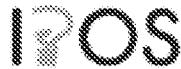


(19)



INTELLECTUAL PROPERTY
OFFICE OF SINGAPORE

(11) Publication number:

SG 182236 A1

30.07.2012

(43) Publication date:

(51) Int. Cl:

;

(12)

Patent Application

(21) Application number: 2012047940

(71) Applicant:

SUCAMPO AG GRABEN 5, ZUG,
CH6300, SWITZERLAND CH

(22) Date of filing: 03.07.2008

(72) Inventor:

UENO, RYUJI 11025 STANMORE DRIVE,
POTOMAC, MONTGOMERY, MARYLAND
20854, UNITED STATES OF AMERICA
US

(30) Priority: US 60/929,557 03.07.2007

US 61/037,517 18.03.2008

(54) **Title:**

PHARMACEUTICAL COMBINATION OF NSAID AND
PROSTAGLANDIN COMPOUND

(57) **Abstract:**

44 ABSTRACT PHARMACEUTICAL COMBINATION OF NSAID AND PROSTAGLANDIN COMPOUND Provided is a pharmaceutical combination of a NSAID and a prostaglandin compound of formula (I). The combination of the invention is useful for the indications for NSAID use such as pain, Alzheimer disease and cancer. Figure for publication: None

ABSTRACT**PHARMACEUTICAL COMBINATION OF NSAID AND PROSTAGLANDIN
COMPOUND**

Provided is a pharmaceutical combination of a NSAID and a prostaglandin compound of formula (I). The combination of the invention is useful for the indications for NSAID use such as pain, Alzheimer disease and cancer.

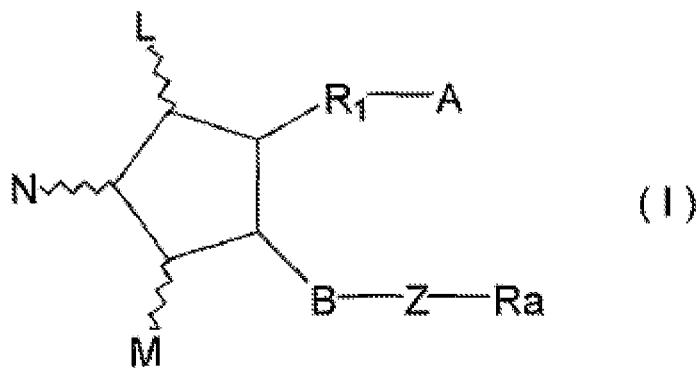


Figure for publication: None

DESCRIPTION

PHARMACEUTICAL COMBINATION OF NSAID
AND PROSTAGLANDIN COMPOUND

5

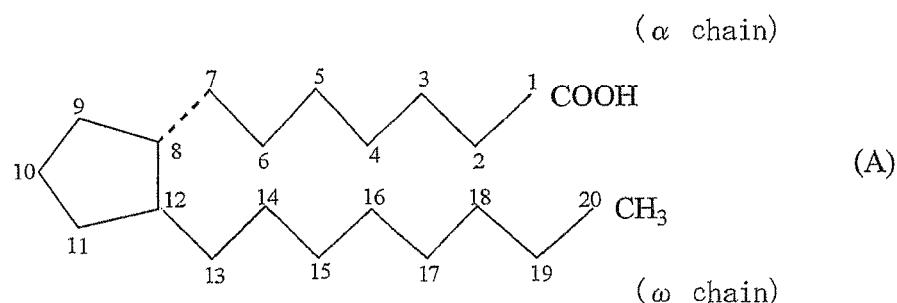
TECHNICAL FIELD

[0001] The present invention relates to a pharmaceutical combination comprising a NSAID and a specific prostaglandin compound.

10 [0002] Non steroidal anti-inflammatory drugs (NSAIDs) are the most frequently prescribed drugs worldwide for the treatment of pain from various etiologies. Commonly used NSAIDs include sulindac, naproxen, indomethacin, mefenamic acid, diclofenac, fenoprofen, and diflunisal. Recently 15 NSAIDs have been considered for treatment and prevention of Alzheimer's disease (Trends Pharmacol Sci. 28(10), 536-543, 2007). In addition, NSAIDs could reduce the risk of cancer through the inhibition of cyclooxygenase-2 (COX-2), and are shown a protective effect in colon, rectum, esophagus, 20 stomach, pancreas, breast, prostate, lung and bladder cancer, suggesting a common mechanistic effect of NSAIDs in all these different cancers (BMC Cancer 2003, 3:28 and Annals of Internal Medicine 146(5), 376-389, 2007). Despite the therapeutic benefits of NSAIDs, their use is 25 commonly limited by an increased risk of

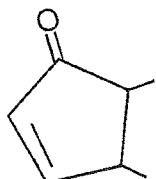
gastrointestinal (GI) side effects such as gastritis and gastric ulcer.

[0003] Prostaglandins (hereinafter, referred to as PGs) are members of class of organic carboxylic acids, which are 5 contained in tissues or organs of human or other mammals, and exhibit a wide range of physiological activity. PGs found in nature (primary PGs) generally have a prostanoic acid skeleton as shown in the formula (A) :

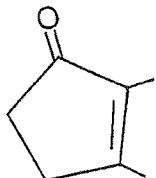


[0004] PGs are classified into several types according to the structure and substituents on the five-membered ring, for example,

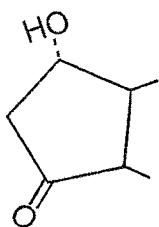
Prostaglandins of the A series (PGAs) ;



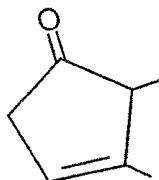
Prostaglandins of the B series (PGBs) ;



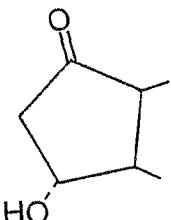
Prostaglandins of the C series (PGCs) ;



Prostaglandins of the D series (PGDs);

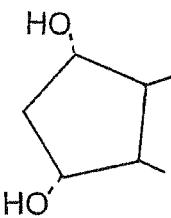


Prostaglandins of the E series (PGEs);



5

Prostaglandins of the F series (PGFs);



and the like. Further, they are classified into PG₁s containing a 13,14-double bond; PG₂s containing, 5,6- and 10 13,14-double bonds; and PG₃s containing 5,6-, 13,14- and 17,18-double bonds. PGs are known to have various pharmacological and physiological activities, for example, vasodilatation, inducing of inflammation, platelet aggregation, stimulating uterine muscle, stimulating

intestinal muscular activity, anti-ulcer effects and the like.

[0005] U.S. Patent Nos. 5,225,439, 5,166,174, 5,284,858, 5,428,062, 5,380,709, 5,876,034 and 6,265,440 describe that 5 certain prostaglandin E compounds are effective for the treatment of ulcers such as duodenal ulcer and gastric ulcer.

[0006] U.S. Patent No. 5,317,032 to Ueno et al. describes prostaglandin analog cathartics, including the 10 existence of bicyclic tautomers and U.S. Patent No. 6,414,016 to Ueno describes the bicyclic tautomers as having pronounced activity as anti-constipation agents. The bicyclic tautomers, substituted by one or more halogen atoms can be employed in small doses for relieving 15 constipation. At the C-16 position, especially, fluorine atoms, can be employed in small doses for relieving constipation.

[0007] U.S. Patent No.7064148 to Ueno et al. describes 20 prostaglandin compound opens and activates chloride channels, especially ClC channels, more especially ClC-2 channel.

[0008] U.S Patent publication No.2003/0166632 to Ueno described ClC-2 channel opener is effective for the treatment of a disease or a condition responsive to opening 25 of ClC-2 channel.

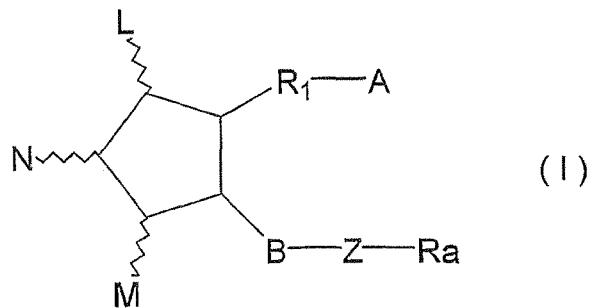
[0009] U.S. Patent publication No.2003/0119898 to Ueno et al. describes specific composition of a halogenated prostaglandin analog for the treatment and prevention of constipation.

5 [0010] U.S. Patent publication No.2004/0138308 to Ueno et al. describes chloride channel opener, especially a prostaglandin compound for the treatment of abdominal discomfort, and the treatment of functional gastrointestinal disorders such as irritable bowel syndrome
10 and functional dyspepsia.

DISCLOSURE OF THE INVENTION

[0011] The present invention relates to a pharmaceutical combination comprising:

- (a) a NSAID and
- 15 (b) a prostaglandin (PG) compound represented by the formula (I):



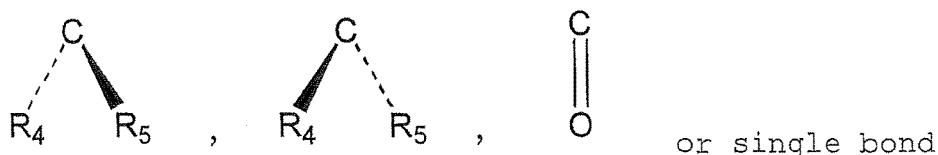
wherein L, M and N are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo,
20 wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have at least one

double bond;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

B is single bond, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{CH}_2-\text{CH}_2-$
5 CH_2- , $-\text{CH}=\text{CH}-\text{CH}_2-$, $-\text{CH}_2-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-\text{CH}_2-$ or $-\text{CH}_2-\text{C}\equiv\text{C}-$;

Z is



wherein R₄ and R₅ are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein
10 R₄ and R₅ are not hydroxy and lower alkoxy at the same time;

R₁ is a saturated or unsaturated bivalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

R_a is a saturated or unsaturated lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy;

cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group, provided that Ra is substituted by halogen or Z is C=O.

[0012] By combining a NSAID and a PG compound of formula 5 (I), the effect of the NSAID is augmented and/or the adverse side effect of the NSAID such as ulcer formation is well suppressed.

[0013] Especially, the present invention also relates to a pharmaceutical combination comprising:

10 (a) a naproxen, and
(b) a prostaglandin (PG) compound represented by the formula (I).

[0014] In another aspect of the invention, there is provided a pharmaceutical composition comprising:

15 (a) a pharmaceutically effective amount of a NSAID such as naproxen, and
(b) a pharmaceutically effective amount of a prostaglandin (PG) compound represented by the formula (I) in association with a pharmaceutically suitable excipient.

20 According to the present invention, the composition may be formulated as separate dosage forms each comprising individual active ingredient or as a single dosage form comprising the two active ingredients.

[0015] Further, the present invention provides a method 25 for treating a condition or disease which is one of the

indications for NSAID use, which comprises administering a combination of:

(a) a pharmaceutically effective amount of a NSAID such as naproxen, and

5 (b) a pharmaceutically effective amount of a prostaglandin (PG) compound represented by the formula (I) to a patient in need thereof.

According to the present invention, the (a) and (b) compounds may be administered simultaneously, 10 separately or sequentially.

DETAILED DESCRIPTION OF THE INVENTION

[0016]

(a) NSAID

The NSAID is selected from but not limited to the group consisting of salicylates, indomethacin, flurbiprofen, diclofenac, ketorolac, naproxen, piroxicam, tebufelone, ibuprofen, etodolac, nabumetone, tenidap, alcofenac, antipyrine, aminopyrine, dipyrone, aminopyrone, phenylbutazone, clofezone, oxyphenbutazone, prexazone, 20 apazone, benzydamine, bucolome, cinchopen, clonixin, ditrazol, epirizole, fenoprofen, floctafeninl, flufenamic acid, glaphenine, indoprofen, ketoprofen, meclofenamic acid, mefenamic acid, niflumic acid, phenacetin, salidifamides, sulindac, suprofen, tolmetin, pharmaceutically acceptable 25 salts thereof, and mixtures thereof. The preferable

example of NSAID is naproxen.

[0017]

(b) The compound of formula (I)

The nomenclature of the prostaglandin compounds used
5 herein is based on the numbering system of the prostanoic
acid represented in the above formula (A).

[0018] The formula (A) shows a basic skeleton of the C-
20 carbon atoms, but the present invention is not limited
to those having the same number of carbon atoms. In the
10 formula (A), the numbering of the carbon atoms which
constitute the basic skeleton of the PG compounds starts at
the carboxylic acid (numbered 1), and carbon atoms in the
α-chain are numbered 2 to 7 towards the five-membered ring,
those in the ring are 8 to 12, and those in the ω-chain are
15 13 to 20. When the number of carbon atoms is decreased in
the α-chain, the number is deleted in the order starting
from position 2; and when the number of carbon atoms is
increased in the α-chain, compounds are named as
20 substitution compounds having respective substituents at
position 2 in place of the carboxy group (C-1). Similarly,
when the number of carbon atoms is decreased in the ω-chain,
the number is deleted in the order starting from position
20; and when the number of carbon atoms is increased in the
ω-chain, the carbon atoms beyond position 20 are named as
25 substituents. Stereochemistry of the compounds is the same

as that of the above formula (A) unless otherwise specified.

[0019] In general, each of the terms PGD, PGE and PGF represents a PG compound having hydroxy groups at positions 9 and/or 11, but in the present specification, these terms 5 also include those having substituents other than the hydroxy group at positions 9 and/or 11. Such compounds are referred to as 9-dehydroxy- 9-substituted-PG compounds or 11-dehydroxy-11-substituted-PG compounds. A PG compound having hydrogen in place of the hydroxy group is simply 10 named as 9- or 11-deoxy-PG compound.

[0020] As stated above, the nomenclature of the PG compounds is based on the prostanoic acid skeleton. However, in case the compound has a similar partial structure as a prostaglandin, the abbreviation of "PG" may 15 be used. Thus, a PG compound of which α -chain is extended by two carbon atoms, that is, having 9 carbon atoms in the α -chain is named as 2-decarboxy-2-(2-carboxyethyl)-PG compound. Similarly, a PG compound having 11 carbon atoms in the α -chain is named as 2-decarboxy-2-(4-carboxybutyl)- 20 PG compound. Further, a PG compound of which ω -chain is extended by two carbon atoms, that is, having 10 carbon atoms in the ω -chain is named as 20-ethyl-PG compound. These compounds, however, may also be named according to the IUPAC nomenclatures.

25 [0021] Examples of the analogs (including substituted

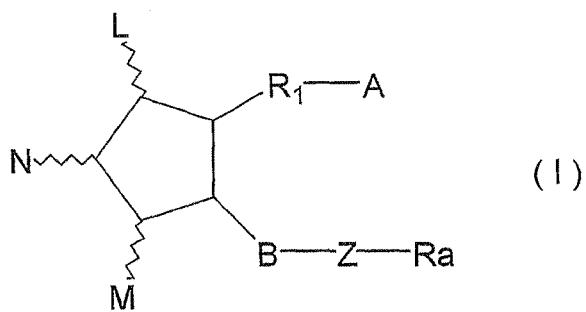
derivatives) or derivatives include a PG compound of which carboxy group at the end of α -chain is esterified; a compound of which α -chain is extended; physiologically acceptable salt thereof; a compound having a double bond at 5 2-3 position or a triple bond at position 5-6, a compound having substituent(s) at position 3, 5, 6, 16, 17, 18, 19 and/or 20; and a compound having lower alkyl or a hydroxy (lower) alkyl group at position 9 and/or 11 in place of the hydroxy group.

10 [0022] According to the present invention, preferred substituents at position 3, 17, 18 and/or 19 include alkyl having 1-4 carbon atoms, especially methyl and ethyl. Preferred substituents at position 16 include lower alkyl such as methyl and ethyl, hydroxy, halogen atoms such as 15 chlorine and fluorine, and aryloxy such as trifluoromethylphenoxy. Preferred substituents at position 17 include lower alkyl such as methyl and ethyl, hydroxy, halogen atoms such as chlorine and fluorine, aryloxy such as trifluoromethylphenoxy. Preferred substituents at 20 position 20 include saturated or unsaturated lower alkyl such as C1-4 alkyl, lower alkoxy such as C1-4 alkoxy, and lower alkoxy alkyl such as C1-4 alkoxy-C1-4 alkyl. Preferred substituents at position 5 include halogen atoms such as chlorine and fluorine. Preferred substituents at 25 position 6 include an oxo group forming a carbonyl group.

Stereochemistry of PGs having hydroxy, lower alkyl or hydroxy(lower)alkyl substituent at position 9 and/or 11 may be α , β or a mixture thereof.

[0023] Further, the above analogs or derivatives may be 5 compounds having an alkoxy, cycloalkyl, cycloalkyloxy, phenoxy or phenyl group at the end of the ω -chain where the chain is shorter than the primary PGs.

[0024] A specific prostaglandin compound used in the present invention is represented by the formula (I):



10

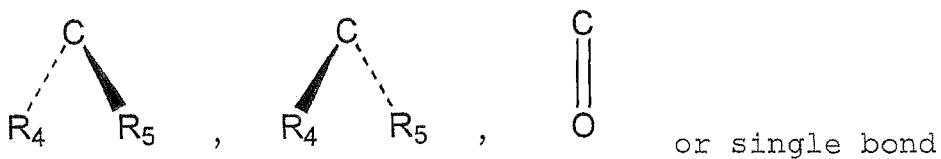
wherein L, M and N are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have at least one 15 double bond;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

B is single bond, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-\text{CH}_2-$, $-\text{CH}_2-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-\text{CH}_2-$ or $-\text{CH}_2-\text{C}\equiv\text{C}-$;

20

Z is



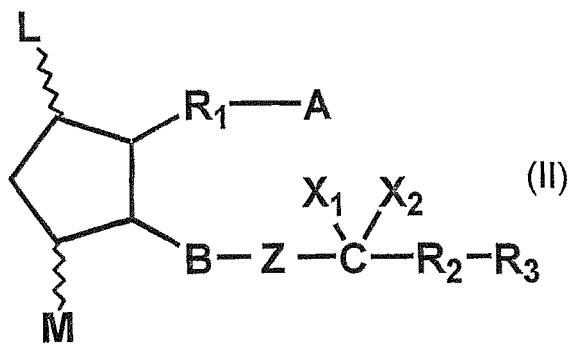
wherein R_4 and R_5 are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein R_4 and R_5 are not hydroxy and lower alkoxy at the same time;

5 time;

R₁ is a saturated or unsaturated bivalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

Ra is a saturated or unsaturated lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy; cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group, provided that Ra is substituted by halogen or Z is C=O.

[0025] A preferred compound used in the present invention is represented by the formula (II):

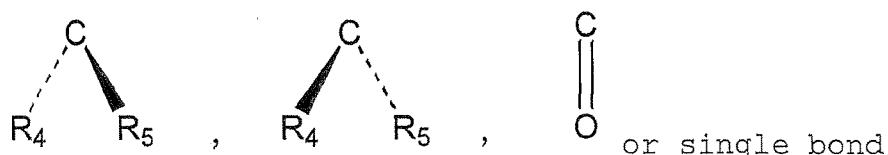


wherein L and M are hydrogen atom, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have one or more double bonds;

A is $-CH_3$, or $-CH_2OH$, $-COCH_2OH$, $-COOH$ or a functional derivative thereof;

B is single bond, $-CH_2-CH_2-$, $-CH=CH-$, $-C\equiv C-$, $-CH_2-CH_2-$
10 CH_2- , $-CH=CH-CH_2-$, $-CH_2-CH=CH-$, $-C\equiv C-CH_2-$ or $-CH_2-C\equiv C-$;

Z is



wherein R_4 and R_5 are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein
15 R_4 and R_5 are not hydroxy and lower alkoxy at the same time;

X_1 and X_2 are hydrogen, lower alkyl, or halogen;

R_1 is a saturated or unsaturated bivalent lower or

medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur;

5 R₂ is a single bond or lower alkylene; and

R₃ is lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group, provided that 10 one of X₁ and X₂ is substituted by halogen or Z is C=O.

[0026] In the above formula, the term "unsaturated" in the definitions for R₁ and Ra is intended to include at least one or more double bonds and/or triple bonds that are isolatedly, separately or serially present between carbon 15 atoms of the main and/or side chains. According to the usual nomenclature, an unsaturated bond between two serial positions is represented by denoting the lower number of the two positions, and an unsaturated bond between two distal positions is represented by denoting both of the 20 positions.

[0027] The term "lower or medium aliphatic hydrocarbon" refers to a straight or branched chain hydrocarbon group having 1 to 14 carbon atoms (for a side chain, 1 to 3 carbon atoms are preferable) and preferably 1 to 10, 25 especially 1 to 8 carbon atoms.

[0028] The term "halogen atom" covers fluorine, chlorine, bromine and iodine.

[0029] The term "lower" throughout the specification is intended to include a group having 1 to 6 carbon atoms 5 unless otherwise specified.

[0030] The term "lower alkyl" refers to a straight or branched chain saturated hydrocarbon group containing 1 to 6 carbon atoms and includes, for example, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, t-butyl, pentyl and 10 hexyl.

[0031] The term "lower alkylene" refers to a straight or branched chain bivalent saturated hydrocarbon group containing 1 to 6 carbon atoms and includes, for example, methylene, ethylene, propylene, isopropylene, butylene, 15 isobutylene, t-butylene, pentylene and hexylene.

[0032] The term "lower alkoxy" refers to a group of lower alkyl-O-, wherein lower alkyl is as defined above.

[0033] The term "hydroxy(lower)alkyl" refers to a lower alkyl as defined above which is substituted with at least 20 one hydroxy group such as hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl and 1-methyl-1-hydroxyethyl.

[0034] The term "lower alkanoyloxy" refers to a group represented by the formula RCO-O-, wherein RCO- is an acyl group formed by oxidation of a lower alkyl group as defined 25 above, such as acetyl.

[0035] The term "cyclo(lower)alkyl" refers to a cyclic group formed by cyclization of a lower alkyl group as defined above but contains three or more carbon atoms, and includes, for example, cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

[0036] The term "cyclo(lower)alkyloxy" refers to the group of cyclo(lower)alkyl-O-, wherein cyclo(lower)alkyl is as defined above.

[0037] The term "aryl" may include unsubstituted or substituted aromatic hydrocarbon rings (preferably monocyclic groups), for example, phenyl, tolyl, xylyl. Examples of the substituents are halogen atom and halo(lower)alkyl, wherein halogen atom and lower alkyl are as defined above.

[0038] The term "aryloxy" refers to a group represented by the formula ArO-, wherein Ar is aryl as defined above.

[0039] The term "heterocyclic group" may include mono- to tri-cyclic, preferably monocyclic heterocyclic group which is 5 to 14, preferably 5 to 10 membered ring having optionally substituted carbon atom and 1 to 4, preferably 1 to 3 of 1 or 2 type of hetero atoms selected from nitrogen atom, oxygen atom and sulfur atom. Examples of the heterocyclic group include furyl, thienyl, pyrrolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, imidazolyl, pyrazolyl, furazanyl, pyranyl, pyridyl, pyridazinyl,

pyrimidyl, pyrazinyl, 2-pyrrolinyl, pyrrolidinyl, 2-imidazolinyl, imidazolidinyl, 2-pyrazolinyl, pyrazolidinyl, piperidino, piperazinyl, morpholino, indolyl, benzothienyl, quinolyl, isoquinolyl, purinyl, quinazolinyl, carbazolyl, 5 acridinyl, phenanthridinyl, benzimidazolyl, benzimidazolinyl, benzothiazolyl, phenothiazinyl. Examples of the substituent in this case include halogen, and halogen substituted lower alkyl group, wherein halogen atom and lower alkyl group are as described above.

10 [0040] The term "heterocyclic-oxy group" means a group represented by the formula HcO^- , wherein Hc is a heterocyclic group as described above.

[0041] The term "functional derivative" of A includes salts (preferably pharmaceutically acceptable salts), 15 ethers, esters and amides.

[0042] Suitable "pharmaceutically acceptable salts" include conventionally used non-toxic salts, for example a salt with an inorganic base such as an alkali metal salt (such as sodium salt and potassium salt), an alkaline earth metal salt (such as calcium salt and magnesium salt), an ammonium salt; or a salt with an organic base, for example, an amine salt (such as methylamine salt, dimethylamine salt, cyclohexylamine salt, benzylamine salt, piperidine salt, ethylenediamine salt, ethanolamine salt, diethanolamine 20 salt, triethanolamine salt, tris(hydroxymethylamino)ethane 25 salt,

salt, monomethyl- monoethanolamine salt, procaine salt and caffeine salt), a basic amino acid salt (such as arginine salt and lysine salt), tetraalkyl ammonium salt and the like. 5 These salts may be prepared by a conventional process, for example from the corresponding acid and base or by salt interchange.

[0043] Examples of the ethers include alkyl ethers, for example, lower alkyl ethers such as methyl ether, ethyl ether, propyl ether, isopropyl ether, butyl ether, isobutyl ether, t-butyl ether, pentyl ether and 1-cyclopropyl ethyl ether; and medium or higher alkyl ethers such as octyl ether, diethylhexyl ether, lauryl ether and cetyl ether; unsaturated ethers such as oleyl ether and linolenyl ether; lower alkenyl ethers such as vinyl ether, allyl ether; 15 lower alkynyl ethers such as ethynyl ether and propynyl ether; hydroxy(lower)alkyl ethers such as hydroxyethyl ether and hydroxyisopropyl ether; lower alkoxy (lower)alkyl ethers such as methoxymethyl ether and 1-methoxyethyl ether; optionally substituted aryl ethers such as phenyl ether, tosyl ether, t-butylphenyl ether, salicyl ether, 20 3,4-di-methoxyphenyl ether and benzamidophenyl ether; and aryl(lower)alkyl ethers such as benzyl ether, trityl ether and benzhydryl ether.

[0044] Examples of the esters include aliphatic esters, 25 for example, lower alkyl esters such as methyl ester, ethyl

ester, propyl ester, isopropyl ester, butyl ester, isobutyl ester, t-butyl ester, pentyl ester and 1-cyclopropylethyl ester; lower alkenyl esters such as vinyl ester and allyl ester; lower alkynyl esters such as ethynyl ester and 5 propynyl ester; hydroxy(lower)alkyl ester such as hydroxyethyl ester; lower alkoxy (lower) alkyl esters such as methoxymethyl ester and 1-methoxyethyl ester; and optionally substituted aryl esters such as, for example, phenyl ester, toyl ester, t-butylphenyl ester, salicyl 10 ester, 3,4-di-methoxyphenyl ester and benzamidophenyl ester; and aryl(lower)alkyl ester such as benzyl ester, trityl ester and benzhydryl ester.

[0045] The amide of A mean a group represented by the formula -CONR'R", wherein each of R' and R" is hydrogen, 15 lower alkyl, aryl, alkyl- or aryl-sulfonyl, lower alkenyl and lower alkynyl, and include for example lower alkyl amides such as methylamide, ethylamide, dimethylamide and diethylamide; arylamides such as anilide and toluidide; and alkyl- or aryl-sulfonylamides such as methylsulfonylamine, 20 ethylsulfonyl-amide and tolylsulfonylamine.

[0046] Preferred examples of L and M include hydrogen, hydroxy and oxo, and especially, M is hydroxy and L is oxo which has a 5-membered ring structure of, so called, PGE type.

25 [0047] Preferred example of A is -COOH, its

pharmaceutically acceptable salt, ester or amide thereof.

[0048] Preferred example of X_1 and X_2 are both being halogen atoms, and more preferably, fluorine atoms, so called 16,16-difluoro type.

5 [0049] Preferred R_1 is a hydrocarbon residue containing 1-10 carbon atoms, preferably 6-10 carbon atoms. Further, at least one carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur.

Examples of R_1 include, for example, the following groups:

10 $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}=\text{CH}-$,

$-\text{CH}_2-\text{C}\equiv\text{C}-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{O}-\text{CH}_2-$,

15 $-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{O}-\text{CH}_2-$,

$-\text{CH}_2-\text{C}\equiv\text{C}-\text{CH}_2-\text{O}-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}=\text{CH}-$,

20 $-\text{CH}_2-\text{C}\equiv\text{C}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}(\text{CH}_3)-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}(\text{CH}_3)-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

$-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$,

25 $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}=\text{CH}-$,

-CH₂-C≡C-CH₂-CH₂-CH₂-CH₂-CH₂- and
-CH₂-CH₂-CH₂-CH₂-CH₂-CH₂-CH(CH₃)-CH₂-.

[0050] Preferred Ra is a hydrocarbon containing 1-10 carbon atoms, more preferably, 1-8 carbon atoms. Ra may 5 have one or two side chains having one carbon atom.

[0051] Most preferred embodiment of the prostaglandin compound is 13,14-dihydro-15-keto-16,16-difluoro-prostaglandin E₁ compound or 13,14-dihydro-15-keto-16,16-difluoro-18-methyl- prostaglandin E₁ compound.

[0052] The configuration of the ring and the α - and/or ω chains in the above formulae (I) and (II) may be the same as or different from that of the primary PGs. However, the present invention also includes a mixture of a compound having a primary type configuration and a compound of a 15 non-primary type configuration.

[0053] In the present invention, the PG compound which is dihydro between 13 and 14, and keto(=O) at 15 position may be in the keto-hemiacetal equilibrium by formation of a hemiacetal between hydroxy at position 11 and keto at 20 position 15.

[0054] For example, it has been revealed that when both of X₁ and X₂ are halogen atoms, especially, fluorine atoms, the compound contains a tautomeric isomer, bicyclic compound.

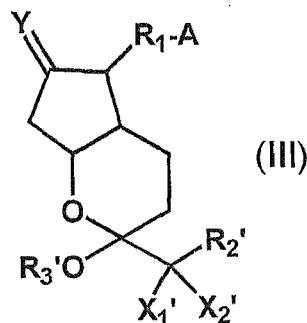
[0055] If such tautomeric isomers as above are present,

the proportion of both tautomeric isomers varies with the structure of the rest of the molecule or the kind of the substituent present. Sometimes one isomer may predominantly be present in comparison with the other.

5 However, it is to be appreciated that the present invention includes both isomers.

[0056] Further, the 15-keto-PG compounds used in the invention include the bicyclic compound and analogs or derivatives thereof.

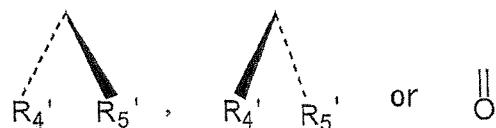
10 [0057] The bicyclic compound is represented by the formula (III):



wherein, A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

15 X_1' and X_2' are hydrogen, lower alkyl, or halogen;

Y is



wherein R_4' and R_5' are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein
20 R_4' and R_5' are not hydroxy and lower alkoxy at the same time.

[0058] R_1 is a saturated or unsaturated divalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

[0059] R_2' is a saturated or unsaturated lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy; cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group.

[0060] R_3' is hydrogen, lower alkyl, cyclo(lower)alkyl, aryl or heterocyclic group.

[0061] Furthermore, while the compounds used in the invention may be represented by a formula or name based on keto-type regardless of the presence or absence of the isomers, it is to be noted that such structure or name does not intend to exclude the hemiacetal type compound.

[0062] In the present invention, any of isomers such as the individual tautomeric isomers, the mixture thereof, or optical isomers, the mixture thereof, a racemic mixture, and other steric isomers may be used in the same purpose.

[0063]

The Pharmaceutically Suitable Excipient

According to the invention, the combination may be formulated in a pharmaceutical composition.. The 5 composition may be formulated as separate dosage forms each comprising individual active ingredient with a pharmaceutically suitable excipient, or as a single dosage form comprising two active ingredients with a pharmaceutically suitable excipient. The pharmaceutically 10 suitable excipient may be, therefore, selected depending on the desired form of the composition. According to the invention, "pharmaceutically suitable excipient" means an inert substance, which is suitable for the form, combined with the active ingredient of the invention.

15 [0064] For example, solid composition for oral administration of the present invention may include tablets, preparations, granules and the like. In such a solid composition, one or more active ingredients may be mixed with at least one inactive diluent, for example, lactose, 20 mannitol, glucose, hydroxypropyl cellulose, microcrystalline cellulose, starch, polyvinyl pyrrolidone, magnesium aluminate metasilicate and the like. According to the usual work-up, the composition may contain additives other than inactive diluent, for example, lubricant such as 25 magnesium stearate; disintegrant such as fibrous calcium

gluconate; stabilizer such as cyclodextrin, for example, α , β - or γ -cyclodextrin; etherified cyclodextrin such as dimethyl- α -, dimethyl- β -, trimethyl- β -, or hydroxypropyl- β -cyclodextrin; branched cyclodextrin such as glucosyl-, 5 maltosyl-cyclodextrin; formylated cyclodextrin, cyclodextrin containing sulfur; phospholipid and the like. When the above cyclodextrins are used, an inclusion compound with cyclodextrins may be sometimes formed to enhance stability. Alternatively, phospholipid may be 10 sometimes used to form a liposome, resulting in enhanced stability.

[0065] Tablets or pills may be coated with film soluble in the stomach or intestine such as sugar, gelatin, hydroxypropyl cellulose, or hydroxypropylmethyl cellulose 15 phthalate as needed. Further, they may be formed as capsules with absorbable substances such as gelatins. Preferably, the composition is formulated in a soft gelatin capsule with liquid contents of the specific prostaglandin compound and a medium chain fatty acid triglyceride. 20 Examples of the medium chain fatty acid triglyceride used in the present invention include a triglyceride of a saturated or unsaturated fatty acid having 6-14 carbon atoms which may have a branched chain. A preferred fatty acid is a straight chain saturated fatty acid, for example 25 caproic acid (C6), caprylic acid (C8), capric acid (C10),

lauric acid (C12) and myristic acid (C14). In addition, two or more medium chain fatty acid triglycerides may be used in combination. Further suitable excipients are disclosed in the published PCT application WO 01/27099.

5 [0066] A liquid composition for oral administration may be pharmaceutically acceptable emulsion, solution, suspension, syrup, or elixir, as well as generally used inactive diluent. Such composition may contain, in addition to the inactive diluent, adjuvants such as
10 lubricants and suspensions, sweetening agents, flavoring agents, preservatives, solubilizers, anti-oxidants and the like. The details of the additives may be selected from those described in any general textbooks in the pharmaceutical field. Such liquid compositions may be
15 directly enclosed in soft capsules. Solutions for parenteral administration, for example, suppository, enema and the like according to the present invention include sterile, aqueous or non-aqueous solution, suspension, emulsion, detergent and the like. The aqueous solution and
20 suspension includes, for example, distilled water, physiological saline and Ringer's solution.

25 [0067] The non-aqueous solution and suspension include, for example, propylene glycol, polyethylene glycol, fatty acid triglyceride, and vegetable oil such as olive oil, alcohols such as ethanol, polysorbate and the like. Such

composition may contain adjuvants such as preservatives, wetting agent, emulsifier, dispersant, anti-oxidants and the like.

[0068] Examples of the injectable compositions of the 5 present invention for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions and emulsions. Diluents for the aqueous solution or suspension may include, for example, distilled water for injection, physiological saline and Ringer's solution.

[0069] Non-aqueous diluents for solution and suspension 10 may include, for example, propylene glycol, polyethylene glycol, vegetable oils such as olive oil, alcohols such as ethanol and polysorbate. The composition may further comprise additives such as preservatives, wetting agents, 15 emulsifying agents, dispersing agents and the like. They may be sterilized by filtration through, e.g. a bacteria-retaining filter, compounding with a sterilizer, or by means of gas or radioisotope irradiation sterilization. The injectable composition may also be provided as a 20 sterilized powder composition to be dissolved in a sterilized solvent for injection before use.

[0070] Another form of the present invention is 25 suppository or pessary, which may be prepared by mixing active ingredients into a conventional base such as cacao butter that softens at body temperature, and nonionic

surfactants having suitable softening temperatures may be used to improve absorbability.

[0071] According to the method of the invention, the combination of the present invention can be administered 5 systemically or locally by means of oral or parental administration, including a suppository, enema and the like. Single or multiple compositions may be administered to achieve the desired dose. According to the method, individual compounds in the combination may be administered 10 simultaneously, separately, or sequentially.

[0072] According to the present invention, a mammalian subject may be treated by the instant invention by administering the compound used in the present invention. The mammalian subject may be any subject including a human. 15 The compound may be applied systemically or topically. Usually, the compound may be administered by oral administration, intravenous injection (including infusion), subcutaneous injection, intra rectal administration, intra vaginal administration, transdermal administration and the 20 like. The dose may vary depending on the strain of the animal, age, body weight, symptom to be treated, desired therapeutic effect, administration route, term of treatment and the like. A satisfactory effect can be obtained by systemic administration 1-4 times per day or continuous 25 administration at combination with the amount of 0.001-

100000 µg, more preferably 0.01-10000 µg, especially 0.1-1000 µg of specific prostaglandin compound, and 0.01-100000mg, more preferably 0.1-10000mg, of NSAID at a daily dose.

5 [0073] The term "combination" used herein means NSAID and the specific prostaglandin compound are both administered to a patient simultaneously in the form of a single entity or dosage, or are both administered to a patient as separate entities either simultaneously or 10 sequentially with no specific time limits, wherein such administration provides therapeutically effective levels of the two components in the body, preferably at the same time.

[0074] The combination of the present invention is useful for the treatment of a disease or condition which is 15 one of the indications for NSAID use. For example, the combination is useful for the treatment of arthritis, rheumatoid arthritis, osteoarthritis, spondylitis, ankylosing spondylitis, juvenile arthritis, juvenile rheumatoid arthritis, tendonitis, bursitis, gout, pain, and 20 dysmenorrhea.

[0075] The term "treatment" used herein includes any means of control such as prevention, care, relief of the condition, attenuation of the condition and arrest of progression.

25 [0076] The combination of the present invention is

useful for the treatment of pain from various etiologies.

[0077] The term "pain from various etiology" includes, but is not limited to, inflammatory pain, hyperalgesia and, in particular, chronic pain, and means in particular pain consequential to trauma, e.g. associated with burns, sprains, fracture or the like, subsequent to surgical intervention, e.g. as post-operative analgesics, chemotherapy-induced pain, as well as inflammatory pain of diverse genesis, e.g. bone and joint pain (osteoarthritis), myofascial pain (muscular injury, fibromyalgia), lower back pain, chronic inflammatory pain, chronic neuropathic pain, e.g. diabetic neuropathy, phantom limb pain and perioperative pain (general surgery, gynecologic surgery) as well as pain associated with, e.g., angina, menstruation or cancer.

[0078] The combination of the present invention is further useful for the treatment of Alzheimer disease or cancers.

[0079] The term "Cancer" includes esophageal cancer, gastric cancer, duodenal cancer, small intestinal cancer, appendiceal cancer, large bowel cancer, colon cancer, rectum cancer, colorectal cancer, anal cancer, pancreatic cancer, liver cancer, gallbladder cancer, spleen cancer, renal cancer, bladder cancer, prostatic cancer, testicular

cancer, uterine cancer, ovarian cancer, mammary cancer, pulmonary cancer and thyroid cancer.

[0080] The further details of the present invention will follow with reference to test examples, which, however, are 5 not intended to limit the present invention.

[0081] Example 1

Male Crl: CD (SD) rats were orally administered with naproxen at 200 mg/kg in combination with Compound A (13,14-dihydro-15-keto-16,16-difluoro-18(S)-methyl-PGE₁ or 10 vehicle. The animal was euthanized by cervical dislocation 5 hours after administration of the dosing preparation, and the stomach was removed. The stomach was filled with 10 mL of saline and fixed in 1% formalin. Then the stomach was opened along the greater curvature and gently washed with 15 saline. The length of the major axis of each gastric ulcer was measured with digital slide gauges, and the total length of the major axes of all gastric ulcers (ulcer index) was obtained.

[0082] As shown in Table 1, the ulcer formation 20 induced by naproxen at 200 mg/kg was reduced by the combined administration of Compound A at 1, 10 and 100 µg/kg with naproxen (200 mg/kg) in a dose-dependent manner. Significant reductions of ulcer formation were observed in the "Compound A (10 µg/kg) + naproxen (200 25 mg/kg) group" and "Compound A (100 µg/kg) + naproxen (200

mg/kg) group" compared to "Vehicle + naproxen (200 mg/kg) group". The combined administration of lansoprazole, a proton pump inhibitor used for the treatment of gastric ulcer in clinic, at 1000 μ g/kg with naproxen did not have a 5 significant effect on the naproxen-induced gastric ulcers.

[0083] The results indicate that Compound A has a potent protective effect against ulcer formation in combination with non-steroidal anti-inflammatory drug (NSAID).

[0084]

10 Table 1 Effect of Compound A in combination with naproxen on naproxen-induced gastric ulcer in rats

Group	n	Ulcer Index (Mean \pm S.E.)	%Inhibition ^a
Vehicle (0.01% Tween 80) + naproxen (200 mg/kg)	10	81.48 \pm 15.05	-
Compound A (1 μ g/kg) + naproxen (200 mg/kg)	10	63.47 \pm 10.87	22%
Compound A (10 μ g/kg) + naproxen (200 mg/kg)	10	12.15 \pm 3.45*	85%
Compound A (100 μ g/kg) + naproxen (200 mg/kg)	10	0.50 \pm 0.40**	99.4%
Lansoprazole (1000 μ g/kg) + naproxen (200 mg/kg)	10	68.09 \pm 7.61	16%

* p<0.05, ** p<0.01 compared to "vehicle + naproxen group" (Dunnett-type test)

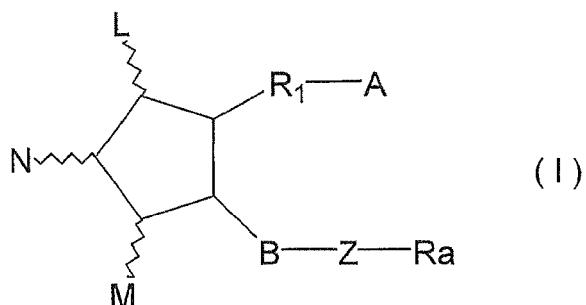
a Calculated as: % inhibition = (1-mean ulcer index in 15 test group/mean ulcer index in control group)X 100

CLAIMS

1. A pharmaceutical combination comprising:

(a) a NSAID, and

5 (b) a prostaglandin (PG) compound represented by the formula (I):

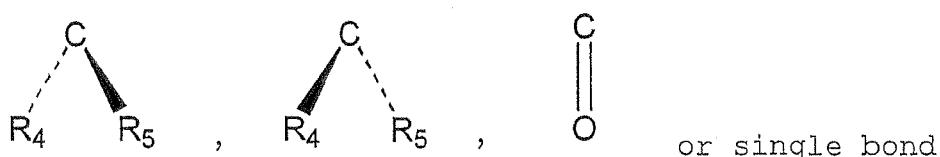


wherein L, M and N are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, 10 wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have at least one double bond;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

15 B is single bond, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-\text{CH}_2-$, $-\text{CH}_2-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-\text{CH}_2-$ or $-\text{CH}_2-\text{C}\equiv\text{C}-$;

Z is



wherein R_4 and R_5 are hydrogen, hydroxy, halogen,

lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein R₄ and R₅ are not hydroxy and lower alkoxy at the same time;

R₁ is a saturated or unsaturated bivalent lower or 5 medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

Ra is a saturated or unsaturated lower or medium 10 aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or 15 heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy; cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group, provided that Ra is substituted by halogen or Z is C=O for the treatment of a condition or disease which is one of 20 the indications for NSAIDs use.

2. The combination as described in Claim 1, which is for the treatment of pain.

3. The combination as described in Claim 2, wherein the pain is chronic pain.

25 4. The combination as described in Claim 1, which is

for the treatment of Alzheimer disease.

5. The combination as described in Claim 1, which is
for the treatment of cancer.

6. The combination as described in any one of Claims
5 1-5, wherein said prostaglandin compound is 16-mono or
dihalogen-prostaglandin compound.

7. The combination as described in any one of Claims
1-5, wherein said prostaglandin compound is 15-keto-
prostaglandin compound.

10 8. The combination as described in any one of
Claims 1-5, wherein said prostaglandin compound is 13,14-
dihydro-16-mono or dihalogen-prostaglandin compound.

9. The combination as described in any one of Claims
1-5, wherein said prostaglandin compound is 13,14-dihydro-
15 15-keto-prostaglandin compound.

10. The combination as described in any one of Claims
1-5, wherein said prostaglandin compound is 13,14-dihydro-
15-keto-16-mono or dihalogen-prostaglandin compound.

11. The combination as described in any one of Claims
20 1-5, wherein said prostaglandin compound is 13,14-dihydro-
16-mono or difluoro-prostaglandin compound.

12. The combination as described in any one of Claims
1-5, wherein said prostaglandin compound is 15-keto-16-mono
or difluoro-prostaglandin compound.

25 13. The combination as described in any one of Claims

1-5, wherein said prostaglandin compound is 13,14-dihydro-15-keto-16-mono or difluoro-prostaglandin compound.

14. The combination as described in any one of Claims 1-5, wherein said prostaglandin compound is 13,14-dihydro-5 16-mono or dihalogen-prostaglandin E compound.

15. The combination as described in any one of Claims 1-5, wherein said prostaglandin compound is 15-keto-16-mono or dihalogen-prostaglandin E compound.

16. The combination as described in any one of Claims 10 1-5, wherein said prostaglandin compound is 13,14-dihydro-15-keto-16-mono or dihalogen-prostaglandin E compound.

17. The combination as described any one of Claims 1-5, wherein said prostaglandin compound is 13,14-dihydro-16,16-difluoro -prostaglandin E₁ compound.

15 18. The combination as described in any one of Claims 1-5, wherein said prostaglandin compound is 13,14-dihydro-15-keto-prostaglandin E₁ compound.

19. The combination as described in any one of Claims 1-5, wherein said prostaglandin compound is 13,14-dihydro-20 15-keto-16,16-difluoro-prostaglandin E₁ compound or 13,14-dihydro-15-keto-16,16-difluoro-18-methyl-prostaglandin E₁ compound.

20. The combination as described in any one of Claims 1-5, wherein said prostaglandin compound is 13,14-dihydro-25 15-keto-16,16-difluoro-prostaglandin E₁ or 13,14-dihydro-

15-keto-16,16-difluoro-18-methyl-prostaglandin E₁.

21. The combination as described in any one of Claims 1-20, wherein the NSAID is selected from the group consisting of salicylates, indomethacin, flurbiprofen, 5 diclofenac, ketorolac, naproxen, piroxicam, tebufelone, ibuprofen, etodolac, nabumetone, tenidap, alcofenac, antipyrine, aminopyrine, dipyrone, aminopyrone, phenylbutazone, clofezone, oxyphenbutazone, prexazone, apazone, benzydamine, bucolome, cinchopen, clonixin, 10 ditrazol, epirizole, fenoprofen, floctafeninl, flufenamic acid, glaphenine, indoprofen, ketoprofen, meclofenamic acid, mefenamic acid, niflumic acid, phenacetin, salidifamides, sulindac, suprofen, tolmetin, a pharmaceutically acceptable salt thereof, and a mixture thereof.

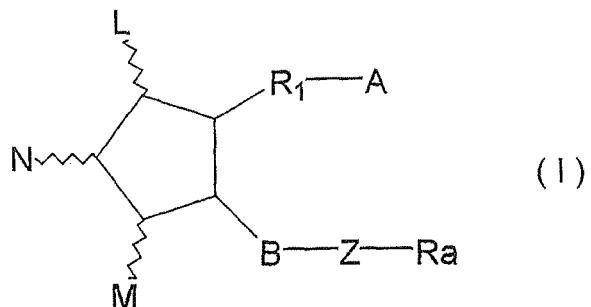
15 22. The combination as described in claim 21, wherein the NSAID is naproxen.

23. The combination as described in any one of Claims 1-22 for simultaneous, separate or sequential use.

24. A pharmaceutical combination comprising:

20 (a) a NSAID, and

 (b) a prostaglandin (PG) compound represented by the formula (I):

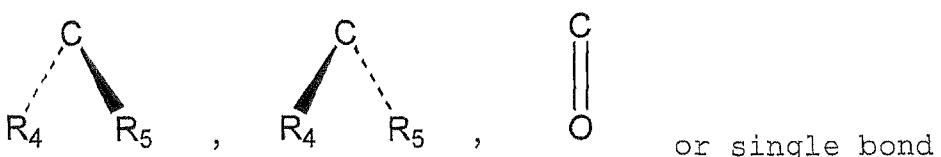


wherein L, M and N are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have at least one double bond;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

B is single bond, $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{CH}_2-\text{CH}_2-$
10 CH_2- , $-\text{CH}=\text{CH}-\text{CH}_2-$, $-\text{CH}_2-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-\text{CH}_2-$ or $-\text{CH}_2-\text{C}\equiv\text{C}-$;

Z is



wherein R_4 and R_5 are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy or hydroxy(lower)alkyl, wherein
15 R_4 and R_5 are not hydroxy and lower alkoxy at the same time;

R_1 is a saturated or unsaturated bivalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy,

oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

Ra is a saturated or unsaturated lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy; cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group, provided that Ra is substituted by halogen or Z is C=O.

25. The combination as described in Claim 24, wherein said prostaglandin compound is 16-mono or dihalogen-prostaglandin compound.

26. The combination as described in Claim 24, wherein said prostaglandin compound is 15-keto-prostaglandin compound.

27. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-16-mono or dihalogen-prostaglandin compound.

28. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-15-keto-prostaglandin compound.

29. The combination as described in Claim 24, wherein

said prostaglandin compound is 13,14-dihydro-15-keto-16-mono or dihalogen-prostaglandin compound.

30. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-16-mono or 5 difluoro-prostaglandin compound.

31. The combination as described in Claim 24, wherein said prostaglandin compound is 15-keto-16-mono or difluoro-prostaglandin compound.

32. The combination as described in Claim 24, wherein 10 said prostaglandin compound is 13,14-dihydro-15-keto-16-mono or difluoro-prostaglandin compound.

33. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-16-mono or dihalogen-prostaglandin E compound.

15 34. The combination as described in Claim 24, wherein said prostaglandin compound is 15-keto-16-mono or dihalogen-prostaglandin E compound.

35. The combination as described in Claim 24, wherein 20 said prostaglandin compound is 13,14-dihydro-15-keto-16-mono or dihalogen-prostaglandin E compound.

36. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-16,16-difluoro-prostaglandin E₁ compound.

37. The combination as described in Claim 24, wherein 25 said prostaglandin compound is 13,14-dihydro-15-keto-

prostaglandin E₁ compound.

38. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-15-keto-16,16-difluoro-prostaglandin E₁ compound or 13,14-dihydro-15-keto-16,16-difluoro-18-methyl-prostaglandin E₁ compound.

39. The combination as described in Claim 24, wherein said prostaglandin compound is 13,14-dihydro-15-keto-16,16-difluoro-prostaglandin E₁ or 13,14-dihydro-15-keto-16,16-difluoro-18-methyl- prostaglandin E₁.

10 40. The combination as described in any one of Claims 24-39, wherein the NSAID is selected from the group consisting of salicylates, indomethacin, flurbiprofen, diclofenac, ketorolac, naproxen, piroxicam, tebufelone, ibuprofen, etodolac, nabumetone, tenidap, alcofenac, 15 antipyrine, aminopyrine, dipyrone, aminopyrone, phenylbutazone, clofezone, oxyphenbutazone, prexazone, apazone, benzydamine, bucolome, cinchopen, clonixin, ditrazol, epirizole, fenoprofen, floctafeninl, flufenamic acid, glaphenine, indoprofen, ketoprofen, meclofenamic acid, 20 mefenamic acid, niflumic acid, phenacetin, salidifamides, sulindac, suprofen, tolmetin a pharmaceutically acceptable salt thereof, and a mixture thereof.

41. The combination as described in claim 40, wherein the NSAID is naproxen.

25 42. The combination as described in any one of Claims

24-41 for simultaneous, separate or sequential use.

43. A pharmaceutical composition comprising the combination as described in any one of Claims 1-42 in association with a pharmaceutically suitable excipient.

5 44. The composition as described in claim 43, wherein said pharmaceutically suitable excipient is orally acceptable.

45. A commercial package comprising a combination as described in any one of claim 1 to 42 together with
10 instructions for simultaneous, separate or sequential use thereof.