim 5, wherein R is hydrogen or methyl.

7. The compound according to claim 5 or 6, wherein the pediatric tumor is a head and neck carcinoma selected from the group consisting of: rhabdomyosarcoma, primitive neuroectodermal tumor (PNET) and neuroblastoma.