



(86) Date de dépôt PCT/PCT Filing Date: 2008/02/11

(87) Date publication PCT/PCT Publication Date: 2008/08/21

(85) Entrée phase nationale/National Entry: 2009/07/07

(86) N° demande PCT/PCT Application No.: US 2008/001755

(87) N° publication PCT/PCT Publication No.: 2008/100449

(30) Priorités/Priorities: 2007/02/09 (US60/900,338);
2007/05/24 (US60/924,663)

(51) Cl.Int./Int.Cl. *A61K 38/18* (2006.01),
A61K 31/00 (2006.01), *A61K 31/05* (2006.01),
A61K 31/335 (2006.01), *A61K 31/35* (2006.01),
A61K 31/365 (2006.01), *A61K 31/395* (2006.01),
A61K 31/4015 (2006.01), *A61K 45/00* (2006.01),
A61P 25/00 (2006.01), *A61P 25/28* (2006.01)

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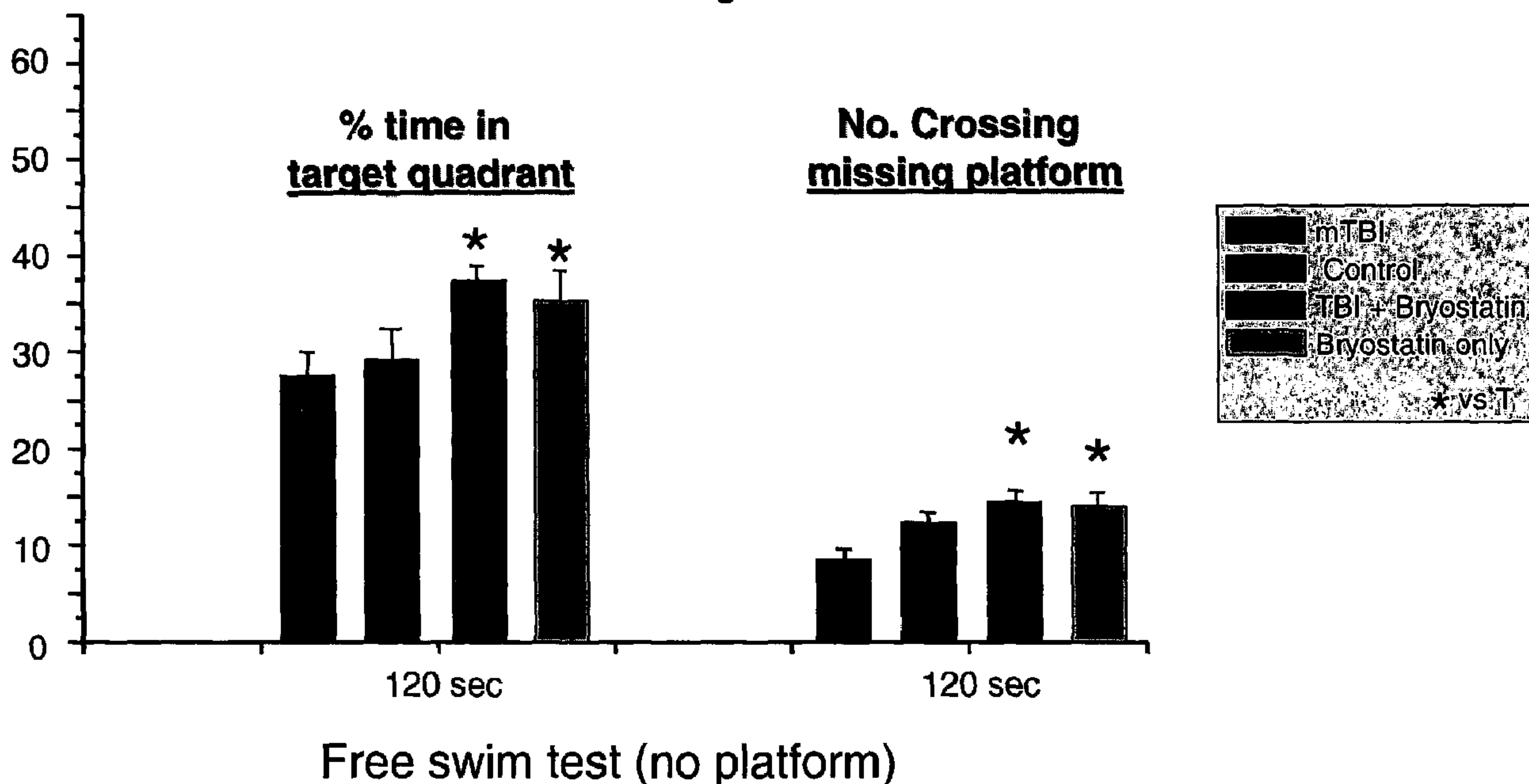
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(54) Titre : EFFETS THERAPEUTIQUES DES BRYOSTATINES, DE LEURS ANALOGUES, ET D'AUTRES
SUBSTANCES CONNEXES SUR L'AFFAIBLISSEMENT DE LA MEMOIRE INDUITE PAR UN TRAUMATISME
CRANIEN ET SUR LES LESIONS CEREBRALES

(54) Title: THERAPEUTIC EFFECTS OF BRYOSTATINS, BRYOLOGS, AND OTHER RELATED SUBSTANCES ON
HEAD TRAUMA-INDUCED MEMORY IMPAIRMENT AND BRAIN INJURY

Figure 3



(57) Abrégé/Abstract:

The invention provides for the use of protein kinase activators or boosters of nerve growth factor (NGF), brain-derived neurotrophic factor (BDNF) or other neurotrophic factors to treat head trauma. Specifically, the present invention provides methods of treating head trauma comprising the steps of identifying a subject having suffered a head trauma and administering to said subject an amount of a pharmaceutical composition comprising a protein kinase C (PKC) activator or 4-methylcatechol acetic acid (MCBA) and a pharmaceutically acceptable carrier effective to treat at least one symptom of head trauma.

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

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
21 August 2008 (21.08.2008)

PCT

(10) International Publication Number
WO 2008/100449 A4

(51) International Patent Classification:

A61K 38/18 (2006.01) **A61K 31/395** (2006.01)
A61K 31/00 (2006.01) **A61K 31/4015** (2006.01)
A61K 31/05 (2006.01) **A61K 45/00** (2006.01)
A61K 31/335 (2006.01) **A61P 25/00** (2006.01)
A61K 31/35 (2006.01) **A61P 25/28** (2006.01)
A61K 31/365 (2006.01)

(21) International Application Number:

PCT/US2008/001755

(22) International Filing Date:

11 February 2008 (11.02.2008)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

60/900,338 9 February 2007 (09.02.2007) US
 60/924,663 24 May 2007 (24.05.2007) US

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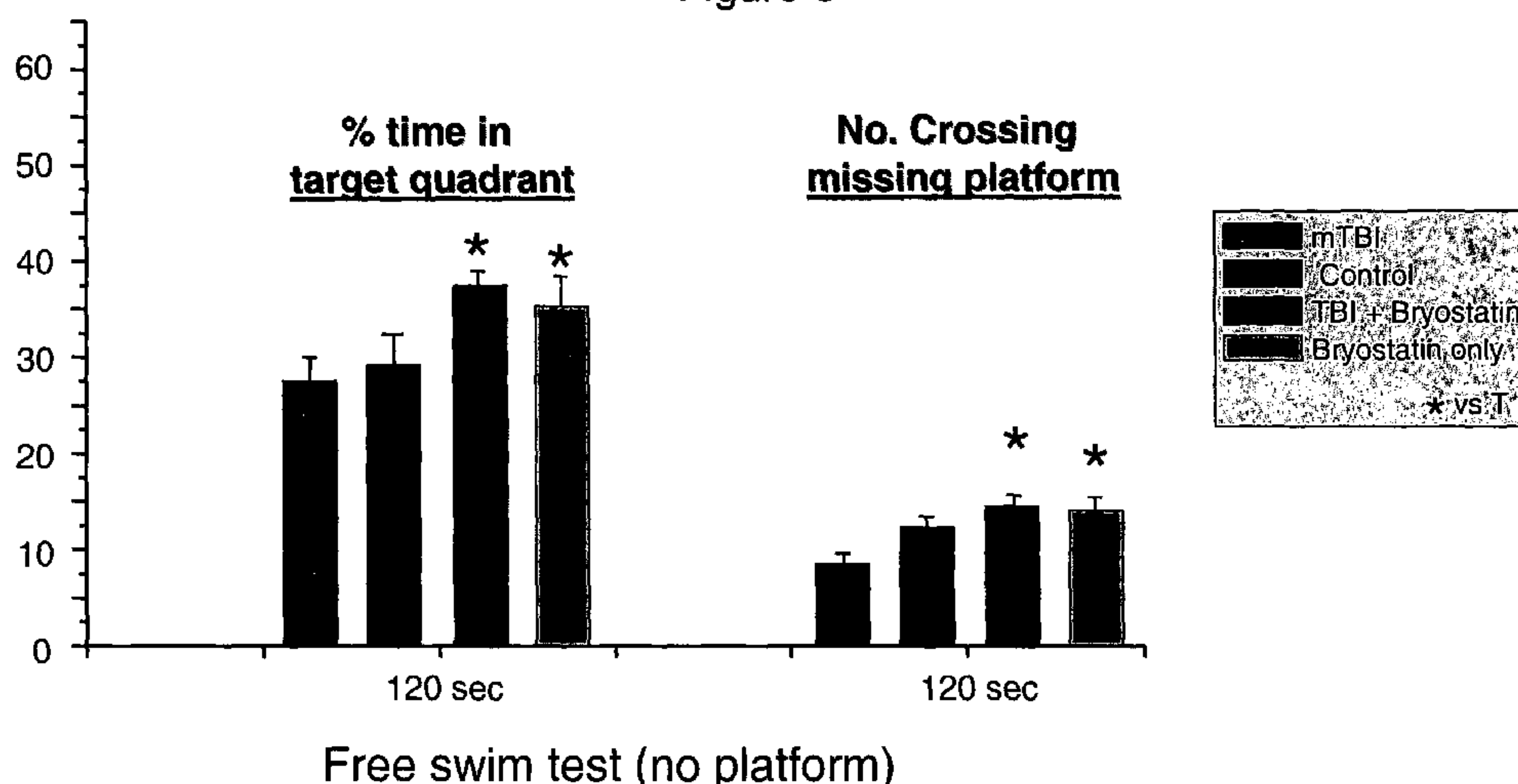
(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, 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, 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,

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WO 2008/100449 A4

WO 2008/100449 A4



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).

— *with amended claims*

Published:

— *with international search report*

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

22 May 2009

Date of publication of the amended claims:

30 July 2009

**THERAPEUTIC EFFECTS OF BRYOSTATINS, BRYOLOGS, AND OTHER RELATED
SUBSTANCES ON HEAD TRAUMA-INDUCED MEMORY IMPAIRMENT AND
BRAIN INJURY**

Inventors: Ofer Zohar and Daniel L. Alkon

This application claims benefit to U. S. Provisional Application Serial No. 60/900,339, filed on February 9, 2007 and U. S. Provisional Application Serial No. 60/924,662, filed on May 24, 2007, all of which are hereby incorporated herein by reference in their entireties.

FIELD OF THE INVENTION

[0001] The present invention relates to the treatment of head trauma with compounds that activate protein kinase C (PKC) or boost nerve growth factor (NGF), brain-derived neurotrophic factor (BDNF) or other neurotrophic factors.

BACKGROUND OF THE INVENTION

A. Head Trauma

[0002] Head injury is a trauma to the head, that may or may not include injury to the brain (see also brain injury). The incidence (number of new cases) of head injury is 300 per 100,000 per year (0.3% of the population), with a mortality of 25 per 100,000 in North America and 9 per 100,000 in Britain. Head trauma is a common cause of childhood hospitalization.

[0003] Head injuries include both injuries to the brain and those to other parts of the head, such as the scalp and skull. Head injuries may be closed or open. A closed (non-missile) head injury is one in which the skull is not broken. A penetrating head injury occurs when an object pierces the skull and breaches the dura mater. Brain injuries may be diffuse, occurring over a wide area, or focal, located in a small, specific area. A head injury may cause a skull fracture, which may or may not be associated with injury to the brain. Some patients may have linear or depressed skull fractures. If intracranial hemorrhage, or bleeding within the brain occurs, a hematoma within the skull can put pressure on the brain. Types of intracranial hematoma include subdural, subarachnoid, extradural, and intraparenchymal hematoma.

Craniotomy surgeries are used in these cases to lessen the pressure by draining off blood. Head trauma is caused by a concussive event.

[0004] Brain injury can be at the site of impact, but can also be at the opposite side of the skull due to a contrecoup effect (the impact to the head can cause the brain to move within the skull, causing the brain to impact the interior of the skull opposite the head-impact). If the impact causes the head to move, the injury may be worsened, because the brain may ricochet inside the skull (causing additional impacts), or the brain may stay relatively still (due to inertia) but be hit by the moving skull.

B. Protein Kinase C

[0005] PKC has been identified as one of the largest gene families of non-receptor serine-threonine protein kinases. Since the discovery of PKC in the early eighties by Nishizuka and coworkers (Kikkawa *et al.* (1982) *J. Biol. Chem.* 257: 13341), and its identification as a major receptor for phorbol esters (Ashendel *et al.* (1983) *Cancer Res.*, 43: 4333), a multitude of physiological signaling mechanisms have been ascribed to this enzyme. The intense interest in PKC stems from its unique ability to be activated *in vitro* by calcium and diacylglycerol (and its phorbol ester mimetics), an effector whose formation is coupled to phospholipid turnover by the action of growth and differentiation factors.

[0006] The activation of PKC has been shown to improve learning and memory. (U.S. Patent Application Serial Nos. PCT/US02/13784; PCT/US03/07102; 60/287,721; 60/362,081; 10/172,005; and 10/476,459; each incorporated herein by reference in its entirety). Prior to the present disclosure, however, the PKC-mediated improvement of learning and memory has not been recognized as a mechanism for the treatment of post-head trauma memory deficits and brain injury. Also, the PKC activators disclosed herein, specifically those compounds that improve learning and memory, were not recognized as possessing brain function-restoring activity after head trauma.

[0007] Head trauma therapy has historically been limited to few treatment options available. Although many types of potential neuroprotectants have been tested in clinical trials, none has been approved for clinical use, because of ineffectiveness especially when used post-head trauma or associated toxicity. The compounds presented in this invention disclosure were

effective when the treatment was started one hour after the head trauma in the animal model at doses that have already been demonstrated to be well tolerated in humans (the bryostatin-1 doses). Compounds that target the protein kinase C (PKC) such as bryostatin-1, a direct PKC activator, and methylcatechol diacetic acid, a derivative of methylcatechol, an enhancer of nerve growth factor (NGF), brain-derived neurotrophic factor (BDNF) or other neurotrophic factors, which is perhaps one of the PKC targets, have been found to have therapeutic value against brain injury and memory impairment induced with head trauma. The development of these substances as therapeutic in the treatment of head trauma is provided by this invention.

SUMMARY OF THE INVENTION

[0008] The present invention provides methods of treating head trauma comprising the steps of identifying a subject having suffered a head trauma and administering to said subject an amount of a pharmaceutical composition comprising a protein kinase C (PKC) activator or 4-methylcatechol acetic acid (MCBA) and a pharmaceutically acceptable carrier effective to treat at least one symptom of head trauma.

[0009] In one embodiment, the PKC activator is FGF-18, a macrocyclic lactone, a benzolactam, a pyrrolidinone, or a combination thereof. In a preferred embodiment, the macrocyclic lactone is a bryostatin or neristatin. In another embodiment, the neristatin is neristatin-1. In another embodiment, the bryostatin is bryostatin-1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17 or 18. More preferably, the bryostatin is bryostatin-1.

[0010] In another preferred embodiment, the pharmaceutical composition comprises 4-methylcatechol acetic acid (MCBA), other derivatives of methylcatechol, or a brain derived neurotrophic factor. MCBA and other derivatives of methylcatechol activate or upregulate nerve growth factor (NGF), brain derived neurotrophic factor (BDNF) or other neurotrophic factors. NGF activates, upregulates or enhances the activity of PKC which in turn upregulates, activates or enhances NGF.

[0011] In one embodiment, administration of the pharmaceutical compositions of the present invention is initiated within 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, or 14 days of said head trauma. In another embodiment, said administration is initiated between 1 and 2 days, 1 and 3

days, 1 and 4 days, 1 and 5 or 1 and 7 days of said head trauma. In another embodiment, the administration of the pharmaceutical compositions of the present invention is initiated within 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, or 24 hours of said head trauma. In yet another embodiment, the administration of the pharmaceutical compositions of the present invention is initiated between 1 and 3, 1 and 5, 1 and 10, 1 and 24, 3 and 5, 3 and 10, 3 and 24, 5 and 10, 5 and 24, or 10 and 24 hours after said head trauma. In yet another embodiment, the administration of the pharmaceutical compositions of the present invention is initiated after 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, or 24 hours after said head trauma. In yet another embodiment, the administration of the pharmaceutical compositions of the present invention is initiated after 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or 21 days after said head trauma.

[0012] In one embodiment, treatment comprising the administration of the pharmaceutical compositions of the present invention is continued for a duration of about 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 weeks.

BRIEF DESCRIPTION OF THE FIGURE

[0013] **Figure 1** depicts the escape latencies of mice after minimal traumatic brain injury (TBI)(30g) followed by 30ug/kg bryostatin injected intraperitoneally.

[0014] **Figure 2** depicts the escape latencies of mice after minimal traumatic brain injury (TBI) (30g):followed by 20ug/kg bryostatin injected intraperitoneally.

[0015] **Figure 3** depicts memory retention in mice after minimal TBI (30g) followed by 20ug/kg bryostatin injected intraperitoneally.

DETAILED DESCRIPTION OF THE INVENTION

A. Definitions

[0016] As used herein, "administration" of a composition includes any route of administration, including oral subcutaneous, intraperitoneal, and intramuscular.

[0017] As used herein, "an effective amount" is an amount sufficient to reduce one or more symptoms associated with a head trauma.

[0018] As used herein, "protein kinase C activator" or "PKC activator" means a substance that increases the rate of the reaction catalyzed by protein kinase C by binding to the protein kinase C.

[0019] As used herein, the term "subject" means a mammal.

[0020] As used herein, the term "pharmaceutically acceptable carrier" means a chemical composition with which the active ingredient may be combined and which, following the combination, can be used to administer the active ingredient to a subject. As used herein, the term "physiologically acceptable" ester or salt means an ester or salt form of the active ingredient which is compatible with any other ingredients of the pharmaceutical composition, which is not deleterious to the subject to which the composition is to be administered.

[0021] As used herein, "pharmaceutically acceptable carrier" also includes, but is not limited to, one or more of the following: excipients; surface active agents; dispersing agents; inert diluents; granulating and disintegrating agents; binding agents; lubricating agents; sweetening agents; flavoring agents; coloring agents; preservatives; physiologically degradable compositions such as gelatin; aqueous vehicles and solvents; oily vehicles and solvents; suspending agents; dispersing or wetting agents; emulsifying agents, demulcents; buffers; salts; thickening agents; fillers; emulsifying agents; antioxidants; antibiotics; antifungal agents; stabilizing agents; and pharmaceutically acceptable polymeric or hydrophobic materials. Other "additional ingredients" which may be included in the pharmaceutical compositions of the invention are known in the art and described, for example in Genaro, ed., 1985, Remington's Pharmaceutical Sciences, Mack Publishing Co., Easton, Pa., which is incorporated herein by reference.

[0022] The formulations of the pharmaceutical compositions described herein may be prepared by any method known or hereafter developed in the art of pharmacology. In general, such preparatory methods include the step of bringing the active ingredient into association with a carrier or one or more other accessory ingredients, and then, if necessary or desirable, shaping or packaging the product into a desired single- or multi-dose unit.

[0023] Although the descriptions of pharmaceutical compositions provided herein are principally directed to pharmaceutical compositions which are suitable for ethical administration to humans, it will be understood by the skilled artisan that such compositions are generally suitable for administration to animals of all sorts. Modification of pharmaceutical compositions suitable for administration to humans in order to render the compositions suitable for administration to various animals is well understood, and the ordinarily skilled veterinary pharmacologist can design and perform such modification with merely ordinary, if any, experimentation. Subjects to which administration of the pharmaceutical compositions of the invention is contemplated include, but are not limited to, humans and other primates, and other mammals.

[0024] Despite progress toward the development of new therapeutic agents and availability of several animal models, there is still a pressing need for improved animal models for screening

B. Animal Models of Learning and Memory

[0025] The area of memory and learning impairment is rich in animal models that are able to demonstrate different features of memory and learning processes. (*See, for example*, Hollister, L.E., 1990, *Pharmacopsychiat.*, 23, (Suppl II) 33-36). The available animal models of memory loss and impaired learning involve measuring the ability of animals to remember a discrete event. These tests include the Morris Water Maze and the passive avoidance procedure. In the Morris Water Maze, animals are allowed to swim in a tank divided into four quadrants, only one of which has a safety platform beneath the water. The platform is removed and the animals are tested for how long they search the correct quadrant verse the incorrect quadrants. In the passive avoidance procedure the animal remembers the distinctive environment in which a mild electric shock is delivered and avoids it on a second occasion. A variant of the passive avoidance procedure makes use of a rodent's preference for dark enclosed environments over light open ones. Further discussion can be found in Crawley, J. N., 1981, *Pharmacol. Biochem. Behav.*, 15, 695-699; Costall, B. et al, 1987, *Neuropharmacol.*, 26, 195-200; Costall, B. et al., 1989, *Pharmacol. Biochem. Behav.*, 32, 777-785; Barnes, J.M. et al., 1989, *Br. J. Pharmacol.*, 98 (Suppl) 693P; Barnes, J.M. et al., 1990, *Pharmacol. Biochem. Behav.*, 35, 955-962.

C. Protein Kinase C (PKC)

[0026] The PKC gene family consists presently of 11 genes which are divided into four subgroups: 1) classical PKC α , β_1 , β_2 (β_1 and β_2 are alternatively spliced forms of the same gene) and γ , 2) novel PKC δ , ϵ , η , and θ , 3) atypical PKC ζ , λ , ι and κ and 4) PKC μ . PKC μ resembles the novel PKC isoforms but differs by having a putative transmembrane domain (reviewed by Blohe *et al.* (1994) *Cancer Metast. Rev.* 13: 411; Ilug *et al.* (1993) *Biochem J.* 291: 329; Kikkawa *et al.* (1989) *Ann. Rev. Biochem.* 58: 31). The α , β_1 , β_2 and γ isoforms are Ca^{2+} , phospholipid and diacylglycerol-dependent and represent the classical isoforms of PKC, whereas the other isoforms are activated by phospholipid and diacylglycerol but are not dependent on Ca^{2+} . All isoforms encompass 5 variable (V1-V5) regions, and the α , β and γ isoforms contain four (C1-C4) structural domains which are highly conserved. All isoforms except PKC α , β and γ lack the C2 domain, the λ , ι and κ isoforms also lack nine of two cysteine-rich zinc finger domains in C1 to which diacylglycerol binds. The C1 domain also contains the pseudosubstrate sequence which is highly conserved among all isoforms, and which serves an autoregulatory function by blocking the substrate-binding site to produce an inactive conformation of the enzyme (House *et al.* (1987) *Science* 238, 1726).

[0027] Because of these structural features, diverse PKC isoforms are thought to have highly specialized roles in signal transduction in response to physiological stimuli (Nishizuka (1989) *Cancer* 10: 1892), as well as in neoplastic transformation and differentiation (Glazer (1994) *Protein Kinase C*, J.F. Kuo, ed., Oxford U. Press at pages 171-198). For a discussion of known PKC modulators see PCT/US97/08141, U.S. Patent Nos. 5,652,232; 6,080,784; 5,891,906; 5,962,498; 5,955,501; 5,891,870 and 5,962,504 (each incorporated herein by reference in its entirety).

[0028] There is increasing evidence that the individual PKC isozymes play different, sometimes opposing, roles in biological processes, providing two directions for pharmacological exploitation. One is the design of specific (preferably, isozyme specific) inhibitors of PKC. This approach is complicated by the fact that the catalytic domain is not the domain primarily responsible for the isotype specificity of PKC. The other approach is to develop isozyme-selective, regulatory site-directed PKC activators. These may provide a way to override the

effect of other signal transduction pathways with opposite biological effects. Alternatively, by inducing down-regulation of PKC after acute activation, PKC activators may cause long term antagonism. Bryostatin is currently in clinical trials as an anti-cancer agent. The bryostatins are known to bind to the regulatory domain of PKC and to activate the enzyme. Bryostatins are examples of isozyme-selective activators of PKC. (see for example WO 97/43268; incorporated herein by reference in its entirety). For a discussion of known PKC modulators see PCT/US97/08141, U.S. Patent Nos. 5,652,232; 6,043,270; 6,080,784; 5,891,906; 5,962,498; 5,955,501; 5,891,870 and 5,962,504 (each of which is incorporated herein by reference in its entirety).

[0029] Several classes of PKC activators have been identified. Phorbol esters, however, are not suitable compounds for eventual drug development because of their tumor promotion activity, (Ibarreta *et al.* (1999) *Neuro Report* 10(5&6): 1035-40). Of particular interest are macrocyclic lactones (i.e. bryostatin class and neristatin class) that act to stimulate PKC. Of the bryostatin class compounds, bryostatin-1 has been shown to activate PKC and proven to be devoid of tumor promotion activity. Bryostatin-1, as a PKC activator, is also particularly useful since the dose response curve of bryostatin-1 is biphasic. Additionally, bryostatin-1 demonstrates differential regulation of PKC isozymes, including PKC α , PKC δ and PKC ϵ . Bryostatin-1 has undergone toxicity and safety studies in animals and humans and is actively investigated as an anti-cancer agent. Bryostatin-1's use in the studies has determined that the main adverse reaction in humans is myalgia. One example of an effective dose is 20 or 30 μ g/kg per dose by intraperitoneal injection.

[0030] Several classes of PKC activators have been identified. Phorbol esters, however, are not suitable compounds for eventual drug development because of their tumor promotion activity, (Ibarreta *et al.* (1999) *Neuro Report* 10(5&6): 1035-40). Of particular interest are macrocyclic lactones (i.e. bryostatin class and neristatin class) that act to stimulate PKC. Of the bryostatin class compounds, bryostatin-1 has been shown to activate PKC and proven to be devoid of tumor promotion activity. Bryostatin-1, as a PKC activator, is also particularly useful since the dose response curve of bryostatin-1 is biphasic. Additionally, bryostatin-1 demonstrates differential regulation of PKC isozymes, including PKC α , PKC δ and PKC ϵ . Bryostatin-1 has undergone toxicity and safety studies in animals and humans and is actively

investigated as an anti-cancer agent. Bryostatin-1's use in the studies has determined that the main adverse reaction in humans is myalgia. One example of an effective dose is 20 or 30 µg/kg per dose by intraperitoneal injection.

[0031] Macrocyclic lactones, and particularly bryostatin-1, are described in U.S. Patent 4,560,774 (incorporated herein by reference in its entirety). Macrocyclic lactones and their derivatives are described elsewhere in U.S. Patent 6,187,568, U.S. Patent 6,043,270, U.S. Patent 5,393,897, U.S. Patent 5,072,004, U.S. Patent 5,196,447, U.S. Patent 4,833,257, and U.S. Patent 4,611,066 (each incorporated herein by reference in its entirety). The above patents describe various compounds and various uses for macrocyclic lactones including their use as an anti-inflammatory or anti-tumor agent. (Szallasi *et al.* (1994) *Journal of Biological Chemistry* 269(3): 2118-24; Zhang *et al.* (1996) *Cancer Research* 56: 802-808; Hennings *et al.* (1987) *Carcinogenesis* 8(9): 1343-1346; Varterasian *et al.* (2000) *Clinical Cancer Research* 6: 825-828; Mutter *et al.* (2000) *Bioorganic & Medicinal Chemistry* 8: 1841-1860)(each incorporated herein by reference in its entirety).

[0032] As will also be appreciated by one of ordinary skill in the art, macrocyclic lactone compounds and their derivatives, particularly the bryostatin class, are amenable to combinatorial synthetic techniques and thus libraries of the compounds can be generated to optimize pharmacological parameters, including, but not limited to efficacy and safety of the compositions. Additionally, these libraries can be assayed to determine those members that preferably modulate α -secretase and/or PKC.

[0033] Combinatorial libraries high throughput screening of natural products and fermentation broths has resulted in the discovery of several new drugs. At present, generation and screening of chemical diversity is being utilized extensively as a major technique for the discovery of lead compounds, and this is certainly a major fundamental advance in the area of drug discovery. Additionally, even after a "lead" compound has been identified, combinatorial techniques provide for a valuable tool for the optimization of desired biological activity. As will be appreciated, the subject reaction readily lend themselves to the creation of combinatorial libraries of compounds for the screening of pharmaceutical, or other biological or medically-related activity or material-related qualities. A combinatorial library for the purposes of the

present invention is a mixture of chemically related compounds, which may be screened together for a desired property; said libraries may be in solution or covalently linked to a solid support. The preparation of many related compounds in a single reaction greatly reduces and simplifies the number of screening processes that need to be carried out. Screening for the appropriate biological property may be done by conventional methods. Thus, the present invention also provides methods for determining the ability of one or more inventive compounds to bind to effectively modulate α -secretase and/or PKC.

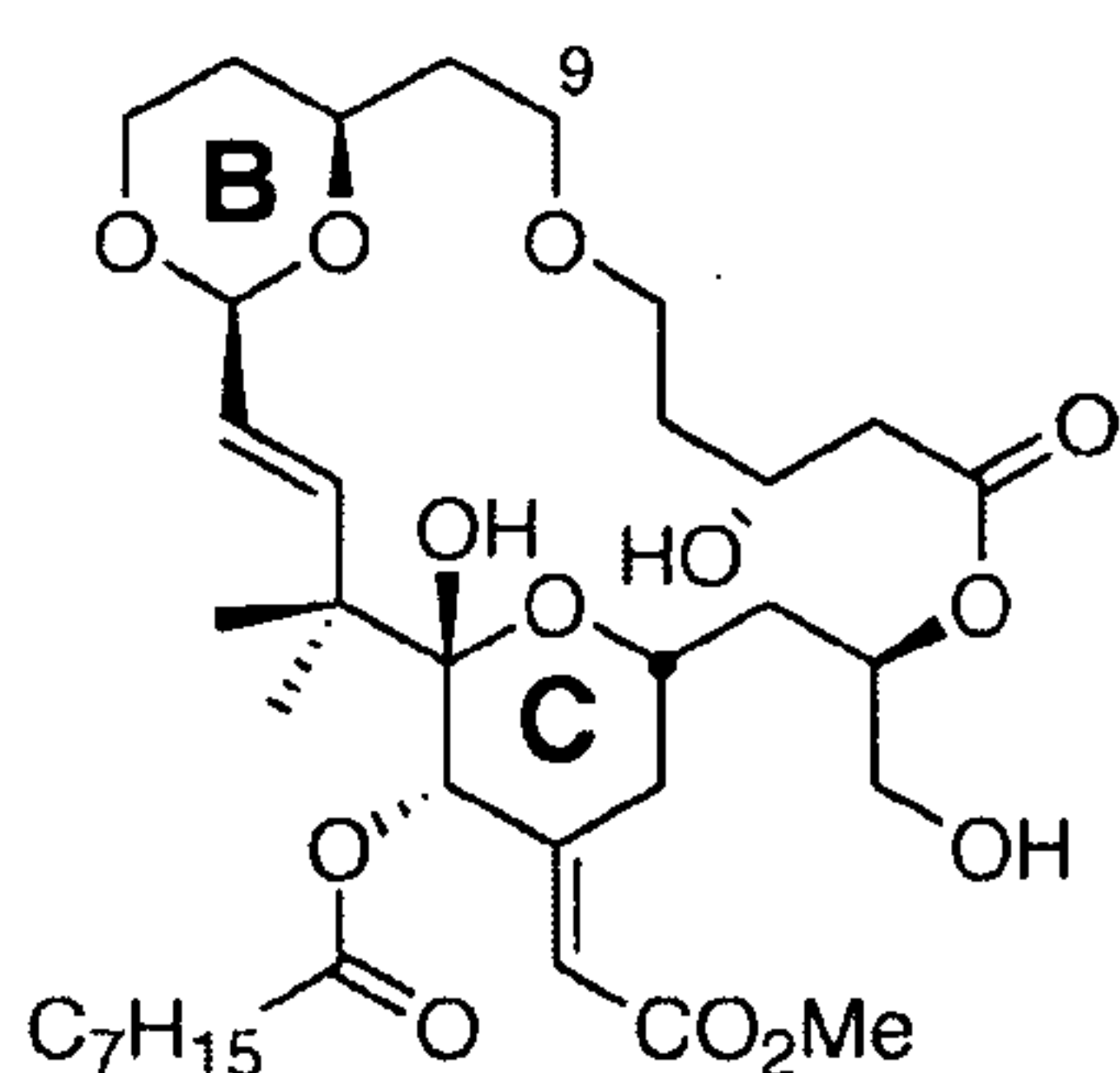
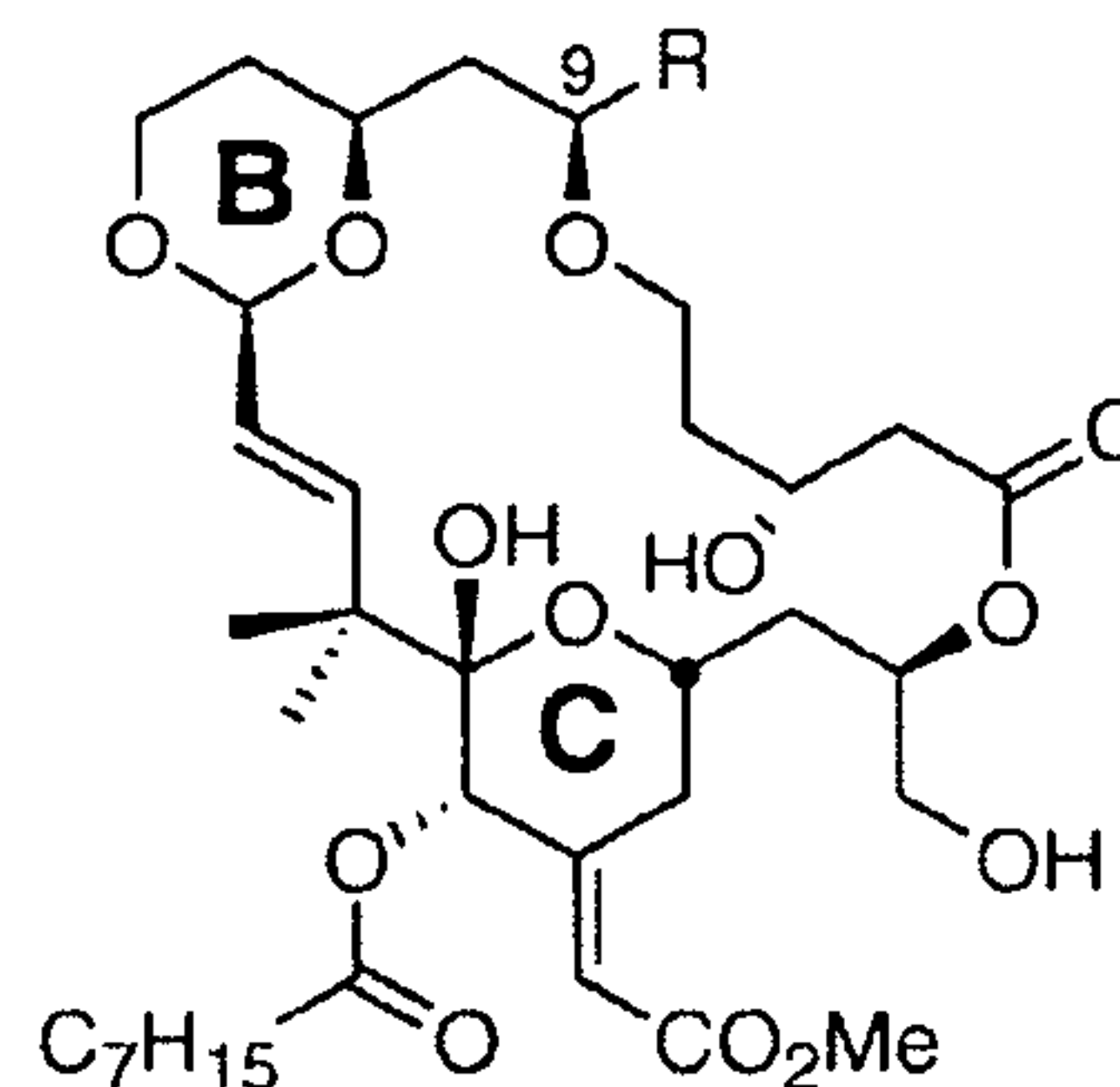
[0034] A variety of techniques are available in the art for generating combinatorial libraries described below, but it will be understood that the present invention is not intended to be limited by the foregoing examples and descriptions. (*See, for example*, Blondelle *et al.* (1995) *Trends Anal. Chem.* 14: 83; U.S. Patents 5,359,115; 5,362,899; U.S. 5,288,514; PCT publication WO 94/08051; Chen *et al.* (1994) *JACCS* 116:266-271; Kerr *et al.* (1993) *JACCS* 115:252; PCT publications W092/10092, W093/09668; W091/07087; and W093/20242; each of which is incorporated herein by reference). Accordingly, a variety of libraries on the order of about 16 to 1,000,000 or more diversomers can be synthesized and screened for a particular activity or property.

[0035] Analogs of bryostatin, commonly referred to as bryologs, are one particular class of PKC activators that are suitable for use in the methods of the present invention. The following Table summarizes structural characteristics of several bryologs, demonstrating that bryologs vary greatly in their affinity for PKC (from 0.25 nM to 10 μ M). Structurally, they are all similar. While bryostatin-1 has two pyran rings and one 6-membered cyclic acetal, in most bryologs one of the pyrans of bryostatin-1 is replaced with a second 6-membered acetal ring. This modification reduces the stability of bryologs, relative to bryostatin-1, for example, in both strong acid or base, but has little significance at physiological pH. Bryologs also have a lower molecular weight (ranging from about 600 to 755), as compared to bryostatin-1 (988), a property which facilitates transport across the blood-brain barrier.

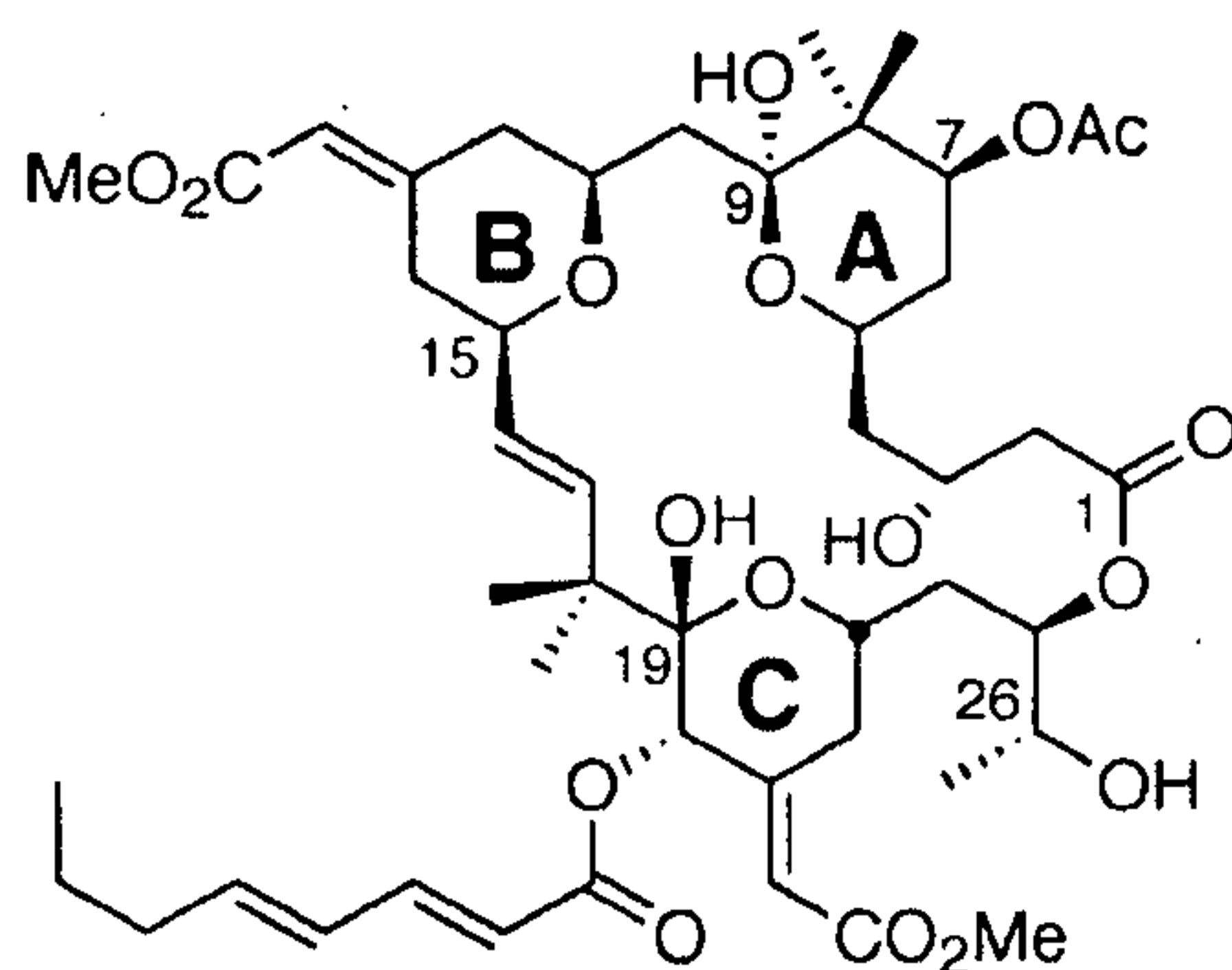
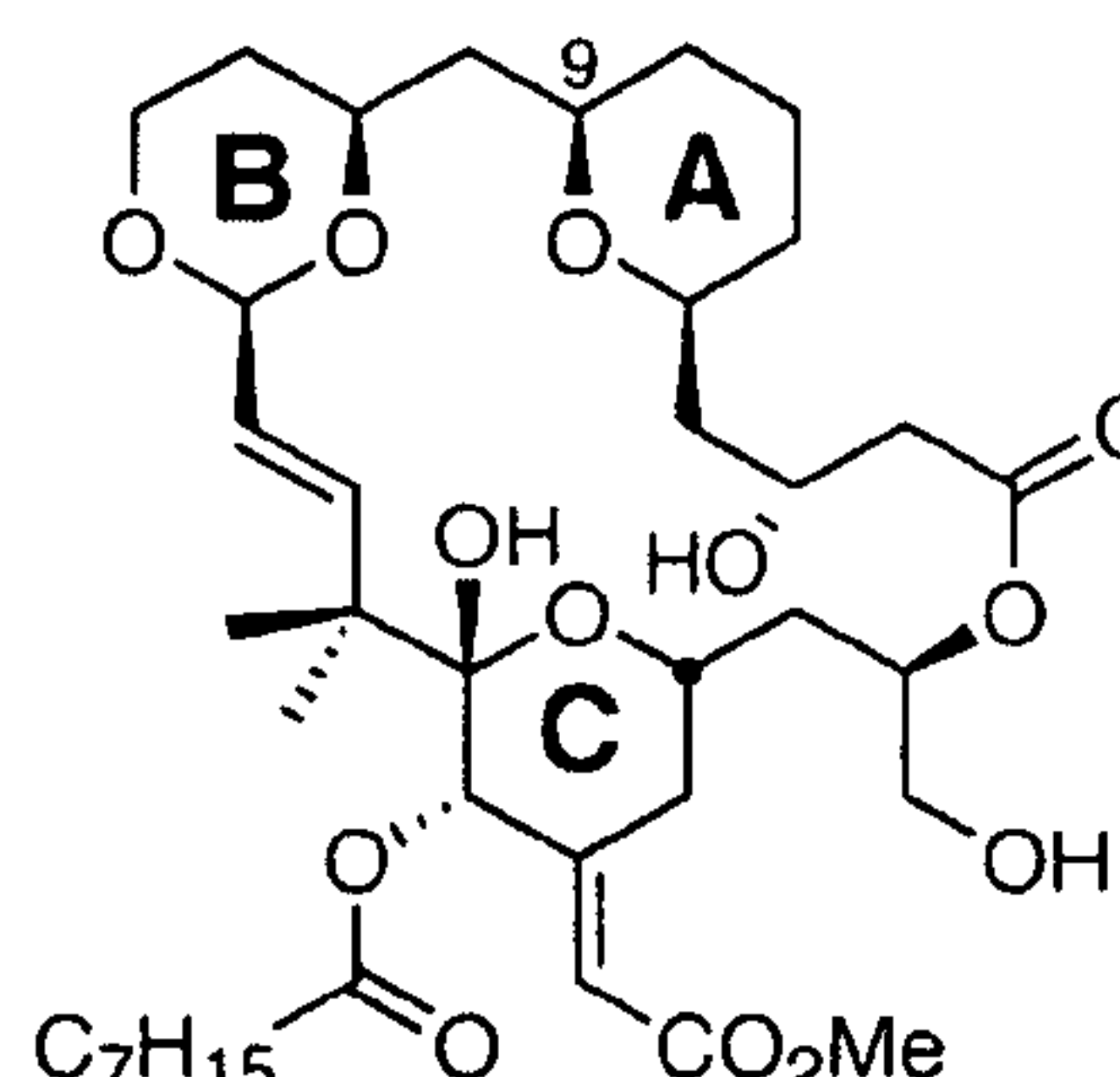
Name	PKC Affin (nM)	MW	Description
Bryostatin 1	1.35	988	2 pyran + 1 cyclic acetal + macrocycle
Analog 1	0.25	737	1 pyran + 2 cyclic acetal + macrocycle
Analog 2	6.50	723	1 pyran + 2 cyclic acetal + macrocycle
Analog 7a	-	642	1 pyran + 2 cyclic acetals + macrocycle

Analog 7b	297	711	1 pyran + 2 cyclic acetals + macrocycle
Analog 7c	3.4	726	1 pyran + 2 cyclic acetals + macrocycle
Analog 7d	10000	745	1 pyran + 2 cyclic acetals + macrocycle, acetylated
Analog 8	8.3	754	2 cyclic acetals + macrocycle
Analog 9	10000	599	2 cyclic acetals

[0036] Analog 1 (Wender et al. (2004) *Curr Drug Discov Technol.* 1: 1; Wender et al. (1998) *Proc Natl Acad Sci U S A* 95: 6624; Wender et al. (2002) *Am Chem Soc.* 124: 13648 (each incorporated herein by reference in their entireties)) possesses the highest affinity for PKC. This bryolog is about 100 times more potent than bryostatin-1. Only Analog 1 exhibits a higher affinity for PKC than bryostatin. Analog 2, which lacks the A ring of bryostatin-1 is the simplest analog that maintains high affinity for PKC. In addition to the active bryologs, Analog 7d, which is acetylated at position 26, has virtually no affinity for PKC.

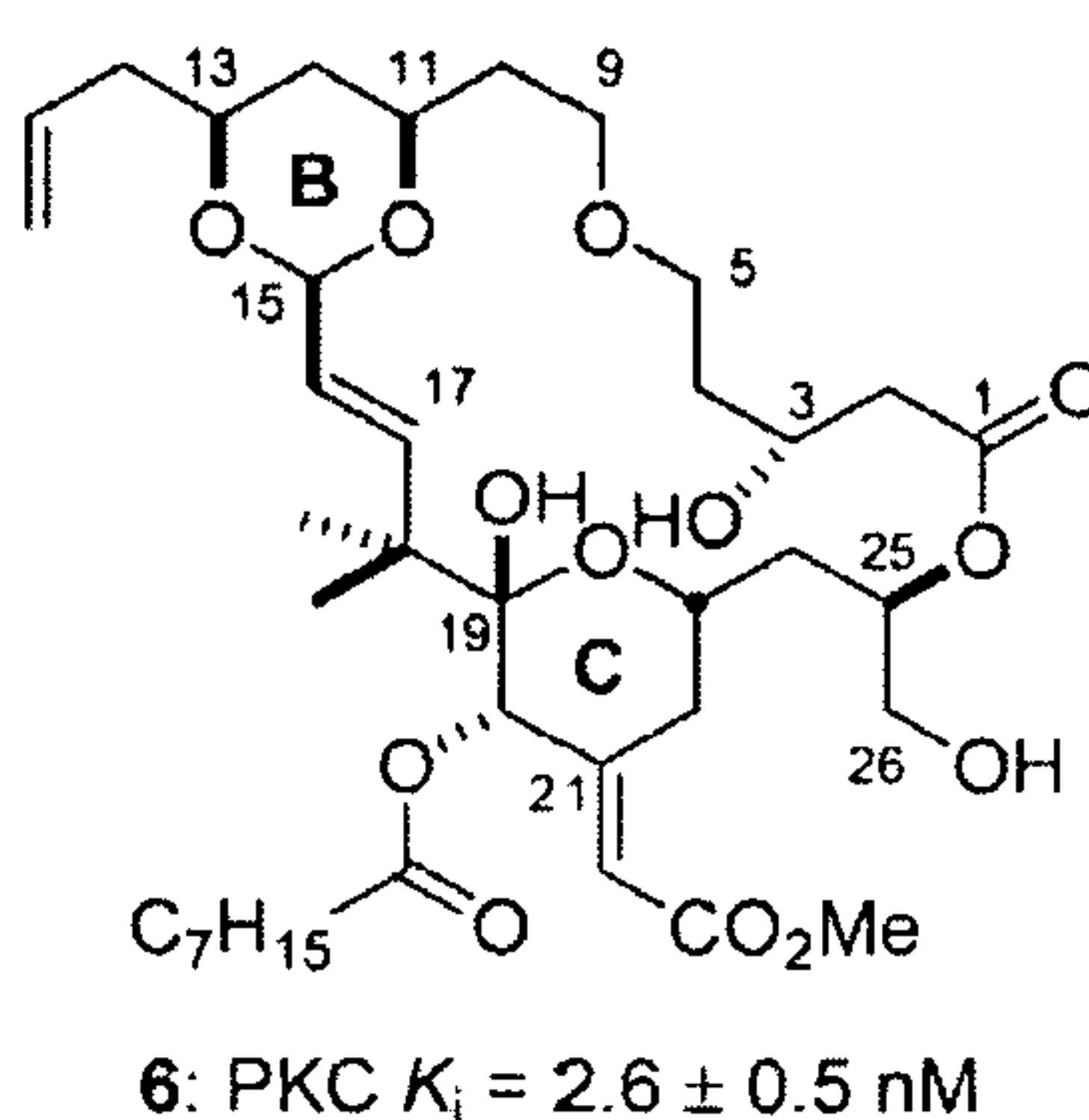
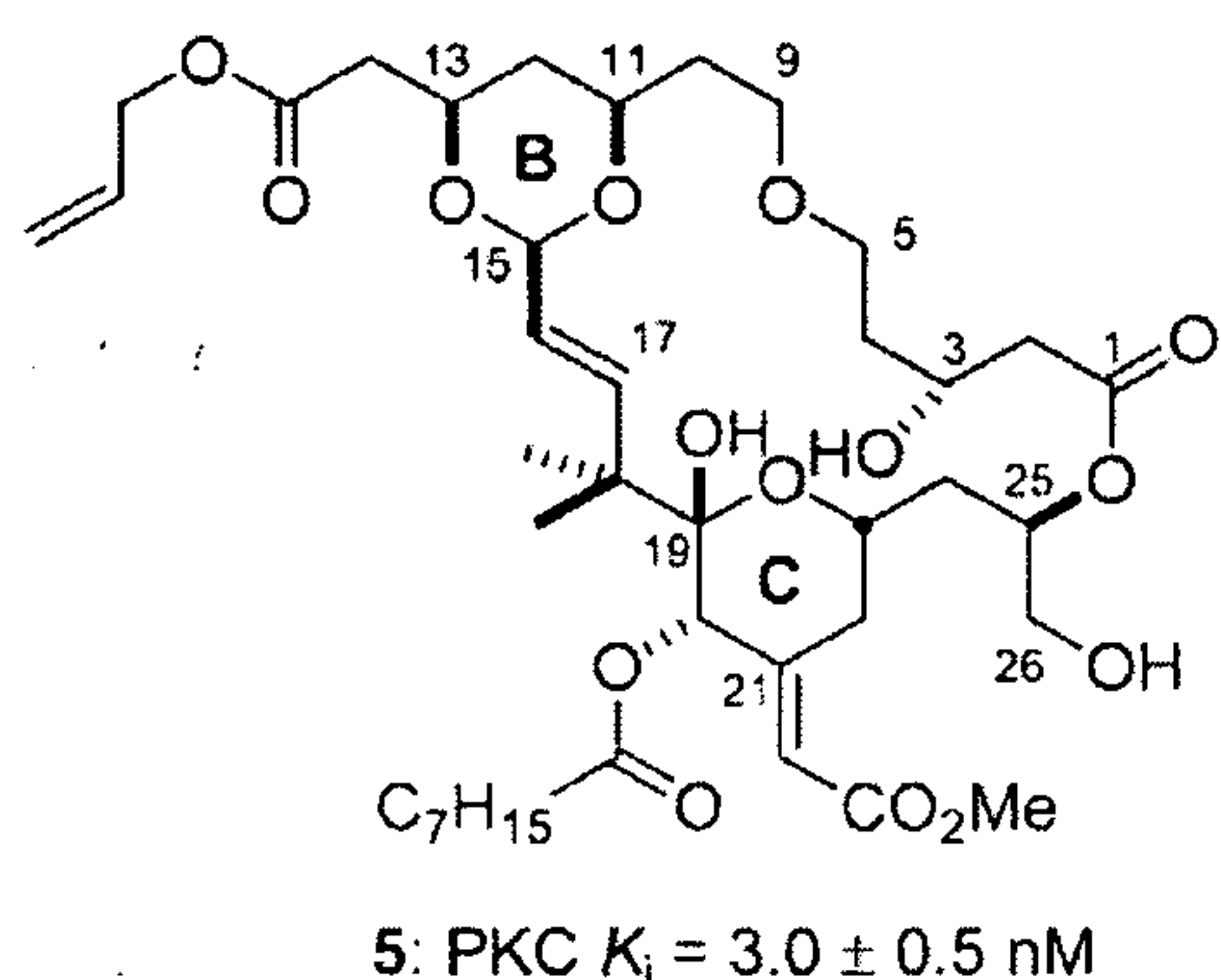
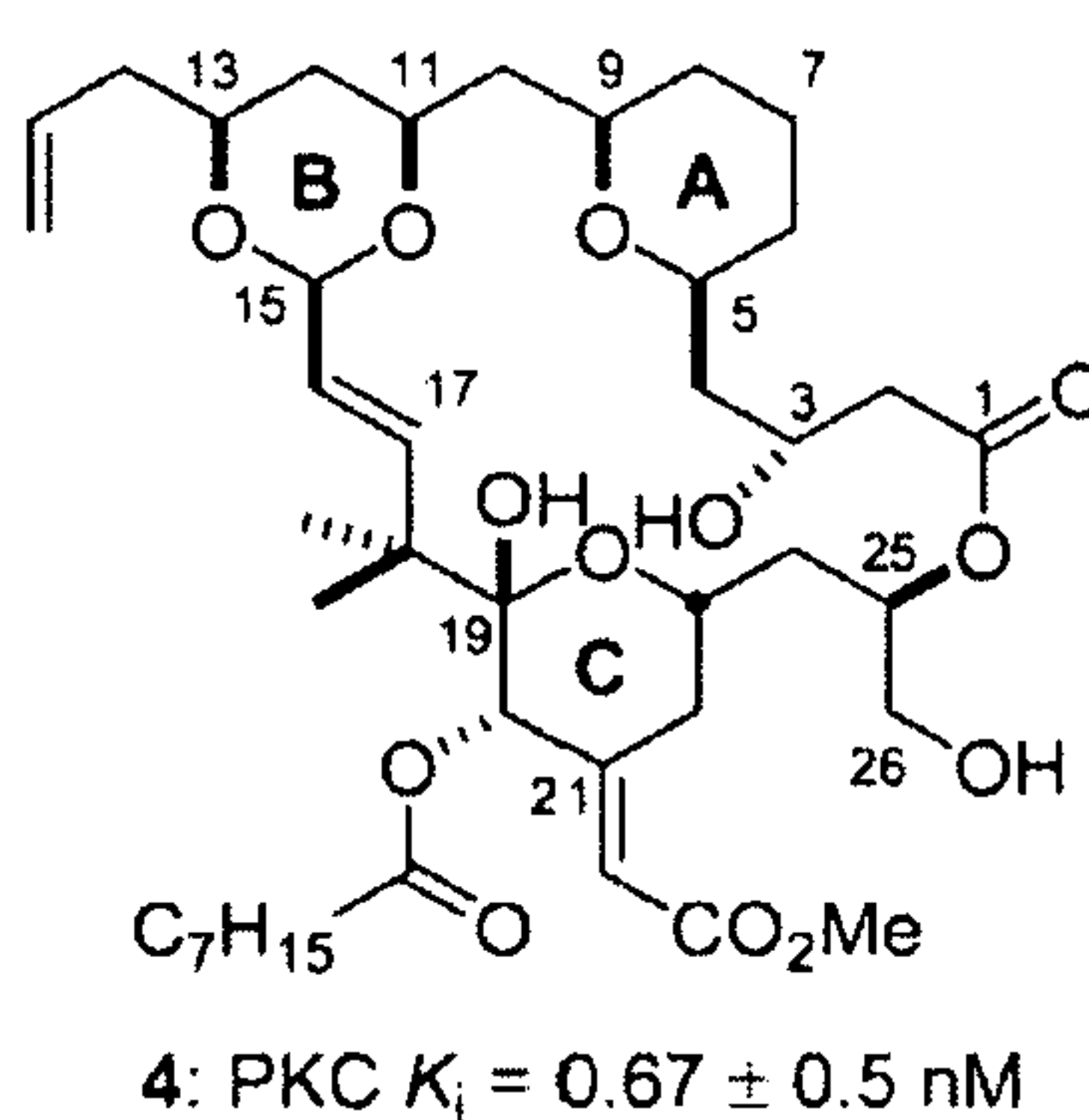
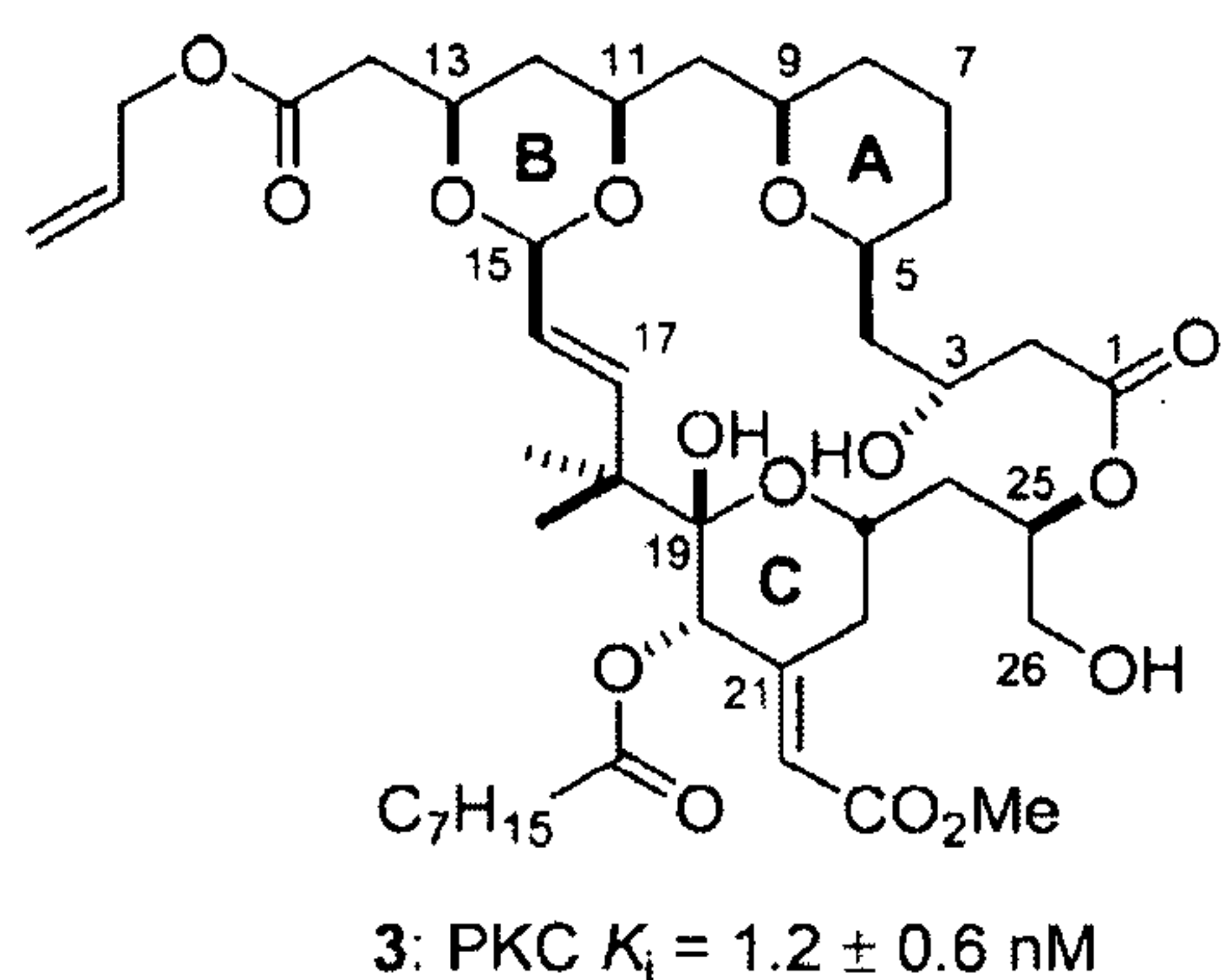
Analog 2; $K_i = 8.0$ nM

3 R = *t*-Bu
 4 R = Ph
 5 R = (CH₂)₃*p*-Br-Ph

Bryostatin 1; $K_i = 1.35$ nMAnalog 1; $K_i = 0.25$ nM

[0037] B-ring bryologs are also suitable for use in the methods of the present invention. These synthetic bryologs have affinities in the low nanomolar range (Wender et al. (2006) *Org*

Lett. 8: 5299 (incorporated herein by reference in its entirety)). The B-ring bryologs have the advantage of being completely synthetic, and do not require purification from a natural source.



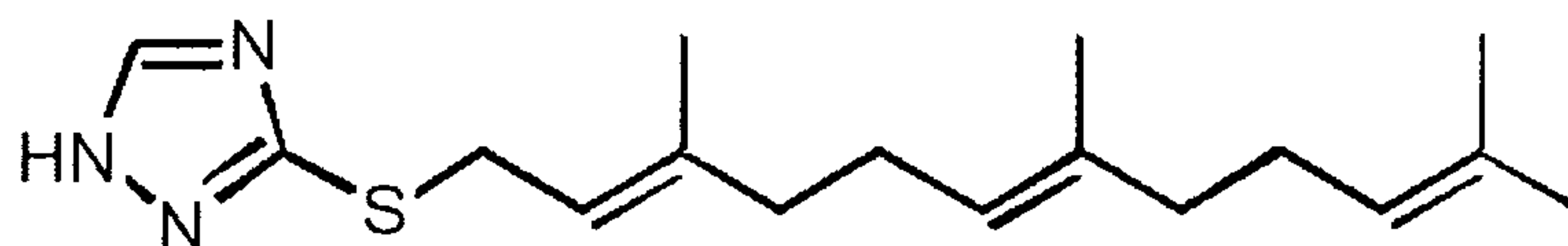
PKC binding affinities for B-ring bryologs

[0038] A third class of suitable bryostatin analogs is the A-ring bryologs. These bryologs have slightly lower affinity for PKC than bryostatin I (6.5, 2.3, and 1.9 nM for bryologs 3, 4, and 5, respectively) but have a lower molecular weight.

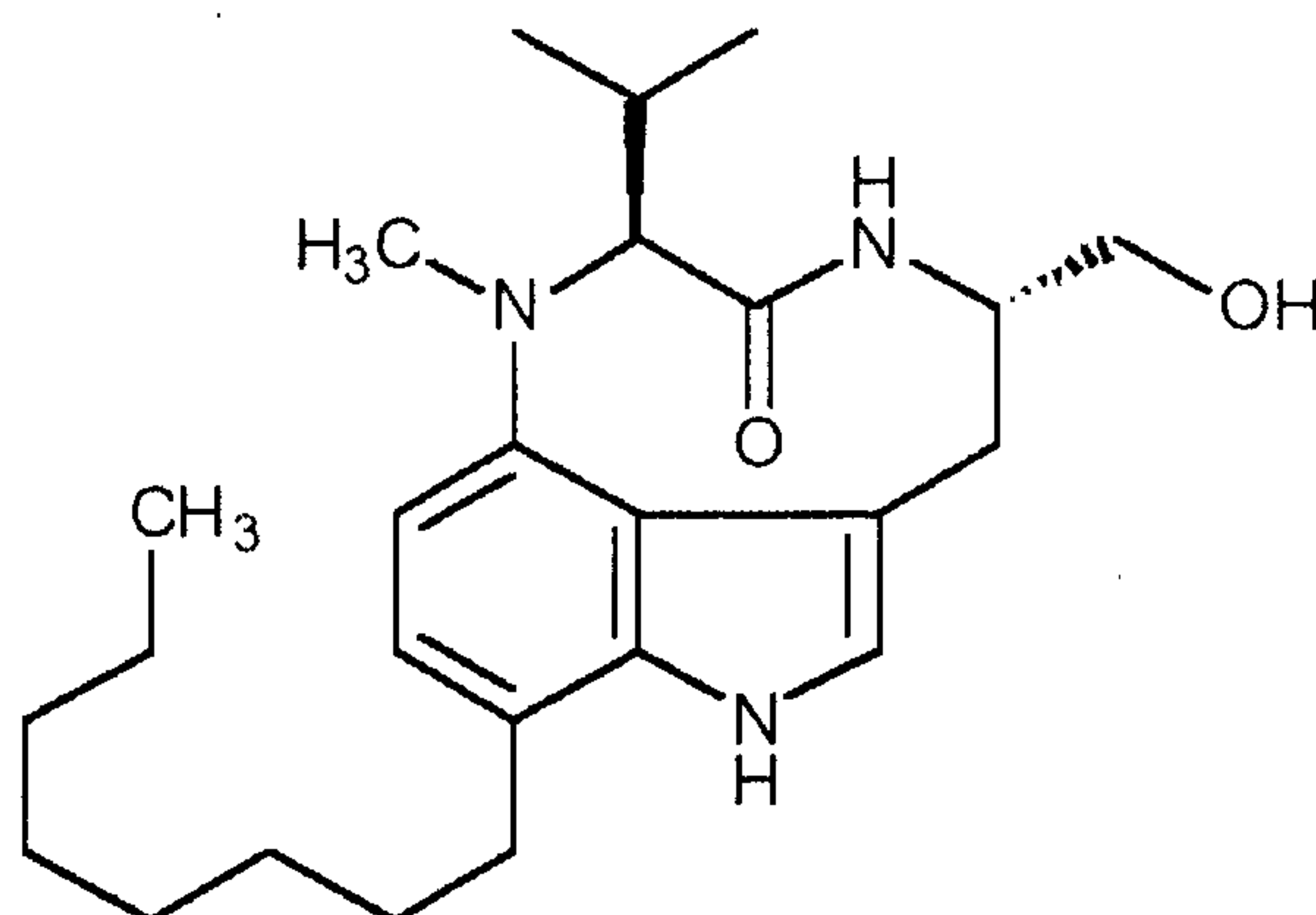
[0039] A number of derivatives of diacylglycerol (DAG) bind to and activate protein kinase C (Niedel et al. (1983) Proc. Natl. Acad. Sci. USA 80: 36; Mori et al. (1982) J. Biochem (Tokyo) 91: 427; Kaibuchi et al. (1983) J. Biol. Chem. 258: 6701). However, DAG and DAG derivatives are of limited value as drugs. Activation of PKC by diacylglycerols is transient, because they are rapidly metabolized by diacylglycerol kinase and lipase (Bishop et al. (1986) J. Biol. Chem. 261: 6993; Chung et al. (1993) Am. J. Physiol. 265: C927; incorporated herein by reference in their entirety). The fatty acid substitution determines the strength of activation. Diacylglycerols having an unsaturated fatty acid are most active. The stereoisomeric

configuration is also critical. Fatty acids with a 1,2-sn configuration are active, while 2,3-sn-diacylglycerols and 1,3-diacylglycerols do not bind to PKC. Cis-unsaturated fatty acids are synergistic with diacylglycerols. In one embodiment of the present invention, the term "PKC activator" expressly excludes DAG or DAG derivatives, such as phorbol esters.

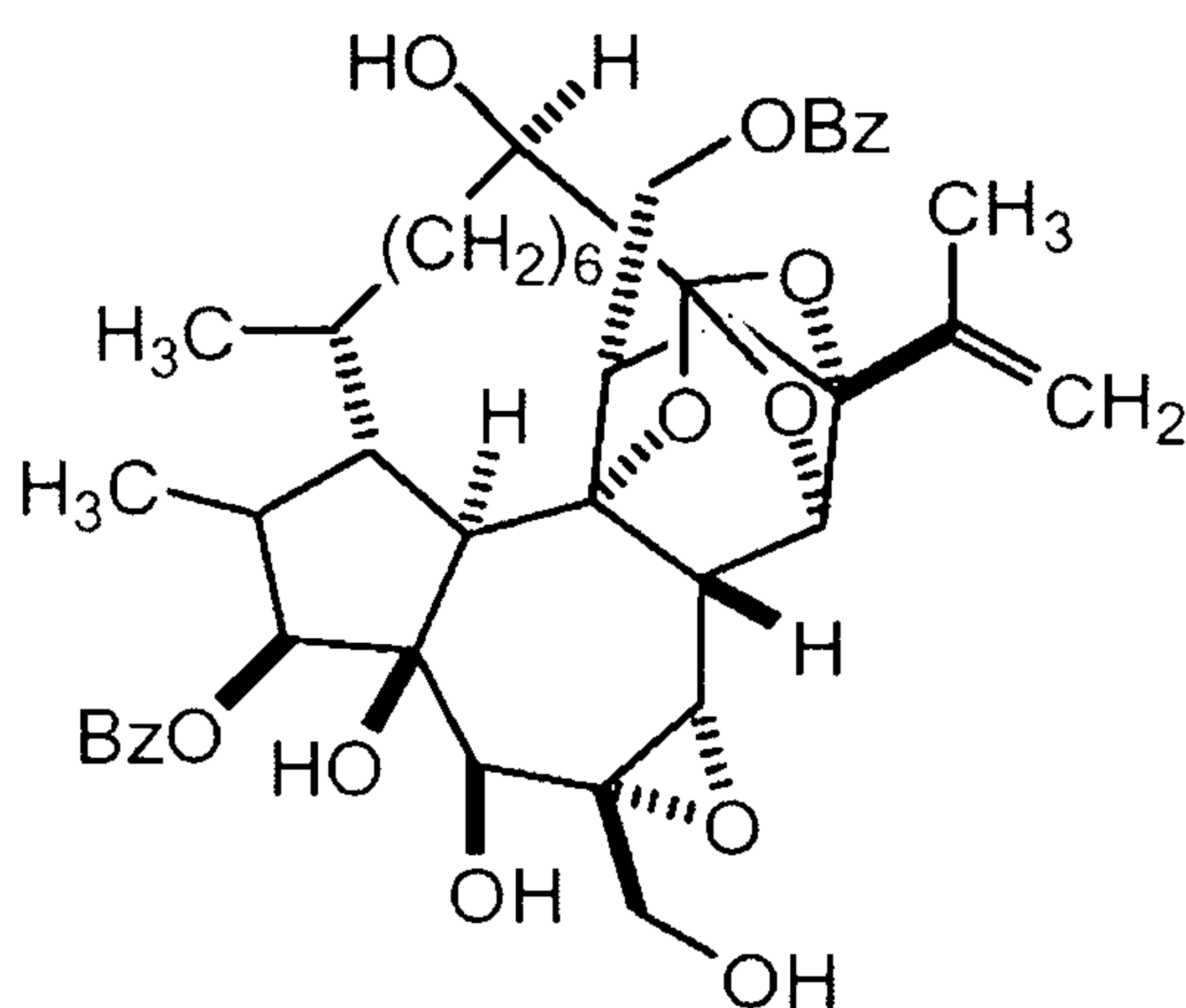
[0040] Isoprenoids are PKC activators suitable for use in the methods of the present invention. Farnesyl thiotriazole, for example, is a synthetic isoprenoid that activates PKC with a K_d of 2.5 μM . Farnesyl thiotriazole, for example, is equipotent with dioleoylglycerol (Gilbert et al. (1995) *Biochemistry* 34: 3916; incorporated herein by reference in its entirety), but does not possess hydrolyzable esters of fatty acids. Farnesyl thiotriazole and related compounds represent a stable, persistent PKC activator. Because of its low MW (305.5) and absence of charged groups, farnesyl thiotriazole would readily cross the blood-brain barrier.



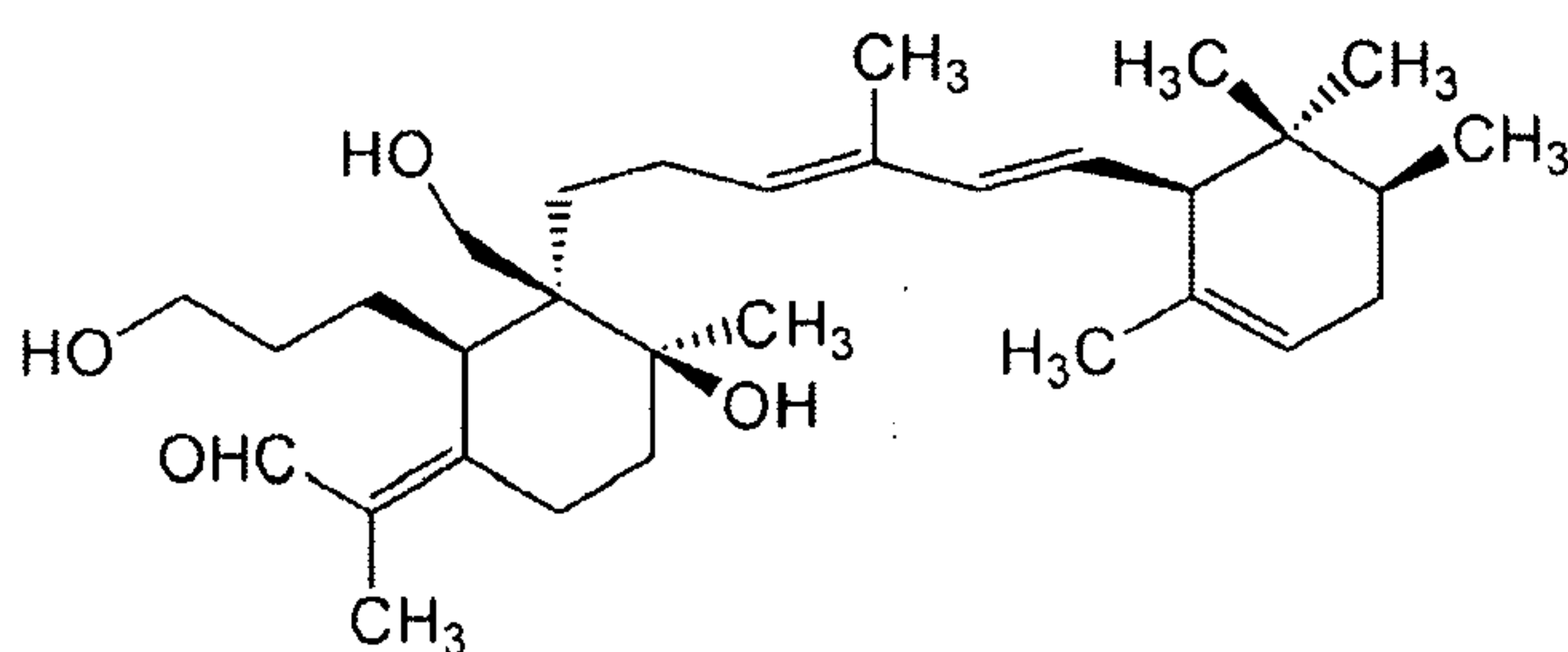
[0041] Octylindolactam V is a non-phorbol protein kinase C activator related to teleocidin. The advantages of octylindolactam V, specifically the (-)-enantiomer, include greater metabolic stability, high potency (Fujiki et al. (1987) *Adv. Cancer Res.* 49: 223; Collins et al. (1982) *Biochem. Biophys. Res. Commun.* 104: 1159; each incorporated herein by reference in its entirety) ($\text{EC}_{50} = 29\text{nM}$) and low molecular weight that facilitates transport across the blood brain barrier.



[0042] Gnidimacrin is a daphnane-type diterpene that displays potent antitumor activity at concentrations of 0.1 - 1 nM against murine leukemias and solid tumors. It acts as a PKC activator at a concentration of ≈ 3 nM in K562 cells, and regulates cell cycle progression at the G1/S phase through the suppression of Cdc25A and subsequent inhibition of cyclin dependent kinase 2 (Cdk2) (100% inhibition achieved at 5 ng/ml). Gnidimacrin is a heterocyclic natural product similar to bryostatin, but somewhat smaller (MW = 774.9).

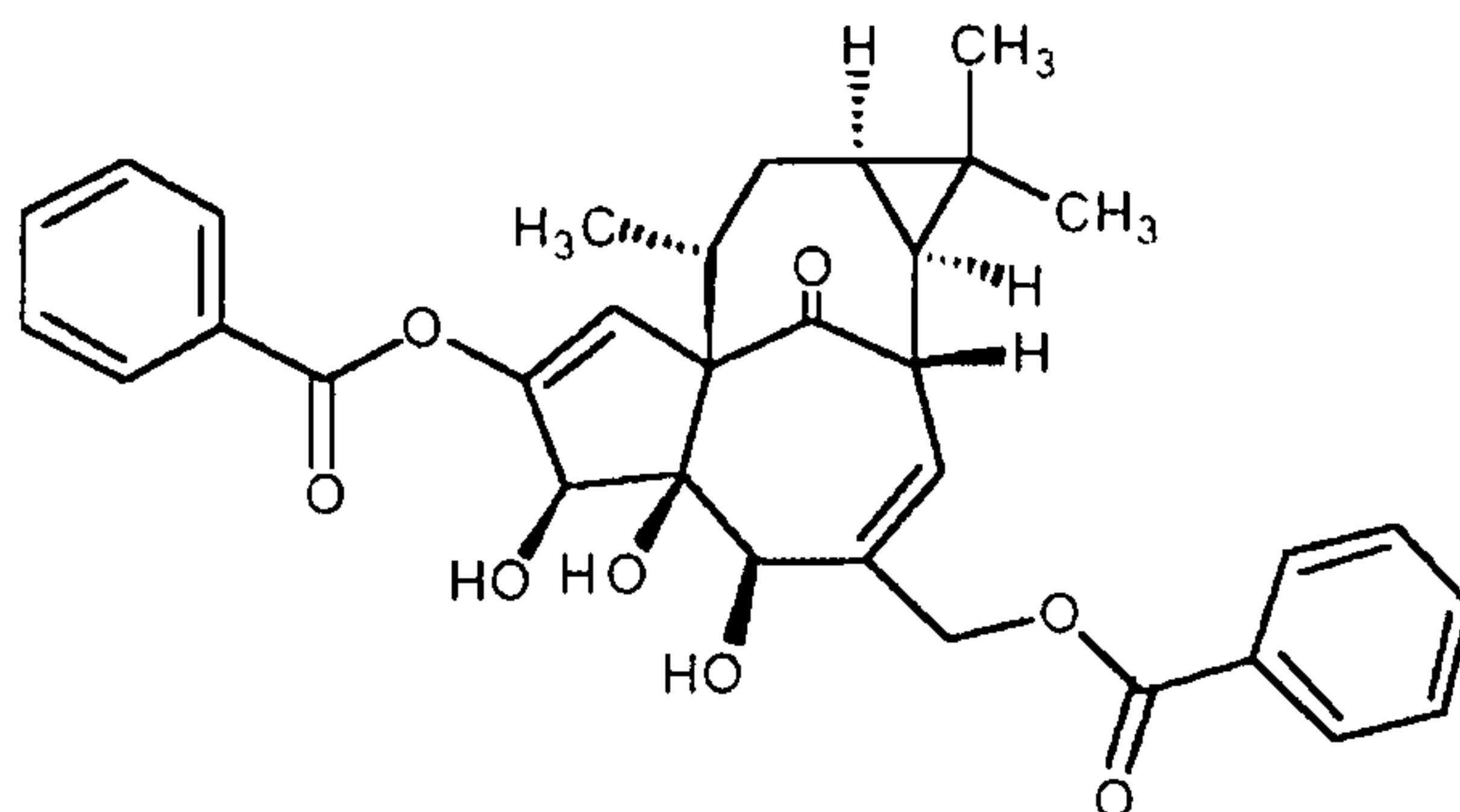


[0043] Iripallidal is a bicyclic triterpenoid isolated from *Iris pallida*. Iripallidal displays anti-proliferative activity in a NCI 60 cell line screen with GI50 (concentration required to inhibit growth by 50%) values from micromolar to nanomolar range. It binds to PKC α with high affinity (K_i = 75.6 nM). It induces phosphorylation of ERK1/2 in a RasGRP3-dependent manner. M.W. 486.7. Iripallidal is only about half the size of bryostatin and lacks charged groups.

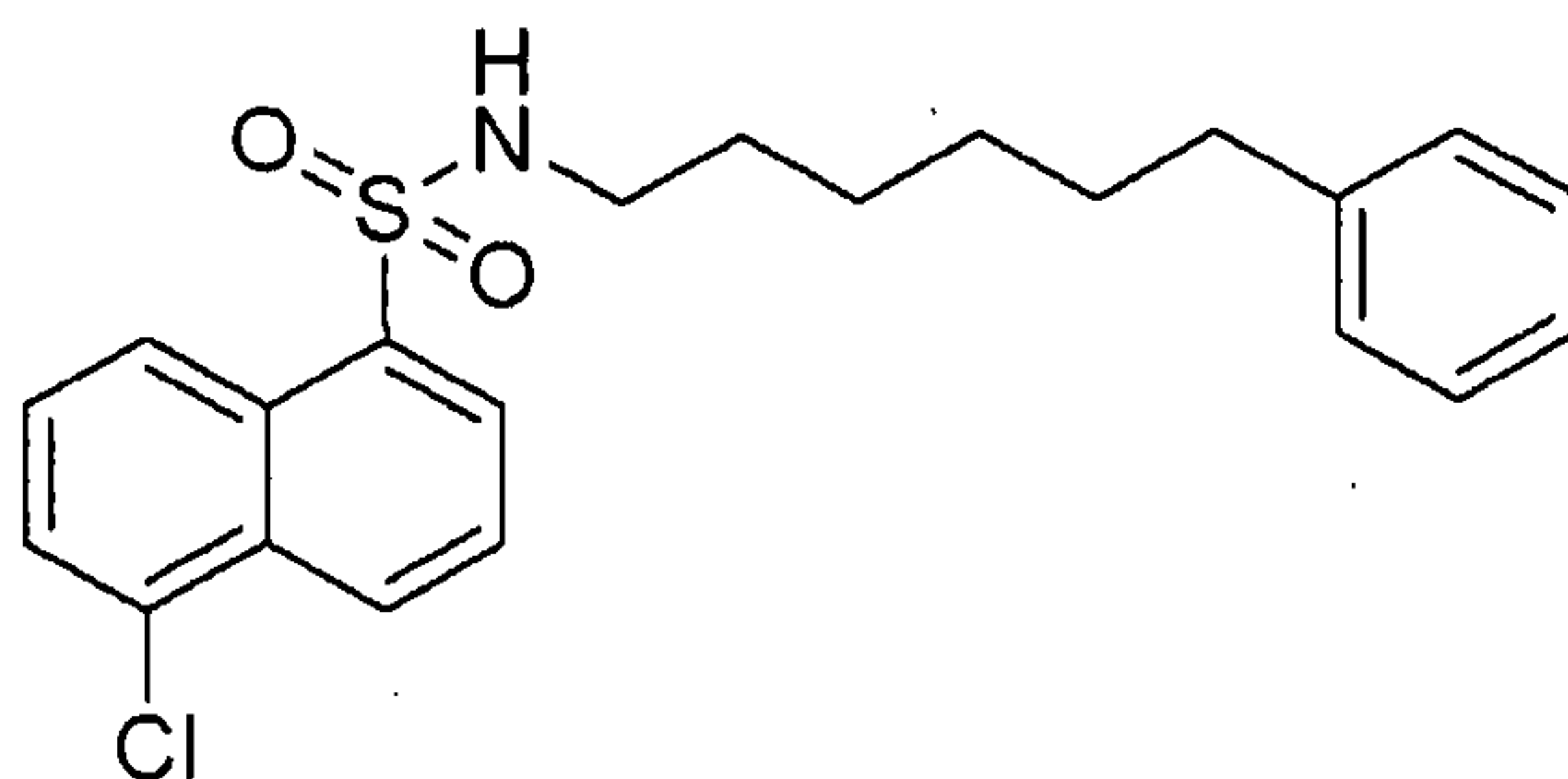


[0044] Ingenol [43] is a diterpenoid related to phorbol but possesses much less toxicity. It is derived from the milkweed plant *Euphorbia peplus*. Ingenol 3,20-dibenzoate, for example, competes with [3H]phorbol dibutyrate for binding to PKC (K_i for binding=240 nM) (Winkler et al. (1995) J.Org.Chem. 60: 1381; incorporated herein by reference). Ingenol-3-angelate

possesses antitumor activity against squamous cell carcinoma and melanoma when used topically (Ogbourne et al. (2007) *Anticancer Drugs*. 18: 357; incorporated herein by reference).



[0045] Naphthalenesulfonamides, including N-(n-heptyl)-5-chloro-1-naphthalenesulfonamide (SC-10) and N-(6-Phenylhexyl)-5-chloro-1-naphthalenesulfonamide, are members of another class of PKC activators. SC-10 activates PKC in a calcium-dependent manner, using a mechanism similar to that of phosphatidylserine (Ito et al. (1986) *Biochemistry* 25: 4179; incorporated herein by reference). Naphthalenesulfonamides act by a different mechanism from bryostatin and would be expected to show a synergistic effect with bryostatin or a member of another class of PKC activators. Structurally, naphthalenesulfonamides are similar to the calmodulin (CaM) antagonist W-7, but are reported to have no effect on CaM kinase.



[0046] The linoleic acid derivative DCP-LA (2-[(2-pentylcyclopropyl)methyl]cyclopropanecarboxylic acid) is one of the few known isoform-specific activators of PKC known. DCP-LA selectively activates PKC ϵ with a maximal effect at 100 nM. (Kanno *et al.* (2006) *J. Lipid Res.* 47: 1146). Like SC-10, DCP-LA interacts with the phosphatidylserine binding site of PKC, instead of the diacylglycerol binding site.

[0047] An alternative approach to activating PKC directly is to increase the levels of the endogenous activator, diacylglycerol. Diacylglycerol kinase inhibitors such as 6-(2-(4-[(4-fluorophenyl)phenylmethylene]-1-piperidinyl)ethyl)-7-methyl-5H-thiazolo[3,2-a]pyrimidin-5-one (R59022) and [3-[2-[4-(bis-(4-fluorophenyl)methylene]piperidin-1-yl)ethyl]-2,3-dihydro-2-thioxo-4(1H)-quinazolinone (R59949) enhance the levels of the endogenous ligand diacylglycerol, thereby producing activation of PKC (Meinhardt *et al.* (2002) *Anti-Cancer Drugs* 13: 725).

[0048] Growth factor activators, such as the 4-methyl catechol derivatives, such as 4-methylcatechol acetic acid (MCBA), that stimulate the synthesis and/or activation of growth factors such as NGF and BDNF, also activate PKC as well as convergent pathways responsible for synaptogenesis and/or neuritic branching.

[0049] All books, articles, patents or other publications and references are hereby incorporated by reference in their entireties. Reference to any compound herein includes the racemate as well as the single enantiomers

EXAMPLES

[0050] The following Examples serve to further illustrate the present invention and are not to be construed as limiting its scope in any way.

EXAMPLE 1: ANIMAL MODEL OF HEAD TRAUMA

[0051] Minimal traumatic brain injury (TBI) was produced in mice by a concussive event using a 30g mass. One hour post trauma, the mice received 20 or 30 μ g bryostatin per kg doses by intraperitoneal injection. The injections were repeated twice weekly for a total of 5 treatments. The effects on learning and memory of bryostatin treatment in treated animals was tested in the Morris Water Maze.

EXAMPLE 2: MORRIS WATER MAZE

[0052] In the Morris Water Maze, animals were allowed to swim in a tank divided into four quadrants, only one of which had a safety platform beneath the water. The platform was removed and the animals were tested for how long they searched the correct quadrant versus the incorrect quadrants. In the passive avoidance procedure the animal remembers the distinctive environment in which a mild electric shock is delivered and avoids it on a second occasion.

EXAMPLE 3: BRYOSTATIN (30 UG/KG) AND TREATMENT OF MINIMAL TBI

[0053] Minimal traumatic brain injury (TBI) was produced in mice by a concussive event using a 30g mass. One hour post trauma, the mice received 30 μ g bryostatin per kg doses by intraperitoneal injection. The injections were repeated twice weekly for a total of 5 treatments. The escape latencies in a Morris Water Maze of mice treated with bryostatin after minimal TBI were compared to animals with minimal TBI, control animals receiving no TBI or bryostatin, and animals receiving bryostatin only. The results are shown in **Figure 1**.

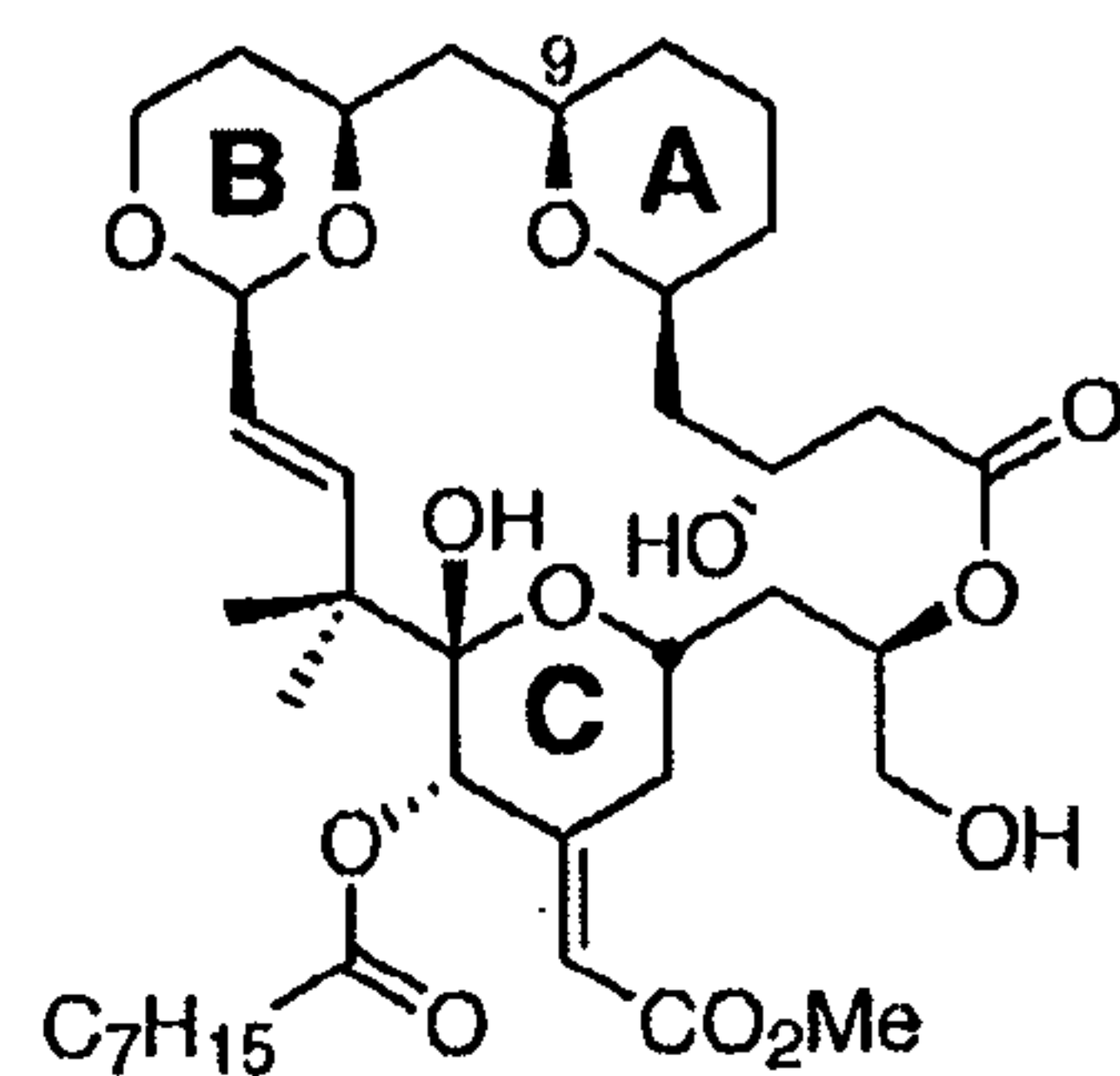
EXAMPLE 4: BRYOSTATIN (20 UG/KG) AND TREATMENT OF MINIMAL TBI

[0054] Minimal traumatic brain injury (TBI) was produced in mice by a concussive event using a 30g mass. One hour post trauma, the mice received 20 μ g bryostatin per kg doses by intraperitoneal injection. The injections were repeated twice weekly for a total of 5 treatments. The escape latencies in a Morris Water Maze of mice treated with bryostatin after minimal TBI were compared to animals with minimal TBI, control animals receiving no TBI or bryostatin, and animals receiving bryostatin only. The results are shown in **Figure 2**. The memory retention of each treatment or control group is tabulated in **Figure 3**.

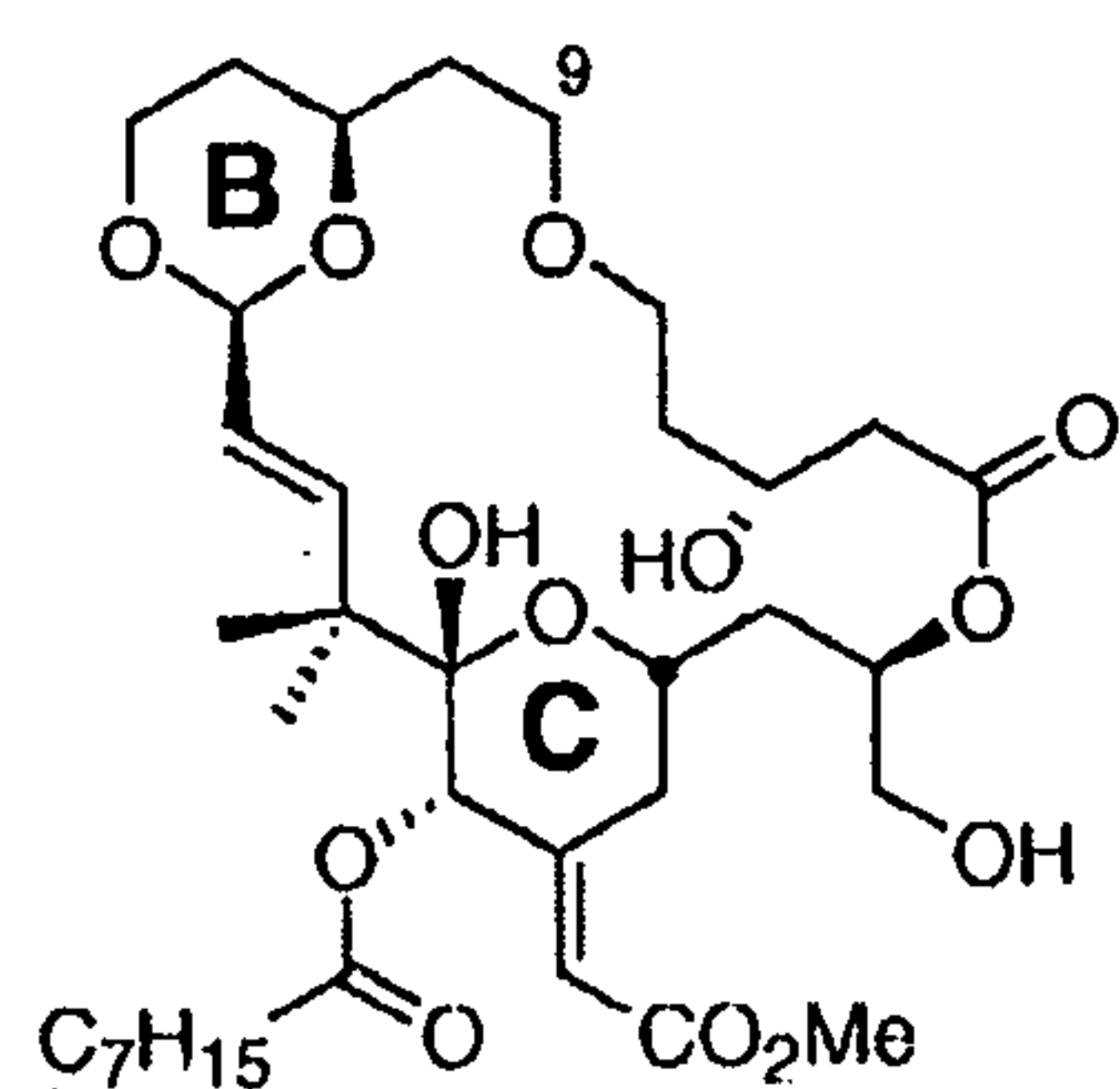
AMENDED CLAIMS**[received by the International Bureau on 08 May 2009 (08.05.09)]**

1. A method of treating head trauma comprising the steps of identifying a subject having suffered a head trauma and administering to said subject an amount of a pharmaceutical composition comprising a protein kinase C (PKC) activator or 4-methylcatechol acetic acid (MCBA), or other derivatives of methylcatechol, and a pharmaceutically acceptable carrier effective to treat at least one symptom of head trauma, wherein the PKC activator is selected from a group consisting of: bryologs, diacylglycerol derivatives other than phorbol esters, isoprenoids, daphnane-type diterpenes, bicyclic triterpenoids, naphthalenesulfonamides, lineolic acid derivatives, diacylglycerol kinase inhibitors, growth factor activators, or a combination thereof.
2. (Cancelled).
3. (Cancelled).
4. (Cancelled).
5. (Cancelled).
6. (Cancelled).
7. The method of claim 1, wherein the pharmaceutical composition comprises 4-methylcatechol acetic acid.
8. The method of claim 1, wherein said administration is initiated within 1 day of said head trauma.
9. The method of claim 1, wherein said administration is initiated within 2 days of said head trauma.
10. The method of claim 1, wherein said administration is initiated within 3 days of said head trauma.

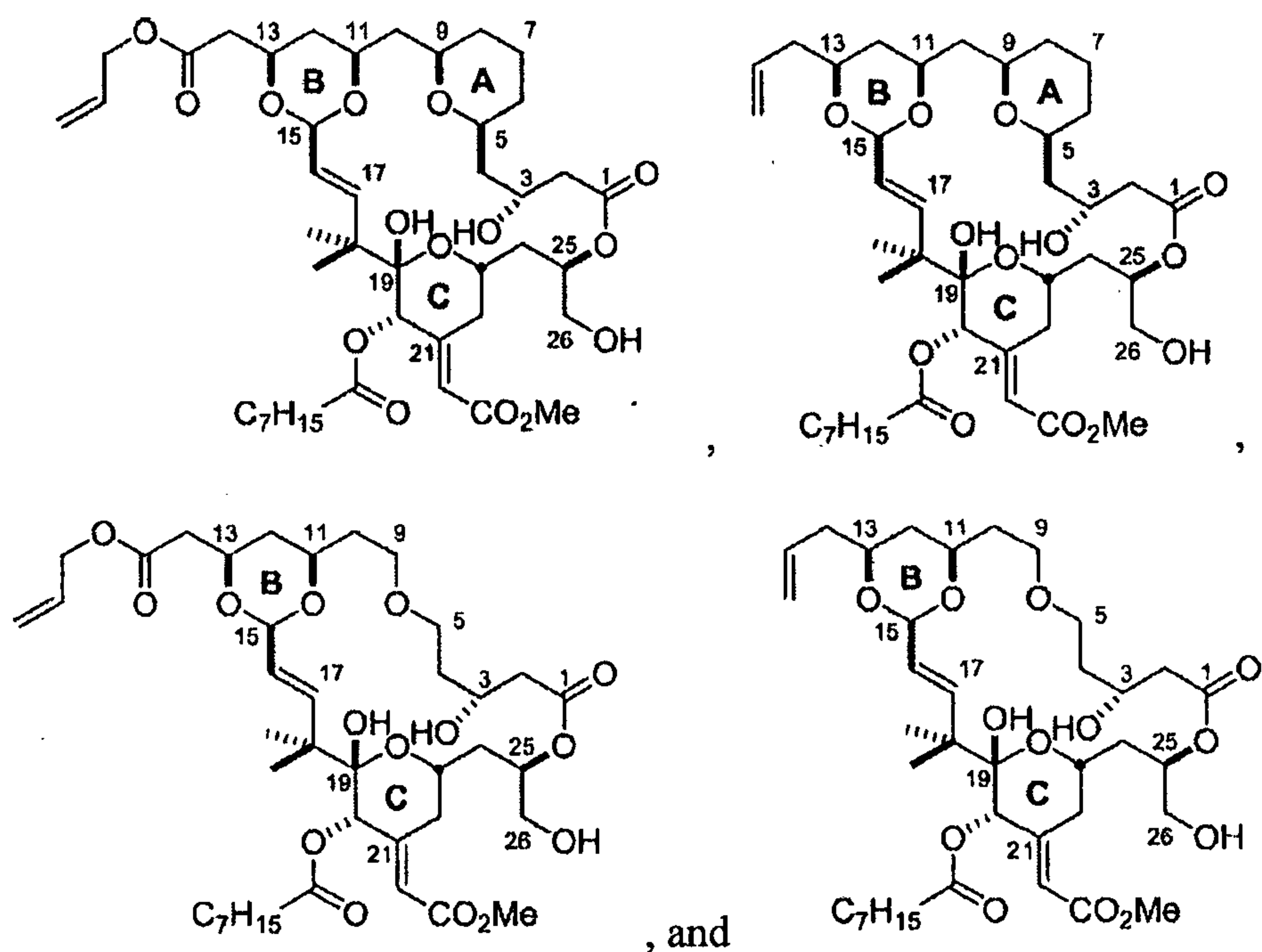
11. The method of claim 1, wherein said administration is initiated between 1 and 2 days of said head trauma.
12. The method of claim 1, wherein said administration is initiated between 1 and 3 days of said head trauma.
13. The method of claim 1, wherein the treatment is continued for a duration of 1 week.
14. The method of claim 1, wherein the treatment is continued for a duration of 2 weeks.
15. The method of claim 1, wherein the treatment is continued for a duration of 3 weeks.
16. The method of claim 1, wherein the treatment is continued for a duration of 4 weeks.
17. The method of claim 1, wherein the treatment is continued for a duration of 6 week.
18. The method of claim 1, wherein said treatment reverses head trauma-induced brain injury.
19. The method of claim 1, wherein said treatment reverses head trauma-induced memory impairment.
20. The method of claim 1, wherein the bryolog is a B-ring bryolog or A-ring bryolog.
21. The method of claim 20, wherein the B-ring or A-ring bryolog has a molecular weight from about 600 to 755 and an affinity for PKC from about 0.25 nM to 10 μ M.
22. The method of claim 1, wherein the bryolog is



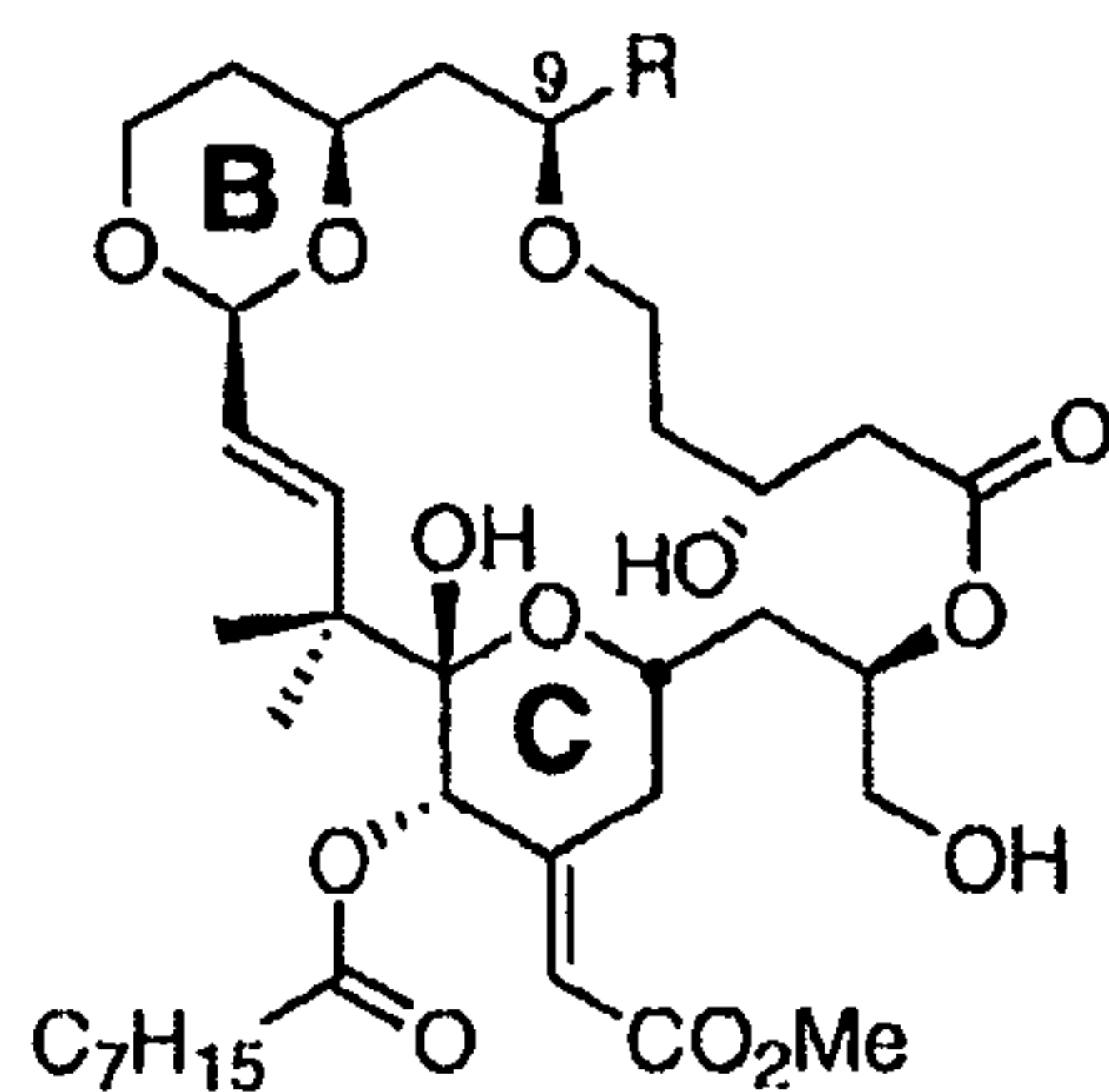
23. The method of claim 1, wherein the bryolog is



24. The method of claim 20, wherein the B-ring byrolog is selected from the group consisting of



25. The method of claim 20, wherein the A-Ring analog is



and R is t-Bu, Ph, or $(\text{CH}_2)_3p\text{-Br-Ph}$.

26. The method of claim 1, wherein the diacylglycerol derivative is comprised of unsaturated fatty acids.

27. The method of claim 26, wherein the fatty acids are in a 1,2-sn configuration.

28. The method of claim 26, wherein the fatty acids are cis-unsaturated fatty acids.

29. The method of claim 1, wherein the PKC activator is octylindolactam V.
30. The method of claim 29, wherein the octylindolactam is the (-)-enantiomer.
31. The method of claim 1, wherein the daphnane-type diterpene is gnidimacrin.
32. The method of claim 1, wherein the bicyclic triterpenoid is iripallidal.
33. The method of claim 1, wherein the diterpenoid is ingenol.
34. The method of claim 1, wherein the diterpenoid is ingenol 3,20-dibenzoate.
35. The method of claim 1, wherein the diterpenoid is ingenol-3-angelate.
36. The method of claim 1, wherein the naphthalenesulfonamide is N-(n-heptyl)-5-chloro-1-naphthalenesulfonamide or N-(6-Phenylhexyl)-5-chloro-1-naphthalenesulfonamide.
37. The method of claim 1, wherein the lineolic acid derivative is 2-[(2-pentylcyclopropyl)methyl]-cyclopropaneoctanoic acid.
38. The method of claim 1, wherein the diacylglycerol kinase inhibitor is 6-(2-(4-[(4-fluorophenyl)phenylmethylene]-1-piperidinyl)ethyl)-7-methyl-5H-thiazolo[3,2-a]pyrimidin-5-one.
39. The method of claim 1, wherein the diacylglycerol kinase inhibitor is [3-[2-[4-(bis-(4-fluorophenyl)methylene]piperidin-1-yl)ethyl]-2,3-dihydro-2-thioxo-4(1H)-quinazolinone.
40. The method of claim 1, wherein the growth factor activator is a 4-methyl catechol derivative.
41. The method of claim 1, wherein the growth factor activator is 4-methylcatechol acetic acid.

Figure 1

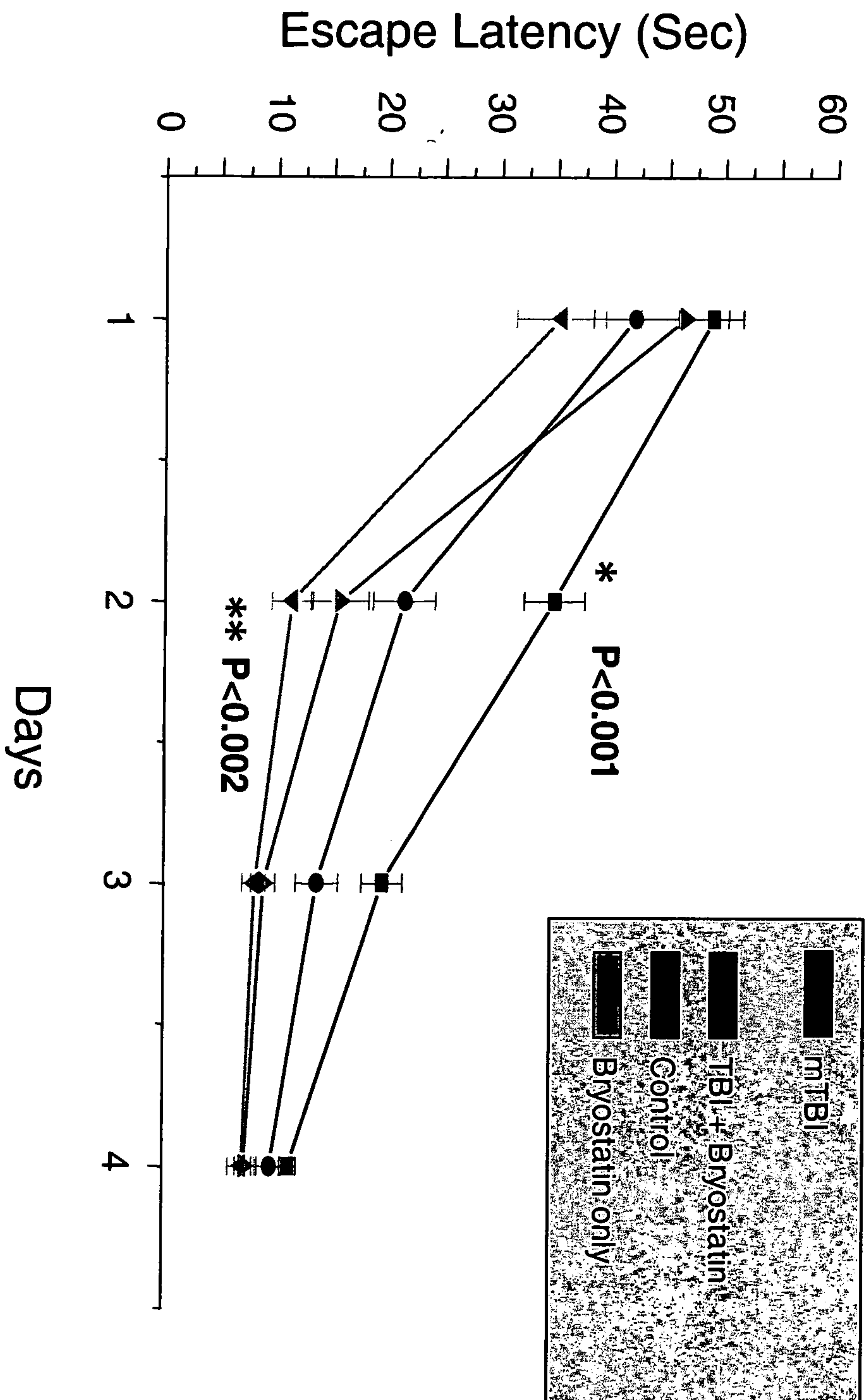


Figure 2

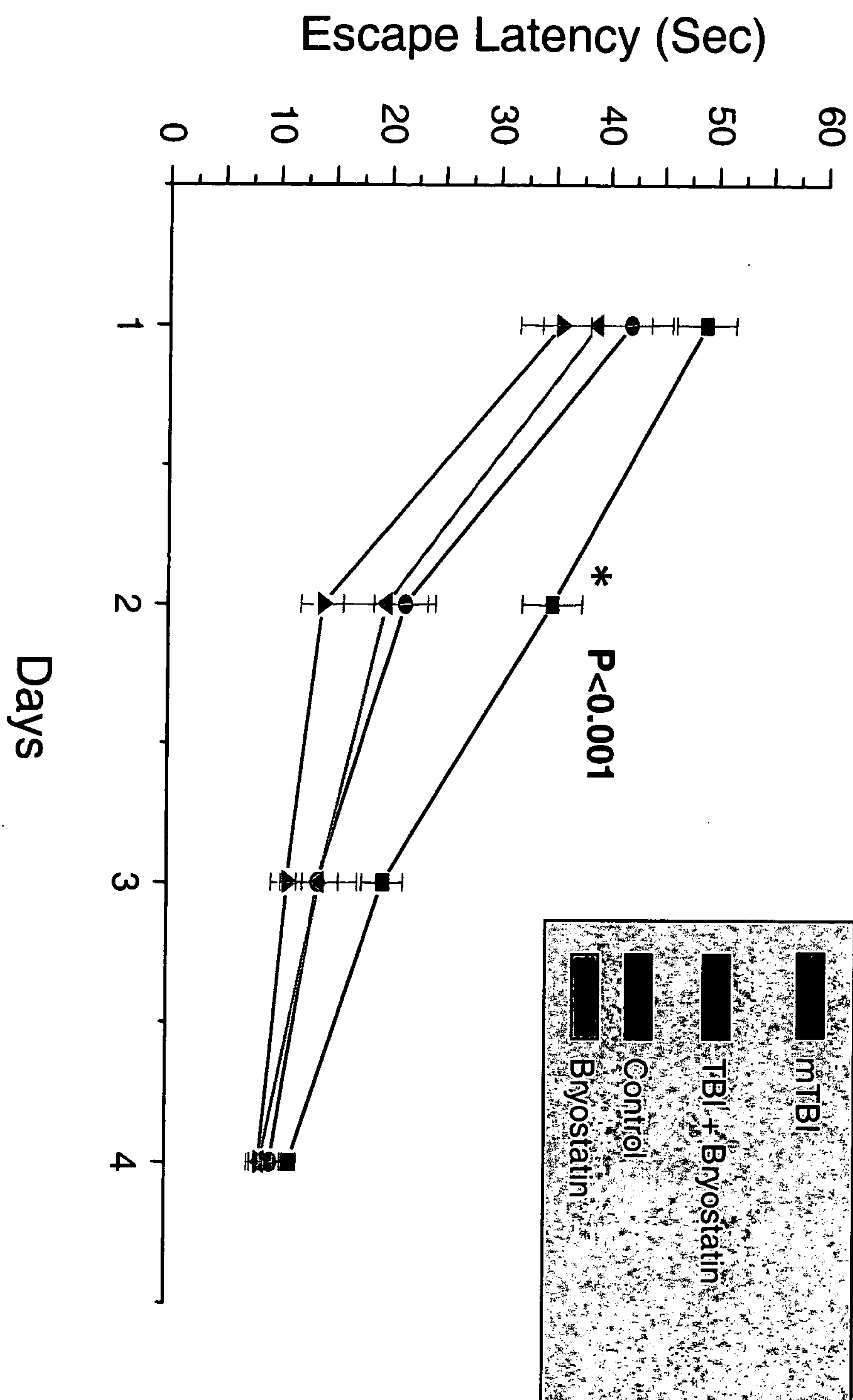


Figure 3

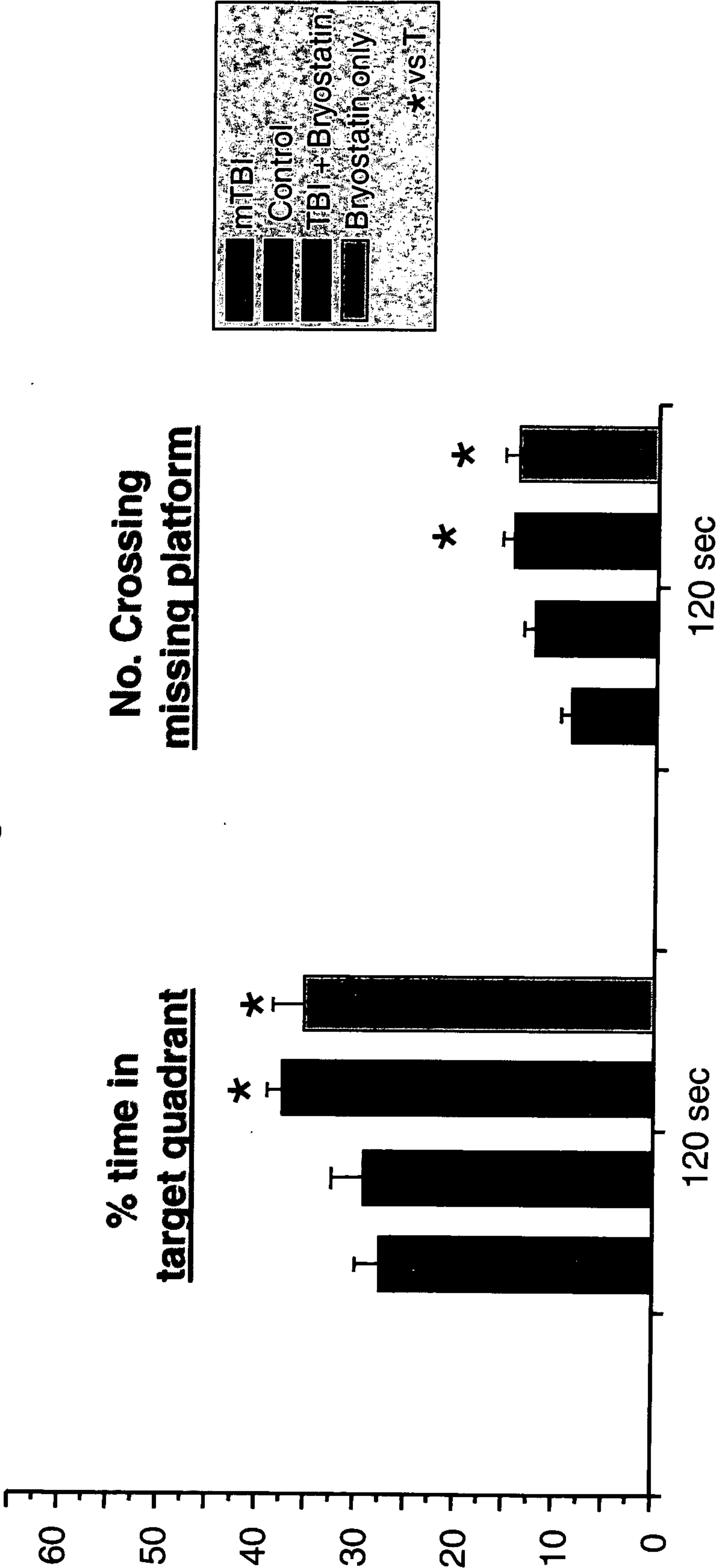


Figure 3

