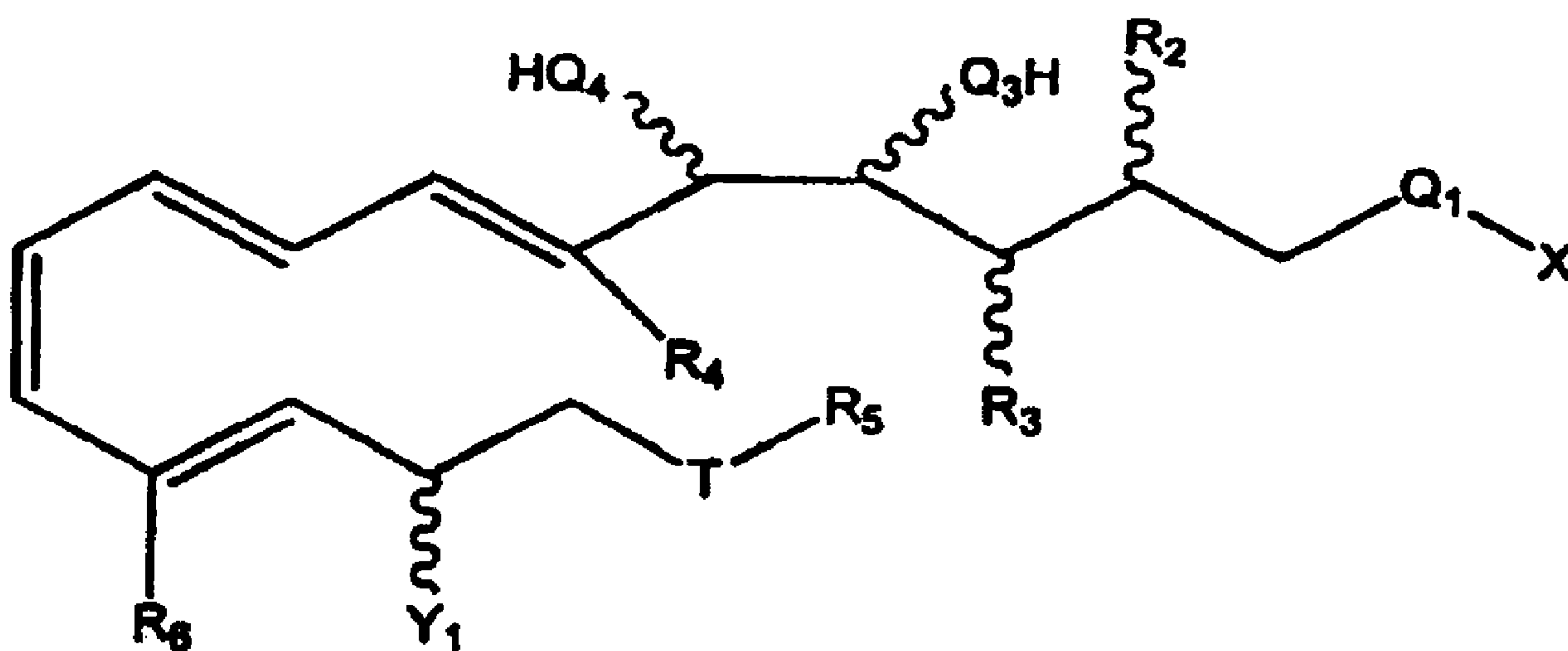




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(72) Inventeurs/Inventors:
SERHAN, CHARLES N., US;
LEVY, BRUCE D., US
(73) Propriétaire/Owner:
BRIGHAM AND WOMEN'S HOSPITAL, INC., US
(74) Agent: RICHES, MCKENZIE & HERBERT LLP

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RESPIRATOIRES INFLAMMATOIRES
(54) Title: LIPOXINS AND THEIR STABLE ANALOGS IN THE TREATMENT OF ASTHMA AND INFLAMMATORY
AIRWAY DISEASES



(57) Abrégé/Abstract:

The use of lipoxin analogs for the treatment or prevention of asthma and asthma related diseases is described. In particular, acetylenic lipoxin analogs are effective for the treatment and prevention of eosinophil recruitment involved with the inflammation processes associated with asthma, asthma like conditions, and lung injuries associated from airway inflammation or infection as brought about by leukocyte-mediated injury from within.



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(71) Applicant (*for all designated States except US*): **THE BRIGHAM AND WOMEN'S HOSPITAL, INC.** [US/US]; 75 Francis Street, Boston, MA 02115 (US).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **SERHAN, Charles, N.** [US/US]; 23 Smith Street, Needham, MA 02492 (US). **LEVY, Bruce, D.** [US/US]; 110 Anawan Avenue, West Roxbury, MA 02132 (US).

(74) Agents: **ROTHENBERGER, Scott, D.** et al.; Dorsey & Whitney LLP, Suite 1500, 50 South Sixth Street, Minneapolis, MN 55402-1498 (US).

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(54) Title: LIPOXINS AND ASPIRIN-TRIGGERED LIPOXINS AND THEIR STABLE ANALOGS IN THE TREATMENT OF ASTHMA AND INFLAMMATORY AIRWAY DISEASES

(57) Abstract: The use of lipoxin analogs for the treatment or prevention of asthma and asthma related diseases is described. In particular, acetylenic lipoxin analogs are effective for the treatment and prevention of eosinophil recruitment involved with the inflammation processes associated with asthma, asthma like conditions, and lung injuries associated from airway inflammation or infection as brought about by leukocyte-mediated injury from within.



WO 03/039533 A1

**Lipoxins and their Stable Analogs in the
Treatment of Asthma and Inflammatory Airway Diseases**

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH

[001] The work leading to this invention was supported in part by National
5 Institutes of Health (NIH) grants GM-38765, P01-DE13499 and K08 HL03788.
The U.S. Government therefore may have certain rights in the invention.

BACKGROUND

10 [003] Asthma is characterized by airway hyper-responsiveness and chronic
airway inflammation¹. Large numbers of eosinophils and T lymphocytes infiltrate
peribronchial tissues in asthmatics², trafficking into the lung an increased
capacity to generate cysteinyl leukotrienes (CysLT's) and T_H2 cytokines^{1, 3, 4}.
CysLT's have been associated with the asthmatic diathesis in both experimental
models and patients with asthma^{5, 6}. One of the many actions of T_H2 cytokines is
to up-regulate the expression of biosynthetic enzymes for eicosanoids - including
leukotrienes and lipoxins (LX's)^{7, 8}.

15 [004] LX's are a separate class of eicosanoids that are distinct in structure and
function⁷, and their biosynthesis is temporally dissociated from the formation and
impact of other eicosanoids⁹. LX's are generated in human tissues, including
airways¹⁰. LX's carry unique counter-regulatory actions that inhibit CysLT-
20 mediated vascular responses¹¹ and promote resolution of cytokine-driven acute

inflammation⁹. When administered to human cells *in vitro* or murine systems *in vivo*, at least two classes of receptors, CysLT1 receptors and LXA₄ receptors (designated ALX), can interact with LX's to mediate their actions^{12, 13}. A role for LX's in asthma has not yet been directly evaluated in well-qualified
5 experimental animal models.

[005] The global prevalence of asthma continues to increase, affecting millions of peoples' daily lives, but treatment is far from ideal¹⁸. Clinical responses to current therapies, such as inhaled corticosteroids and leukotriene modifiers are heterogeneous¹⁹ and even with optimal treatment there is a substantial burden of
10 unaddressed disease.

[006] With further evidence for serious toxicity from exogenous corticosteroids, new anti-inflammatory strategies are needed for asthma and other allergic illnesses.

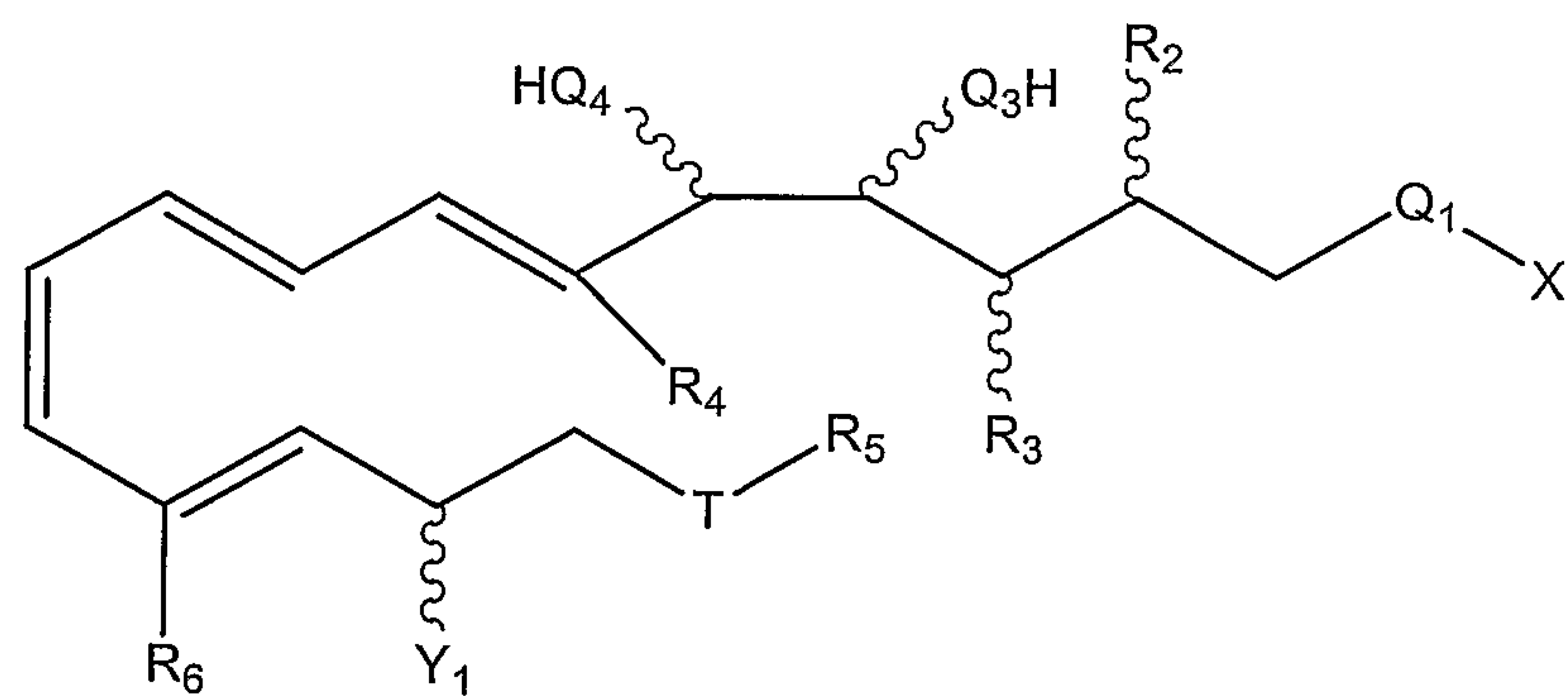
SUMMARY

15 [007] It has been surprisingly discovered that lipoxins (LX's) of the invention, discussed *infra*, decrease both airway inflammation and hyper-responsiveness in response to allergen sensitization and aerosol challenge. Consequently, the lipoxins disclosed herein are useful for the treatment and/or prevention of asthma and related airway and respiratory inflammatory disorders.

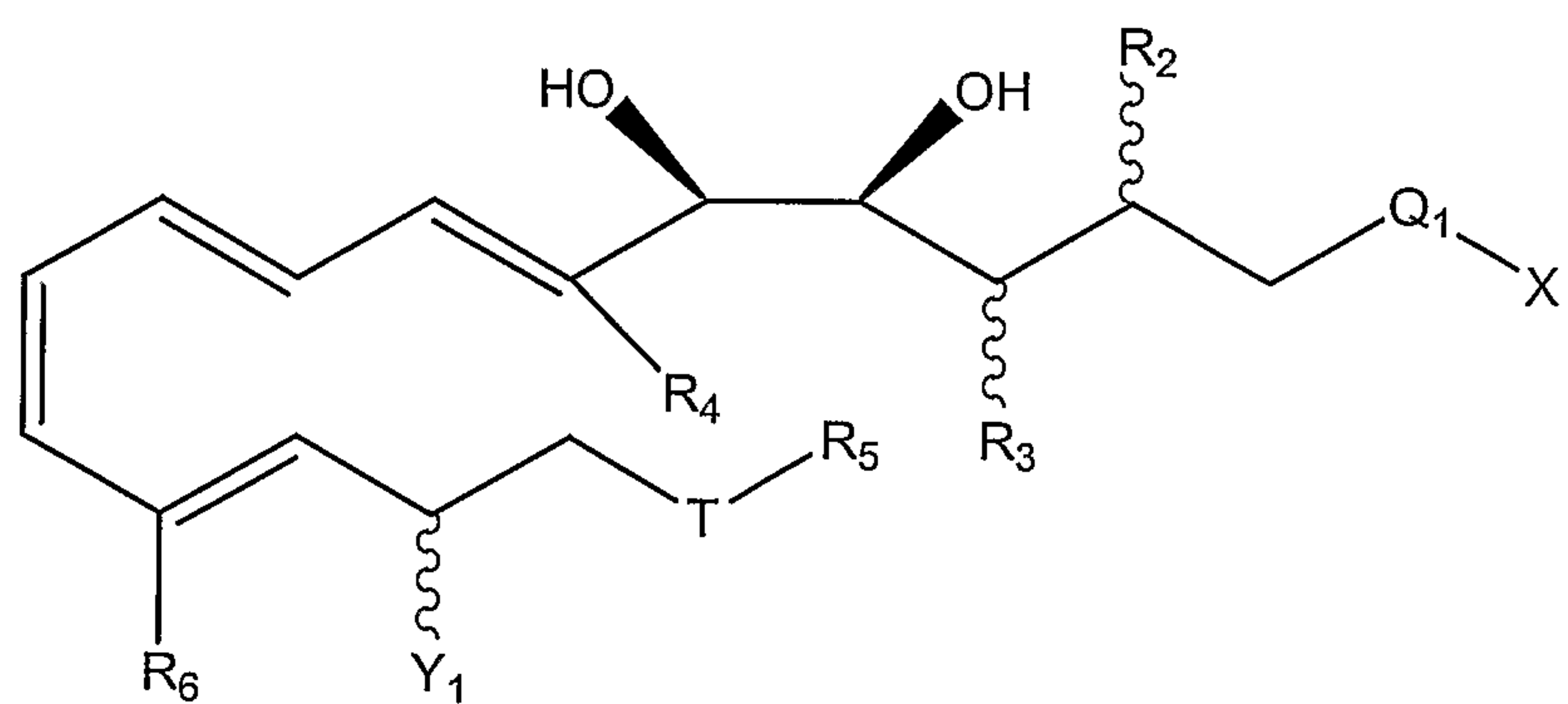
20 [008] The present invention pertains to methods for treating or preventing conditions associated with asthma, a related airway or respiratory inflammatory disorder. The methods include administration to a subject, an effective amount of a lipoxin analog, having the formula described *infra*, such that the asthma or related airway or respiratory inflammatory disorder is treated.

25 [009] Exemplary compounds useful in the treatment of the above-identified ailments include:

[010] Formula (I)

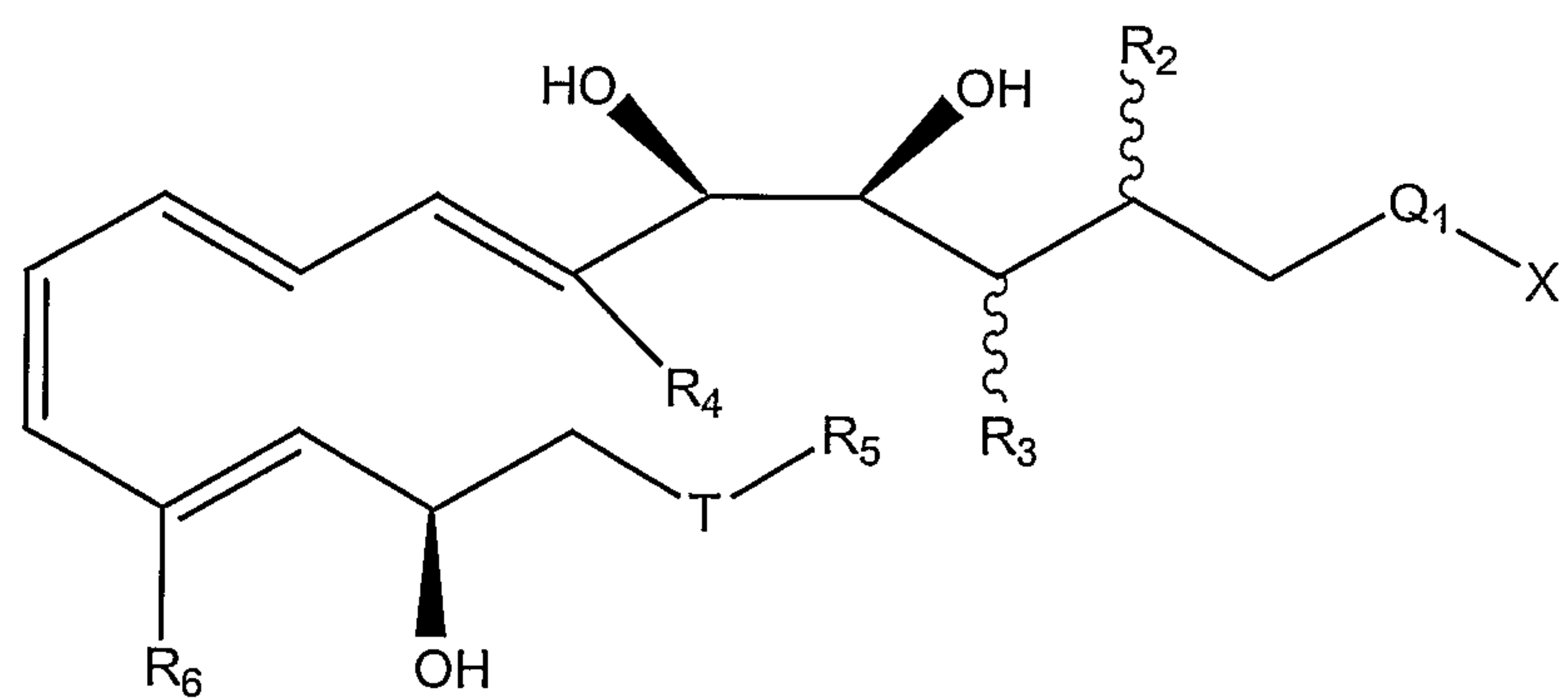


[011] Formula (II)

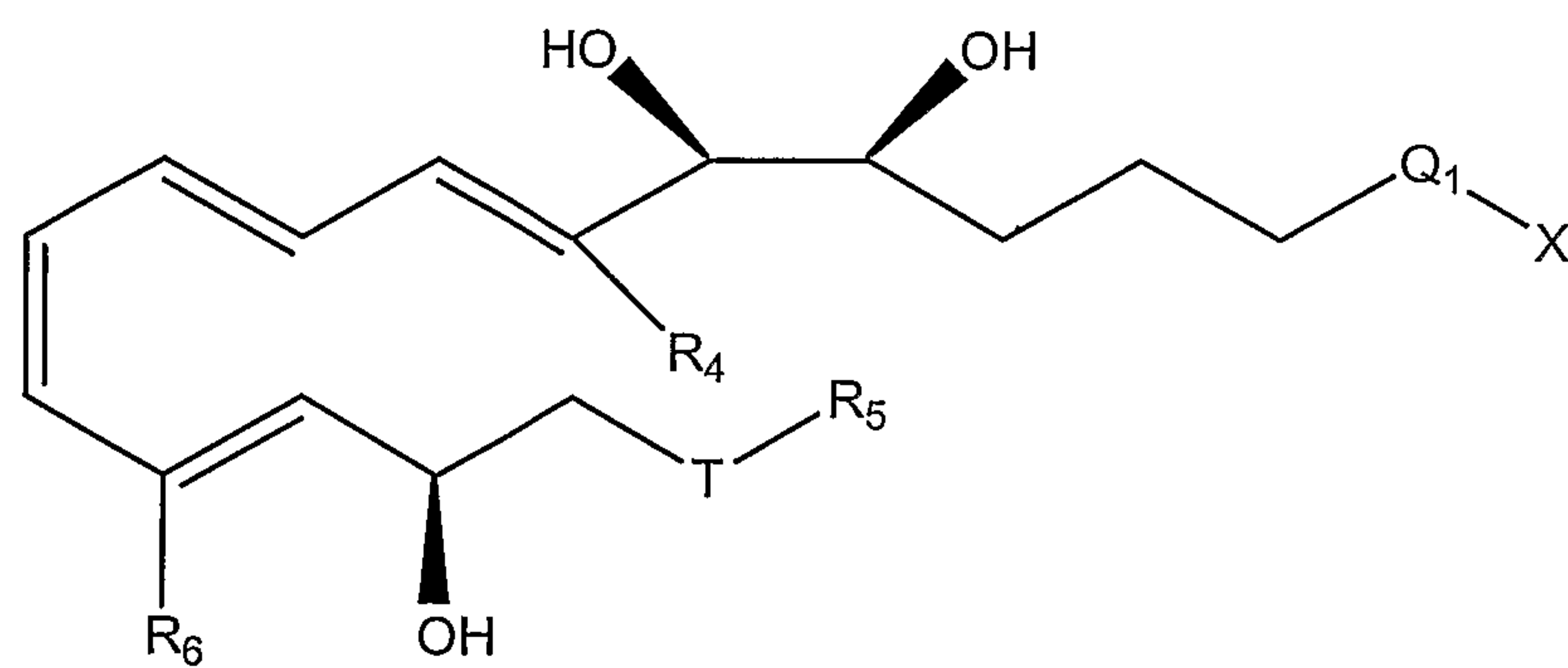


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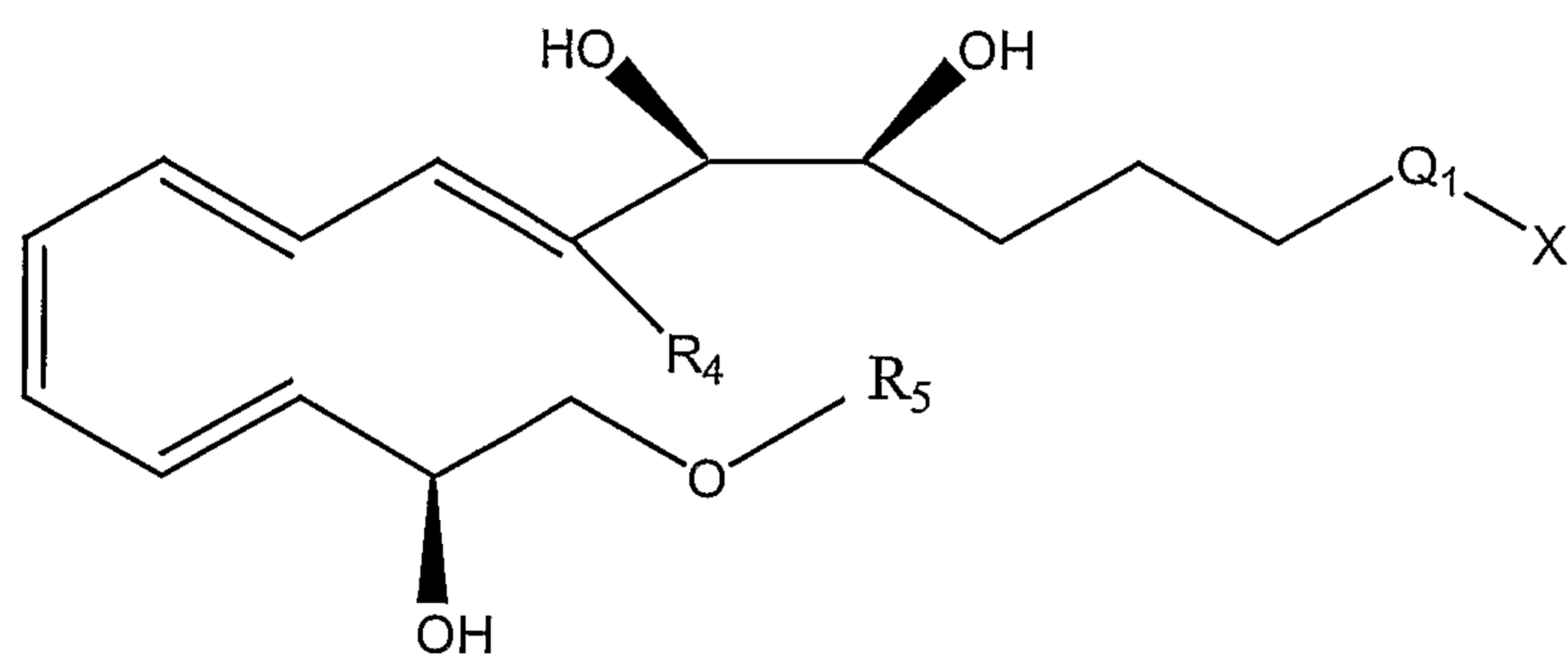
[012] Formula (III)



10 [013] Formula (IV)

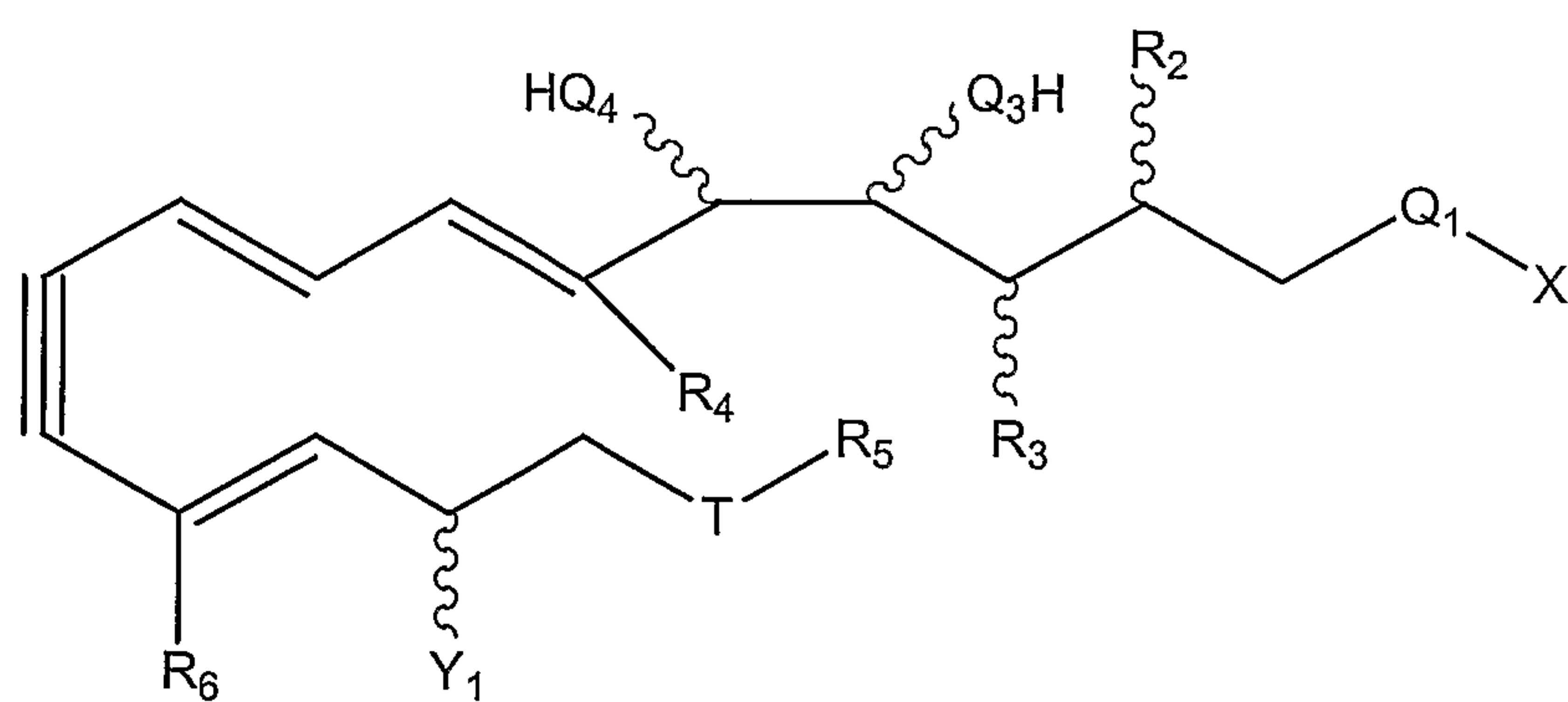


[014] Formula (V)

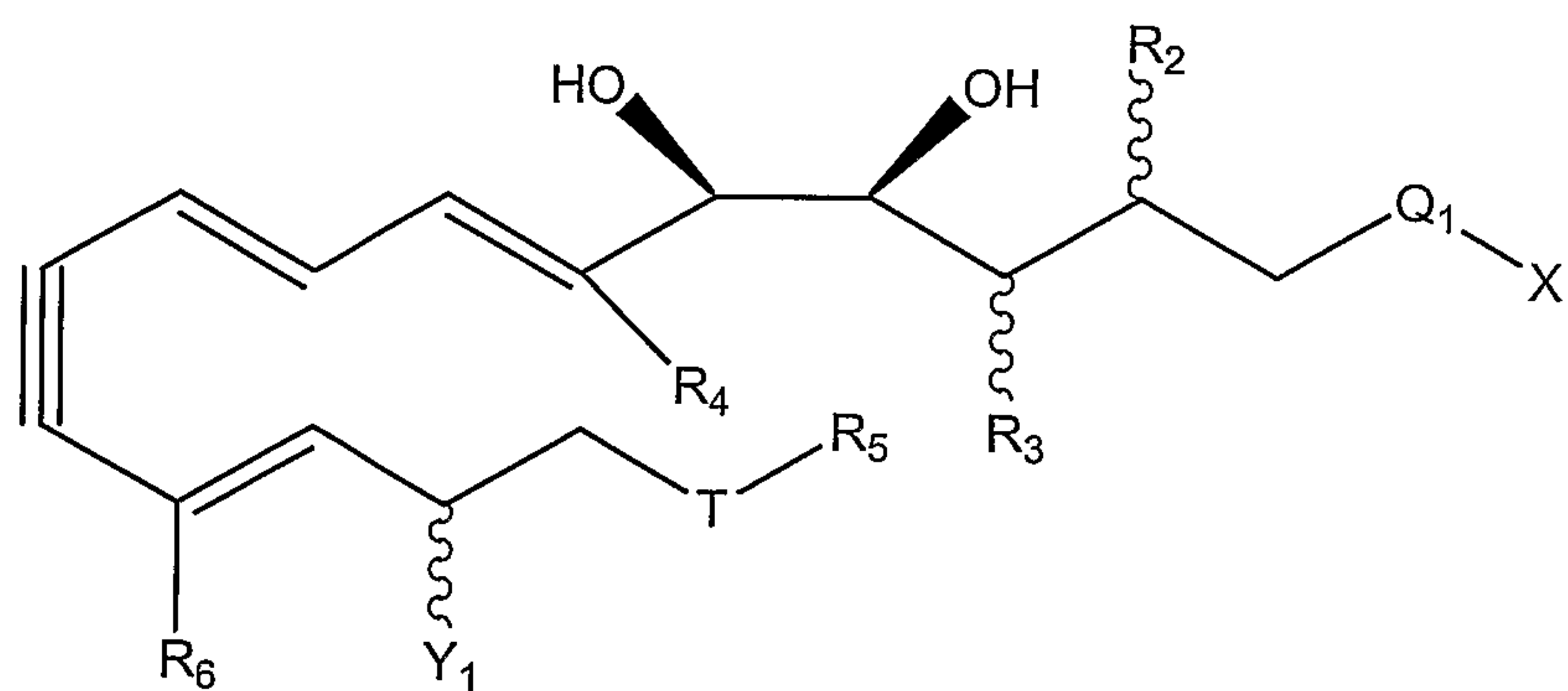


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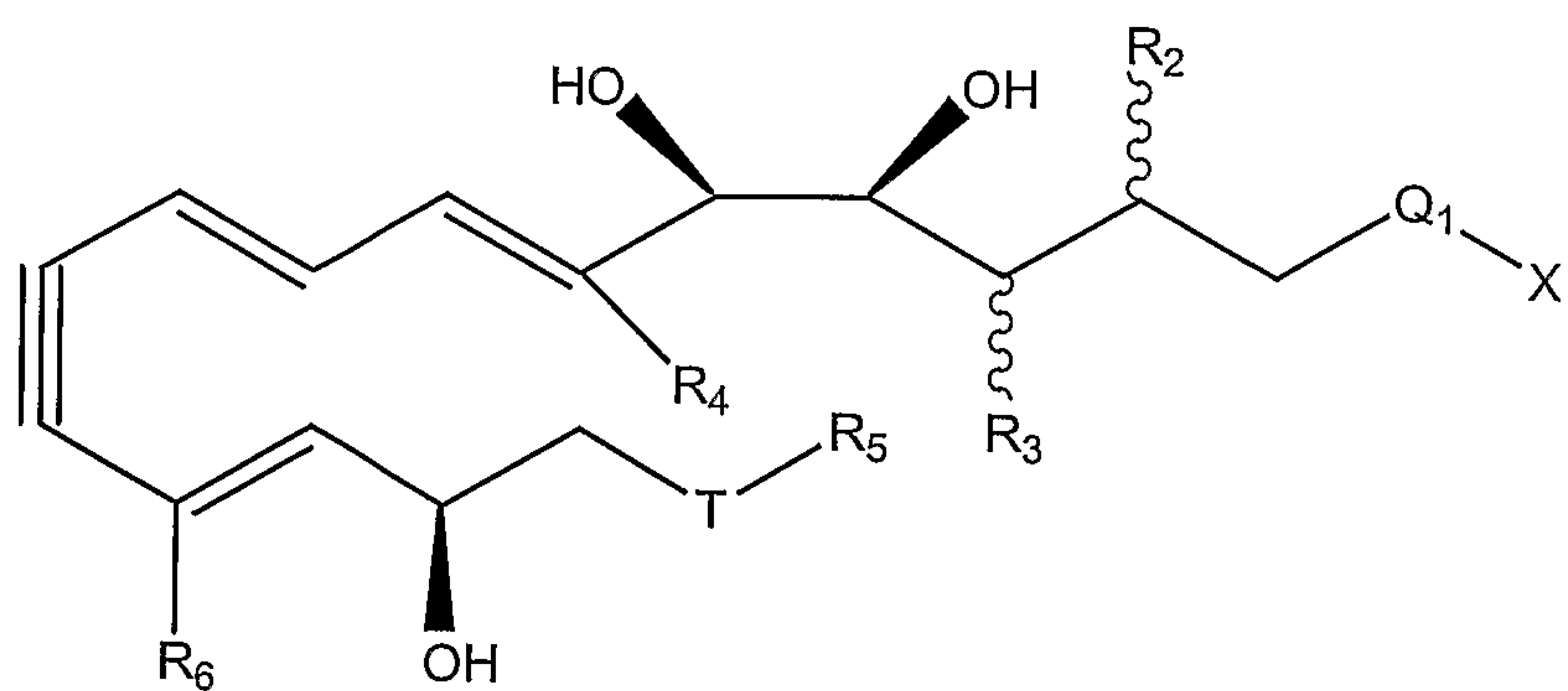
[015] Formula (VI)



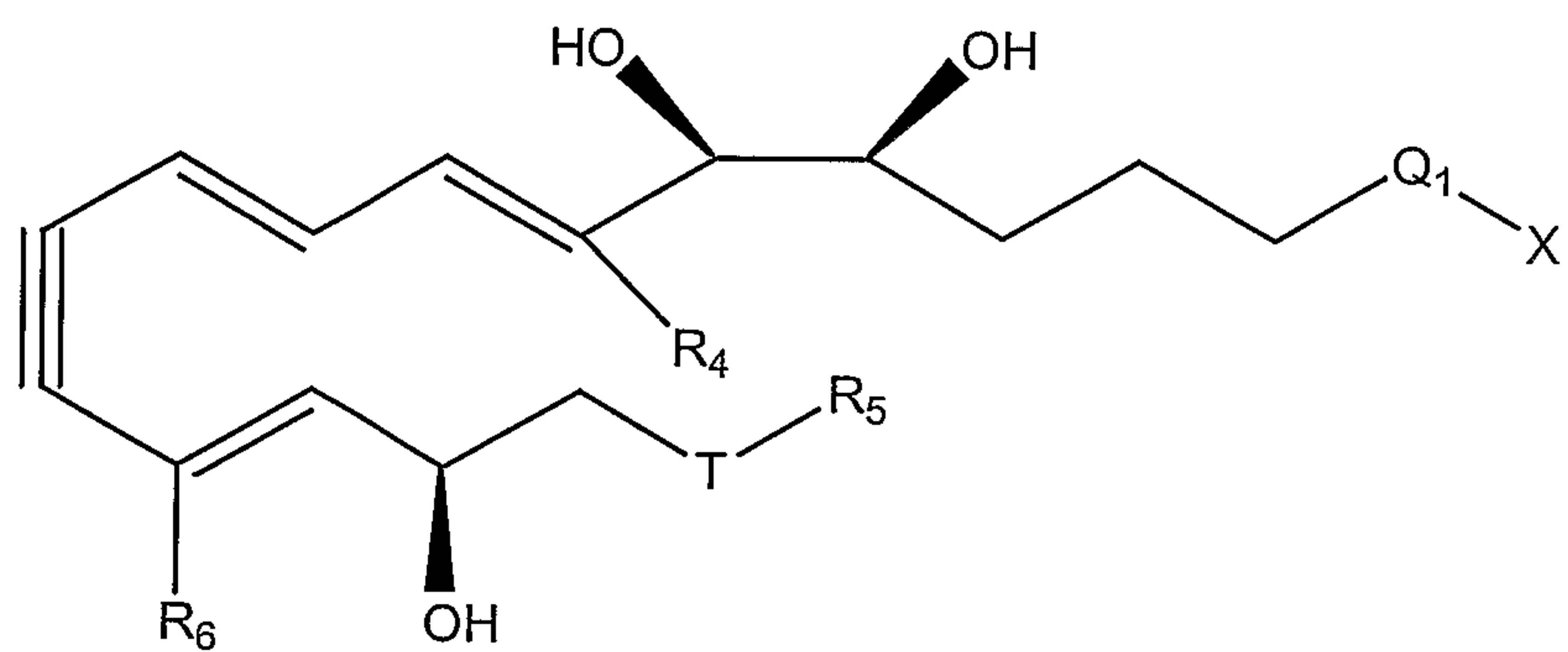
10 [016] Formula (VII)



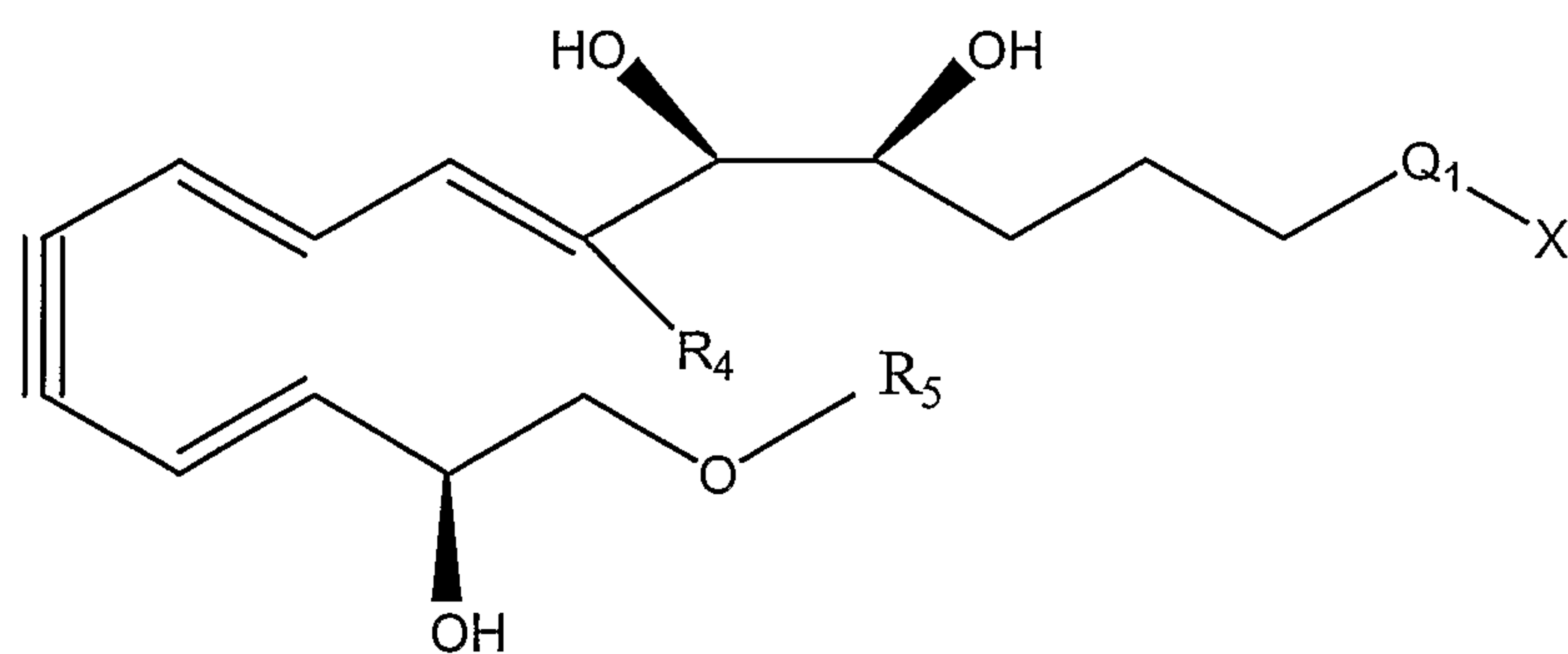
[017] Formula (VIII)



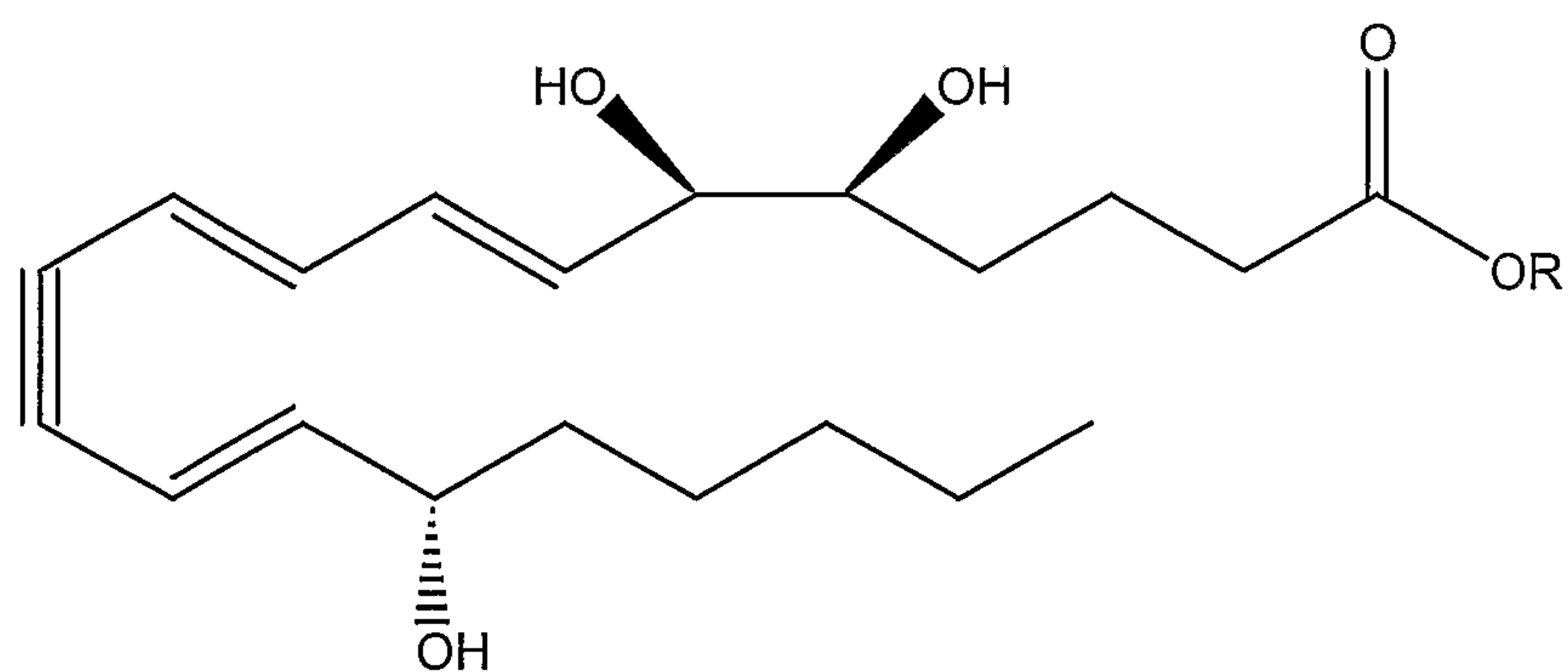
5 [018] Formula (IX)



[019] Formula (X)



[020] Formula (XI)



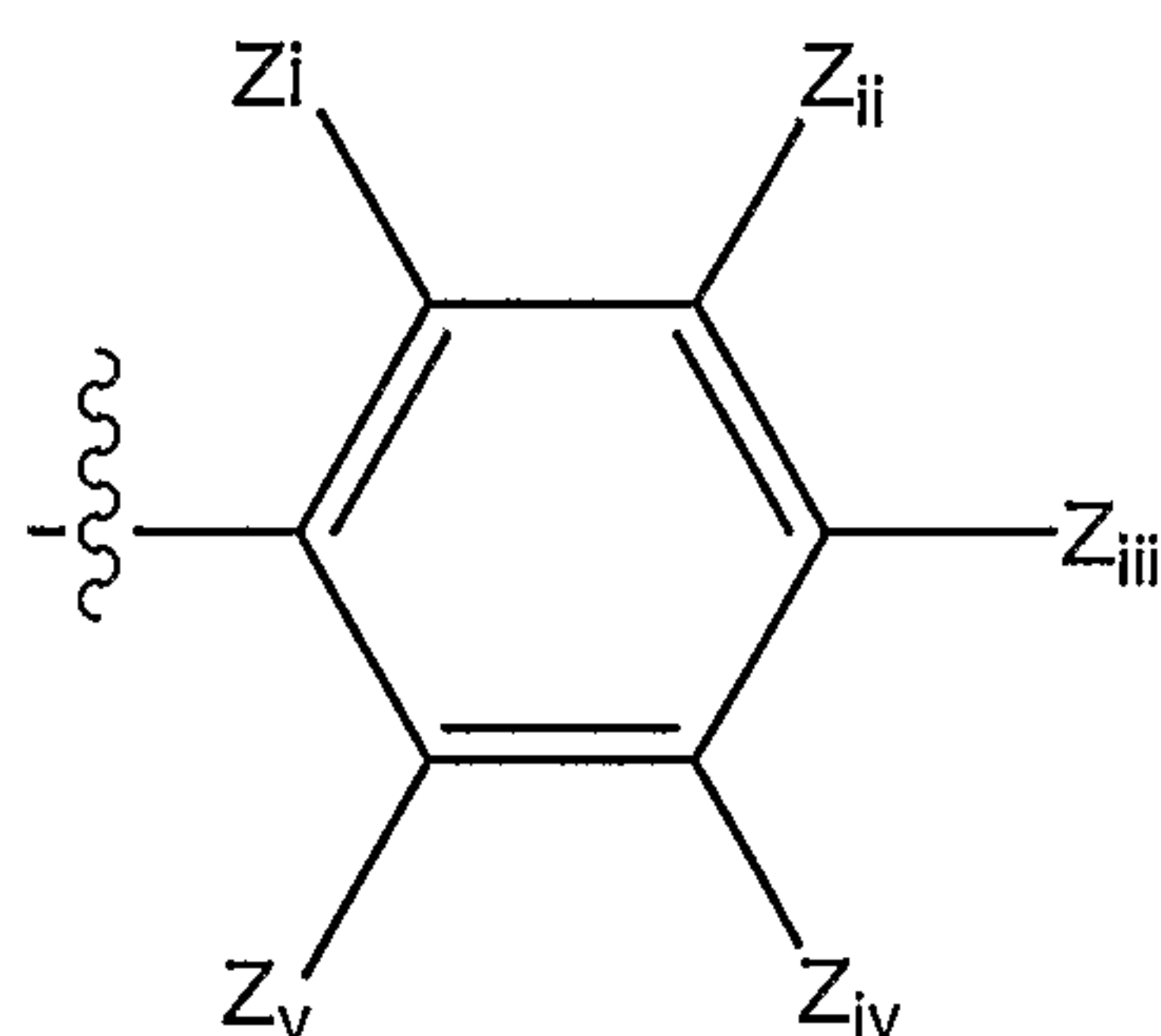
5

wherein X, if present, is R₁, OR₁, or SR₁;

wherein R₁, if present, is

- 10
- (i) a hydrogen atom;
 - (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
 - (iii) a cycloalkyl of 3 to 10 carbon atoms;
 - (iv) an aralkyl of 7 to 12 carbon atoms;
 - 15 (v) phenyl;

(vi) substituted phenyl



5 wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , –CN, –C(=O)– R_1 , – SO_3H , a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

(vii) a detectable label molecule; or

10 (viii) a straight or branched chain alkenyl of 2 to 8
 carbon atoms, inclusive;

wherein Q_1 , if present, is (C=O), SO_2 or (CN), provided when Q_1 is
 CN, then X is absent;

15 wherein Q_3 , if present, and Q_4 , if present, are each independently O, S
 or NH;

wherein one of R_2 , if present, and R_3 , if present, is a hydrogen atom and
 the other is

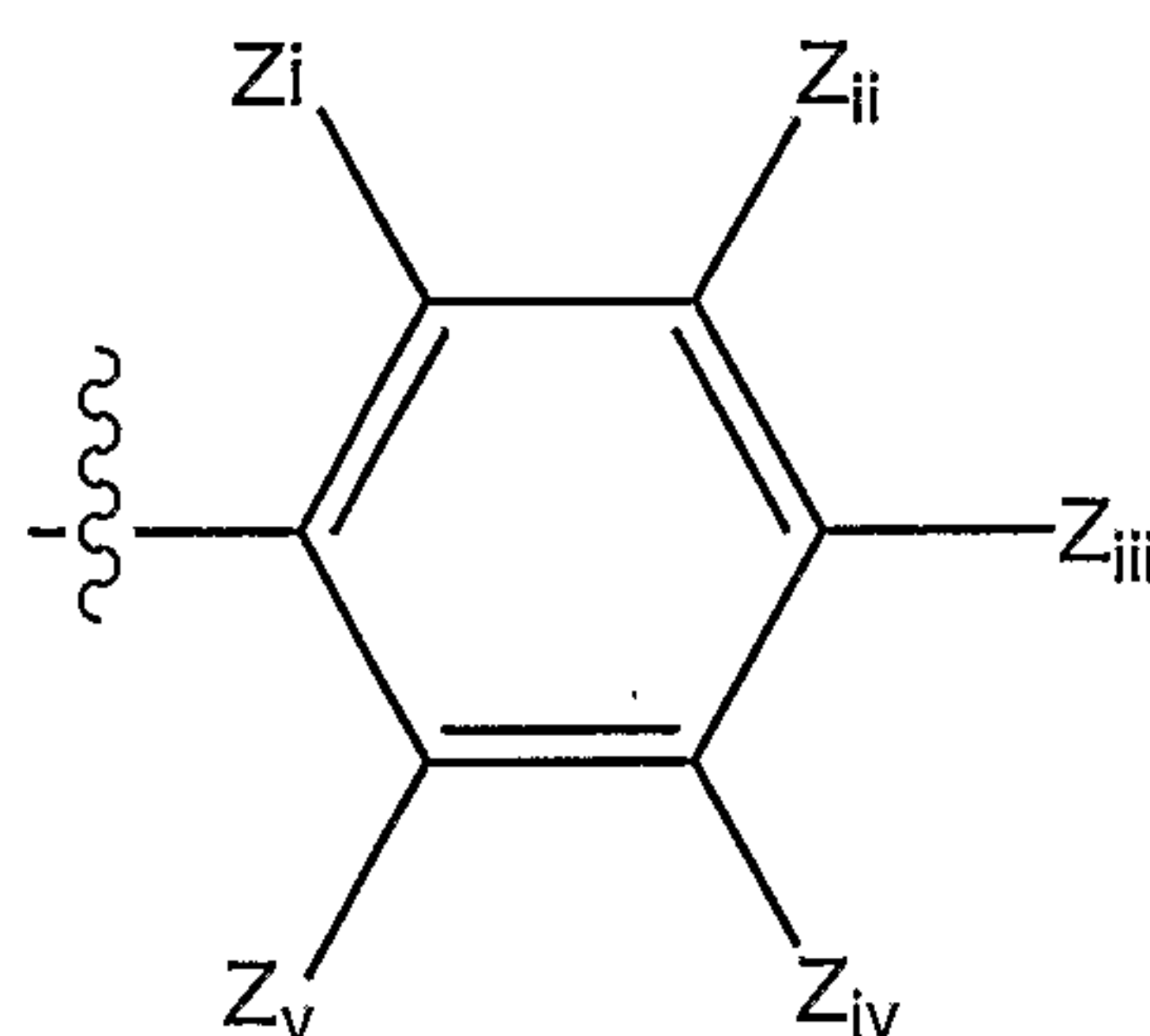
- 5
- (a) H;
- (b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;
- (d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be straight chain or branched; or
- (e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene of 0 to 6 carbon atoms, inclusive, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 , if present, is

- 15
- (a) H;
- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 , if present, is

20



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from —
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, —
 5 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group;

wherein Y_1 , if present, is $-\text{OH}$, methyl, $-\text{SH}$, an alkyl of 2 to 4 carbon
 10 atoms, inclusive, straight chain or branched, an alkoxy of 1 to 4 carbon
 atoms, inclusive, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is
 cyano, nitro or a halogen;

wherein R_6 , if present, is

- 15
- (a) H ;
 - (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain
 or branched;

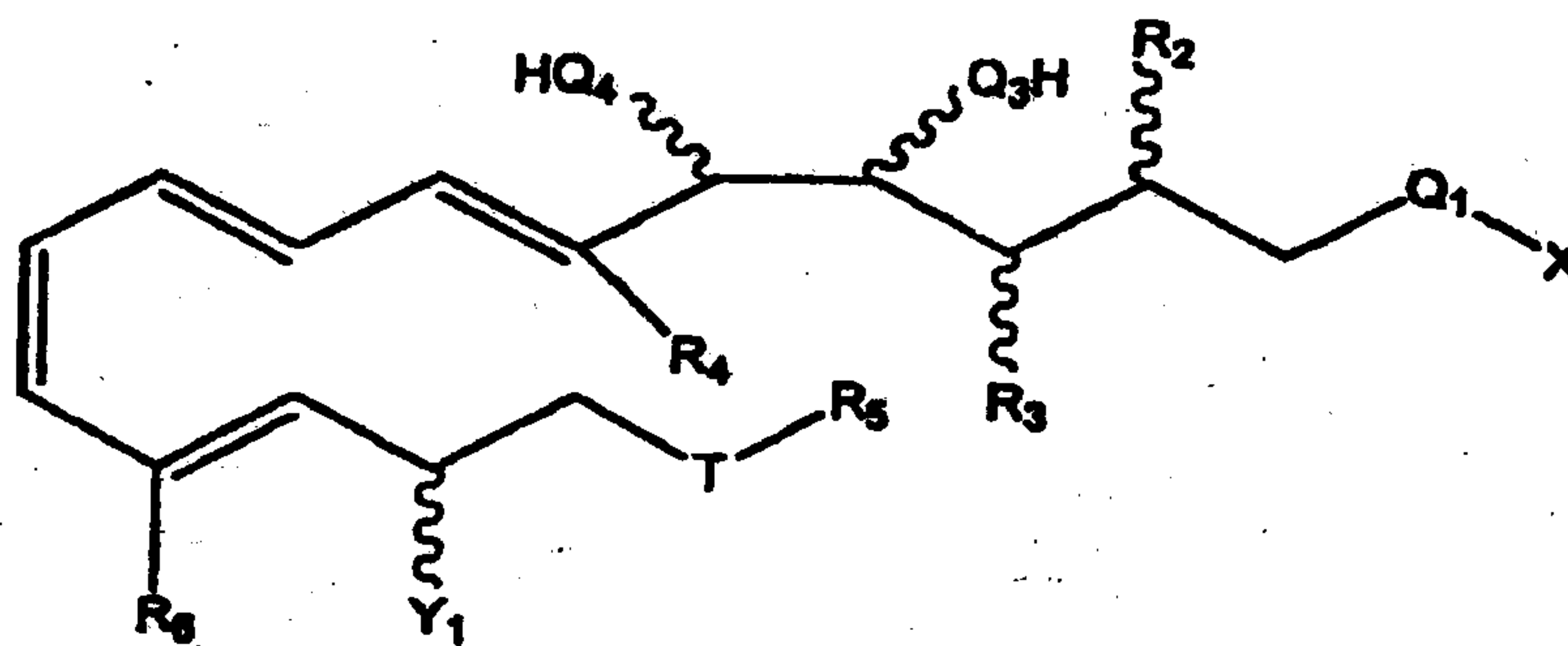
wherein T, if present, is O or S, and pharmaceutically acceptable salts thereof.

[021] In certain embodiments, the methods of the invention are performed *in vitro* or *in vivo*.

[022] In another aspect, the present invention is directed to a packaged pharmaceutical composition for treating the activity or conditions described herein in a subject. The packaged pharmaceutical composition includes a container holding a therapeutically effective amount of at least one lipoxin compound having one of the formulae described *infra* and instructions for using the lipoxin compound for treating the activity or condition in the subject.

10 In another aspect, the present invention provides use of a lipoxin analog for treating or preventing disorders of the respiratory tract or lung selected from the group consisting of chronic bronchitis, bronchiectasis, eosinophilic lung diseases, allergic bronchopulmonary aspergillosis, bronchiolitis, bronchiolitis obliterans, bronchiolitis obliterans with organizing pneumonia, eosinophilic granuloma, Wegener's granulomatosis, sarcoidosis,

15 hypersensitivity pneumonitis, lung injury associated with parasite infection and chronic obstructive pulmonary disease (COPD), said lipoxin analog having the formula

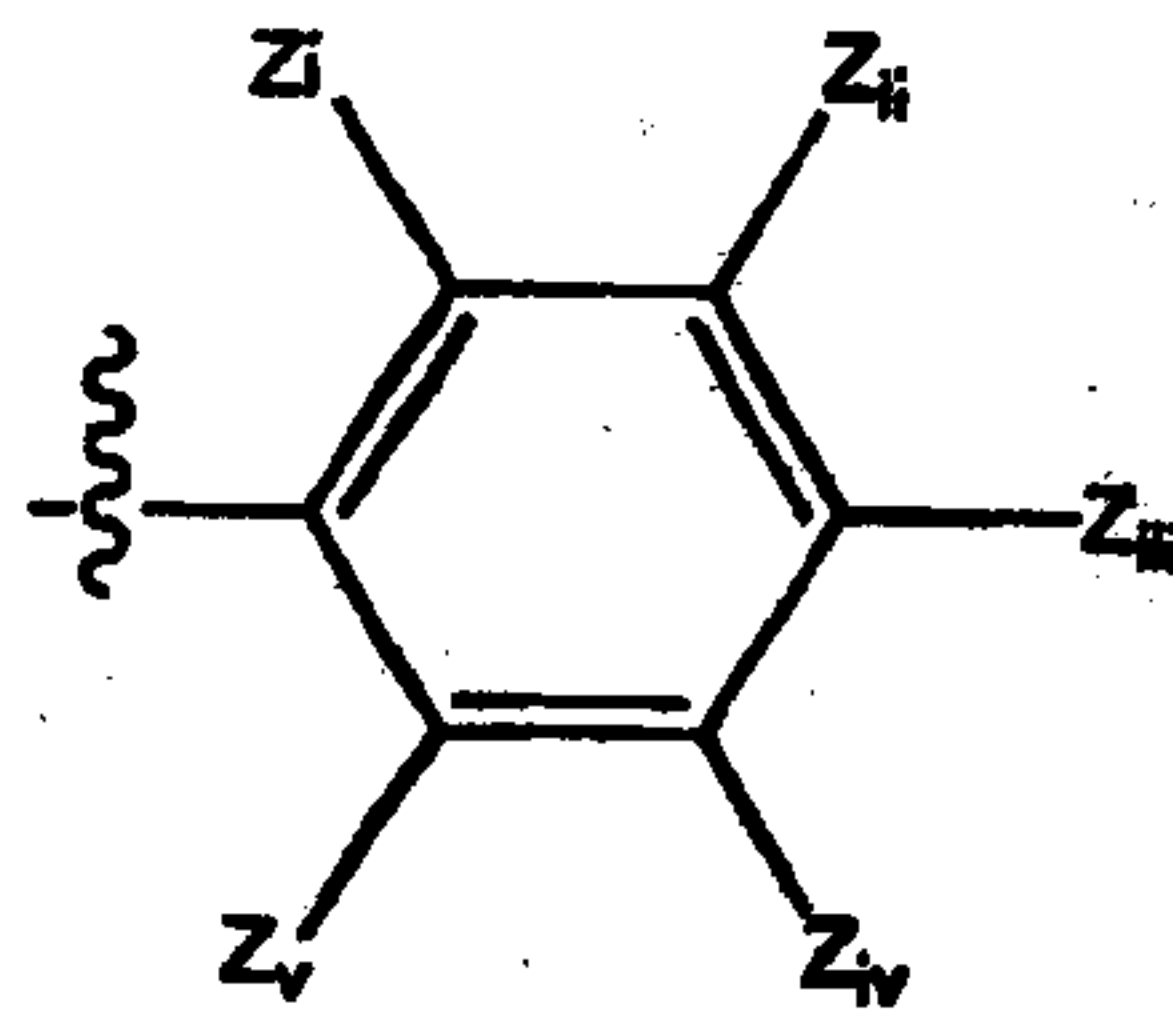


wherein X is R₁, OR₁, or SR₁;

20 wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;

- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$,

- 5 $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is absent; wherein Q_3

- 10 and Q_4 are each independently O , S or NH ;

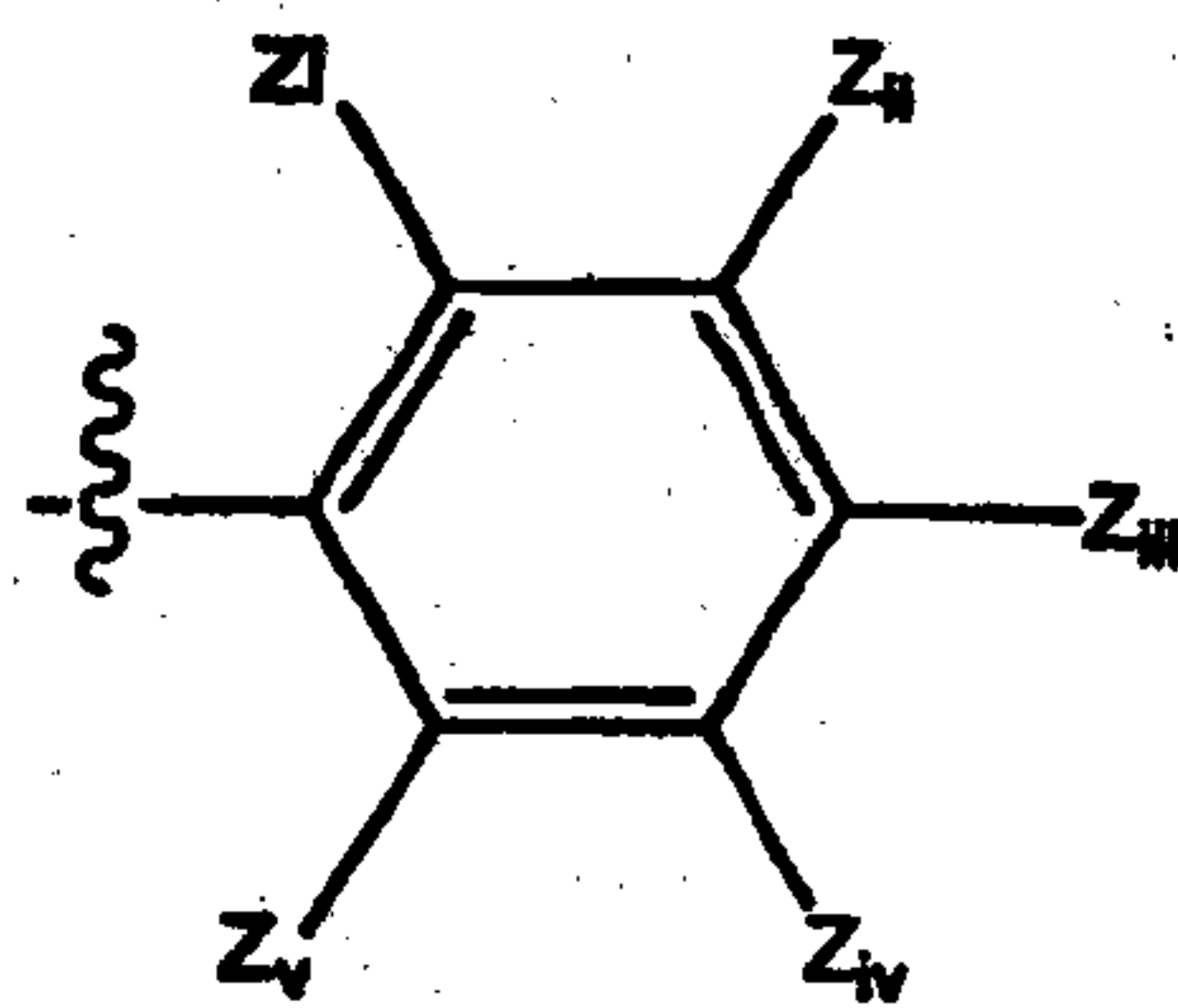
wherein one of R_2 and R_3 is a hydrogen atom and the other is

- (a) H ;
- (b) an alkyl of 1 to 8 carbon atoms, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms;
- 15 (d) an alkenyl of 2 to 8 carbon atoms, which may be straight chain or branched;
or
- (e) $R_a Q_2 R_b$ wherein Q_2 is $-\text{O}-$ or $-\text{S}-$; wherein R_a is alkylene of 0 to 6 carbon atoms, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, which may be straight chain or branched, provided when
20 R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

- (a) H ;
- (b) an alkyl of 1 to 6 carbon atoms, which may be a straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

wherein Y_1 is $-\text{OH}$, methyl, $-\text{SH}$, an alkyl of 2 to 4 carbon atoms, straight chain or branched, an alkoxy of 1 to 4 carbon atoms, or CH_2Z_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano, nitro or a halogen;

wherein R_6 is

- (a) H ;
- (b) an alkyl from 1 to 4 carbon atoms, straight chain or branched;

wherein T is O or S , and pharmaceutically acceptable salts thereof; with the proviso that the lipoxin analog is not 16-phenoxy lipoxin A_4 .

BRIEF DESCRIPTION OF THE DRAWINGS

[023] The invention will be more fully understood from the following detailed description taken in conjunction with the accompanying drawings, in which:

[024] Figure 1. Inhibition of airway hyper-responsiveness with LXa. OVA sensitized mice were treated with LXa (μg) (o) or vehicle ($\bullet$) prior to OVA aerosol challenge. Airway reactivity was determined by methacholine-dependent change in lung resistance (a) and calculation of ED_{200} (b). Results are expressed as mean $\pm$ SEM ($n \geq 7$ in (a) and $n \geq 6$ (b)). $*P < 0.05$ by Student's t-test compared to control animals.

[025] Figure 2. Lung histopathology from LXa treated mice. Mice were sensitized and aerosol challenged with OVA in the absence (left column) or presence (right column) of LXa (10 μg). Representative ($n=3$) lung tissue sections (magnifications: x40 (a,b), x100 (d),

x200 (*c,e*) and x400 (*f*) were obtained from formalin-fixed, paraffin-embedded lung tissue, prepared and stained with hematoxylin and eosin. Arrows denote eosinophils; *br*, bronchus; *v*, vessel.

5 [026] Figure 3. LXa selectively inhibits airway leukocyte infiltration and inflammatory mediators. Bronchoalveolar lavage fluids were obtained from OVA sensitized and challenged mice. Leukocytes in bronchoalveolar lavage fluid were enumerated and identified after Wright-Giemsa stain (*a*). Mediator

profile of (b) specific cytokines (IL-5, IL-12, IL-13, eotaxin and TNF α), and lipid mediators, including PGE₂ (c), LTB₄ and cysLT's (d) were determined by ELISA's in materials from control animals and those receiving LXa (10 μ g, i.v.). Results are expressed as mean $\pm$ SEM ($n \geq 6$, $d \geq 2$). * $P < 0.05$ by Student's t-test compared to control animals.

[027] Figure 4. *In situ* hybridization of murine ALX in lung. Murine LXA₄ receptor expression was detected in perivascular leukocyte-rich infiltrates (upper panels) and airway epithelia (lower panels) in OVA sensitized and aerosol challenged (OVA/OVA) mouse lung (representative bright-field images, magnification x400, $n=3$). Experimental (PBS/PBS) and sense probe controls are shown for comparison.

[028] Figure 5. Expression of human ALX in transgenic mice decreases pulmonary inflammation. a, Lung histopathology from human ALX transgenic mice. Non-transgenic (left column) and human ALX-tg mice (right column) were sensitized and aerosol challenged with OVA. Representative ($n=6$) lung tissue sections (magnifications: x100 (a,b) and x200 (c,d)) were obtained from formalin-fixed, paraffin-embedded lung tissue, prepared and stained with hematoxylin and eosin. br, bronchus; v, vessel. b-e, BAL fluids were obtained, leukocytes enumerated and identified after Wright-Giemsa stain. Percent inhibition in cell number (b) and changes in mediator profile of specific cytokines and lipid mediators (c) were determined. Serum total IgE levels were determined by immunoassay in samples from mice sensitized/challenged with either OVA or buffer (PBS) (d). Airway reactivity was determined by methacholine (100 mg/ml)-dependent change in Penh (e). Results are expressed as mean $\pm$ SEM ($n=4$, $d \geq 2$). * $P < 0.05$ by Student's t-test compared to control animals.

[029] Figure 6. Expression of human ALX in transgenic mice prevents eosinophil trafficking. LTB₄ (1 μ g) and PGE₂ (1 μ g) were topically applied (13-16h) to mouse ear skin and EPO activity in tissue biopsies was determined by bromination of HPA (see *Methods*). Values represent the mean $\pm$ SEM for pmoles Br-HPA per μ g total protein for Non-tg littermates ($n=6$) or BLT-tg ($n=2$)

mice in the absence or presence of topical LXA (10 µg per ear) and 2 separate ALX-tg lines ($n=4$). * $P < 0.05$ by Student's T-test compared to littermate controls.

5 [030] Figure 7. Depicts regulation of allergic airway inflammation by 15-epi,16-para-fluoro-phenoxy-LXA₄-methyl ester in comparison to a commercially available drug (Montelukast) used to treat chronic asthma. This graph demonstrates that the LXA₄ methyl ester is statistically more effective than the Montelukast in terms of lymphocyte reduction.

DETAILED DESCRIPTION

10 [031] The features and other details of the invention will now be more particularly described and pointed out in the claims. It will be understood that the particular embodiments of the invention are shown by way of illustration and not as limitations of the invention. The principle features of this invention can be employed in various embodiments without departing from the scope of the
15 invention.

[032] The prevalence of asthma continues to increase and its optimal treatment remains a challenge. The present invention pertains to methods that involve the actions of lipoxin A₄ and its leukocyte receptor in pulmonary inflammation using a murine model of asthma. Allergen challenge initiated airway biosynthesis of
20 lipoxin A₄ and increased expression of its receptor. Administration of a stable analogs of lipoxin A₄ blocked both airway hyper-responsiveness and pulmonary inflammation, as evidenced by decreased recruitment of leukocytes as well as reduced mediator levels, including interleukin-5, interleukin-13, eotaxin, prostanoids and cysteinyl leukotrienes. Moreover, transgenic expression of
25 human lipoxin A₄ receptors in murine leukocytes led to significant inhibition of both pulmonary inflammation and eicosanoid-initiated eosinophil tissue infiltration. Inhibition of airway hyper-responsiveness and allergic airway inflammation with a lipoxin A₄ stable analogs highlights a unique counter-regulatory profile for the lipoxin A₄ system and its leukocyte receptor in airway

responses and suggest that lipoxin and related pathways present novel multi-pronged therapeutic approaches for consideration in human asthma.

[033] Lipoxin A₄ (LXA₄) is rapidly generated from arachidonic acid during inflammatory responses, and, unlike leukotrienes (LT), LXA₄ and its stable
5 analogs carry anti-inflammatory properties that promote resolution of acute exudative inflammation. The present invention demonstrates that LXA₄ also regulates airway inflammation and bronchial reactivity, characteristic features of asthma. High levels of both cysteinyl LT and prostaglandin E₂ (PGE₂) in bronchoalveolar lavage (BAL) fluids from ovalbumin (OVA) sensitized and
10 aerosol were found in challenged mice. LXA₄ was also identified in these same samples, yet in concentrations 1-2 log orders less (15.01 +/- 3.26 pg/ml, mean +/- SEM, n=5) than LT and PG, respectively. When an LXA₄ stable analog (including its 11,12 acetylenic version) was given 1-2 h prior to OVA aerosol challenge, a marked reduction in tissue eosinophils, neutrophils, lymphocytes and
15 vascular injury was observed. The LX analogs also led to decreased concentrations of IL-5, IL-13 and eotaxin, but not IL-12 or TNF in OVA sensitized and challenged animals. Lipid mediator formation was also selectively down-regulated by LX analog administration as levels of PGE₂ and cysteinyl LT's, but not LTB₄ were decreased. In addition to inflammatory responses,
20 administration of the LX analog significantly inhibited bronchoconstriction in response to methacholine with an ED₂₀₀ that approximated that of unchallenged mice. Because LXA₄ can interact with cysLT1 receptors, it was determined that LX administration alone would promote airway hyperreactivity *in vivo*, and no significant differences were present in ED₂₀₀ compared to control mice. Since
25 LXA₄ displayed potent inhibition of eosinophil tissue infiltration, it was next determined the impact of over-expression of its receptors (ALXR) on agonist-initiated eosinophil accumulation. Thirteen hours after application of LTB₄ and PGE₂ to mouse ear skin, punch biopsies were obtained for histology and quantitation of eosinophil peroxidase activity. Compared to littermate control
30 animals, increased expression of ALXR profoundly inhibited eosinophil infiltration in two separate transgenic lines. Together, these findings indicate that

both increased LXA₄ bioavailability via utilization of a stable analog and increased ALXR expression can mediate potent inhibition of airway inflammation and eosinophil responses *in vivo*. Hence, this invention reports a novel use for lipoxins, aspirin-triggered lipoxins and their structural analogs in
5 airway inflammation, asthma and related disorders.

Eicosanoid formation during antigen-initiated airway inflammation.

[034] After systemic sensitization to ovalbumin (OVA, 10 µg i.p.), male Balb/c mice (5-7 weeks) were exposed to aerosolized OVA (6%, 25 min) on four
10 successive days. 24 hours after the last aerosol, bronchial responsiveness to intravenous methacholine was determined (as in refs. 14, 15), bronchoalveolar lavage was performed and tissues were harvested for microscopy. Animals so treated demonstrate both airway hyper-responsiveness to methacholine and inflammation (*vide infra*), including leukocytic infiltration and elaboration of
15 cytokines, chemokines and bioactive lipids. After allergen sensitization and aerosol challenge, high levels of both CysLT (139.0 +/- 27.3 pg CysLT/ml, mean +/- SEM, n=9) and PGE₂ (1117.7 +/- 103.8 pg PGE₂/ml, mean +/- SEM, n=5) in bronchoalveolar lavage (BAL) (Table 1) were identified. In these same BAL fluids, LXA₄ was also present (15.0 +/- 3.3 pg LXA₄/ml, mean +/- SEM, n=5) at
20 levels similar to LTB₄ (6.4 ± 2.3 pg LTB₄/ml, mean +/- SEM, n=9), but in 10- to 100-fold lower concentrations than CysLT's and PGE₂, respectively (Table 1). LXA₄ was not detected in cell-free BAL supernatants from non-immunized mice.

Lipoxin A₄ prevents airway hyper-responsiveness to methacholine.

25 [035] To determine if LXA₄ would protect mice from OVA-induced airway hyper-responsiveness, 10 µg/mouse (i.v.) of a LXA₄ analog (LXa) was administered that resists metabolic inactivation and blocks neutrophil accumulation and exudate formation in murine dorsal air pouches^{7, 16}. When given at least 60 min prior to OVA challenge, LXa significantly inhibited

bronchoconstriction in response to methacholine in a dose-dependent manner (Figures 1a & b). The ED₂₀₀ to methacholine for mice given LXa (10 µg) approximated the responses of control mice that had been sensitized but not challenged with OVA (Figure 1b). Neither deleterious physiologic nor behavioral effects were evident in LXa treated mice. It was also determined that LXa administration alone would promote airway hyper-reactivity *in vivo*, and no significant differences in ED₂₀₀ compared to control mice (Figure 1b) were found.

Lipoxin A₄ inhibits key parameters of allergic pulmonary inflammation.

[036] In addition to dampening airway hyper-responsiveness in OVA-allergic mice, administration of LXa significantly reduced leukocyte infiltration, in particular, tissue eosinophils and lymphocytes as well as vascular injury (Figure 2). In BAL, total leukocytes, eosinophils and lymphocytes were sharply reduced in a dose-dependent fashion (Figure 3a). LXa also led to decreased T_H2 cytokines IL-5, IL-13 as well as eotaxin in BAL fluids from OVA sensitized and challenged animals (Figure 3b). Inhibition appeared selective, since levels of IL-12 and TNF were not similarly reduced when determined in the same samples of BAL fluid (Figure 3b). LXa also regulated levels of the lipid mediators, as both PGE₂ and CysLT's, but not LTB₄ were decreased in these mice (Figures 3c & d). These results indicate that administration of LXA₄ mimetics can significantly inhibit the generation of allergic pulmonary inflammation, including leukocyte infiltration and formation of specific mediators of interest in airway pathophysiology, including key cytokines and eicosanoids.

Expression of human LXA₄ receptors dampens pulmonary inflammation.

[037] Allergen sensitization and challenge with OVA increased LXA₄ receptor (ALX) expression in infiltrating leukocytes and airway epithelial cells observed with *in situ* hybridization (Figure 4). To assess whether OVA allergic pulmonary inflammation and airway reactivity were amenable to regulation by ALX, the responses of transgenic mice expressing human ALX using the CD11b promoter

on their leukocytes (see *Methods*) was assessed. After OVA sensitization and aerosol challenge, human ALX transgenic animals (ALX-tg) displayed reduced airway and vascular injury, when compared to their age and gender-matched littermate controls (Non-tg). A clear decrease in leukocytic infiltrates was
5 observed in peribronchial and perivascular spaces (Figure 5a). BAL from allergen challenged ALX-tg mice also had reduced numbers of total leukocytes, eosinophils and lymphocytes, with 63%, 68% and 85% inhibition, respectively (Figure 5b). Levels of several key pro-inflammatory peptide and lipid mediators were also decreased in ALX-tg mice, including IL-13 (77% inhibition), IL-5
10 (71% inhibition) and CysLT (74% inhibition) (Figure 5c). In addition, allergen sensitization in the ALX-tg animals was blunted, as monitored by total serum IgE levels (Figure 5d). Despite inhibition of allergic inflammation in the OVA sensitized and challenged ALX-tg mice, significant differences in airway hyper-responsiveness were not observed (Figure 5e) when compared to age and gender-matched littermates (peak Penh was 8.37 ± 1.89 (ALX-tg) and 8.25 ± 1.90 (Non-tg), mean $\pm$ S.E. for n=6).

Expression of human LXA₄ receptors inhibits murine eosinophil tissue infiltration.

20 [038] With levels of both tissue leukocytes and pro-inflammatory mediators reduced in allergic ALX-tg mouse lungs (Figure 5), the direct impact of human ALX expression on murine eosinophil recruitment *in vivo* was examined. After topical application of LTB₄ (1 μ g) and PGE₂ (1 μ g) to mouse ear skin, there was a significant increase in eosinophil tissue infiltration as quantified by eosinophil
25 peroxidase (EPO) activity present within 6 mm skin punch biopsies (Figure 6). Compared to littermate non-tg controls, expression of human ALX markedly reduced the number of eosinophils recruited in ear skin tissues from two separate murine transgenic lines (Figure 6). Of note, endogenous LXA₄ was recovered from inflamed ears and was without significantly different levels between ALX-tg and Non-tg animals (16.0 $\pm$ 1.5 and 11.5 $\pm$ 1.9 pg/mg protein, respectively).
30

Increased expression of human LTB₄ receptors on leukocytes (also using CD11b promoter) led to a similar amplitude of eosinophil infiltration in this model as Non-tg littermate controls. Topical administration of as little LXA as 10 µg to mouse ears inhibited eicosanoid-stimulated skin EPO activity in both Non-tg mice as well as those with increased expression of LTB₄ receptors (Figure 6). Together, these findings indicate that LXA₄-ALX interactions can mediate potent inhibition of eosinophil responses *in vivo*.

Sensitization and challenge protocols

10 [039] Five to seven week old male FVB (Charles River Laboratories, Wilmington, Massachusetts) mice were housed in isolation cages under viral antibody-free conditions. After Harvard Medical Area IRB approval (Protocol #02570), mice were sensitized with intraperitoneal injections of ovalbumin (OVA) (Grade III; Sigma Chemical Co., St. Louis, Missouri) (10 mg) plus 1 mg
15 aluminum hydroxide (ALUM) (J.T. Baker Chemical Co.; Phillipsburg, New Jersey) as adjuvant in 0.2 ml PBS on days 0 and 7. On days 14, 15, 16 and 17, the mice received an analog of LXA₄ (15-epi,16-para-fluoro-phenoxy-LXA₄-methyl ester) (10 mg, 21.7 nmoles), montelukast (13 mg, 21.7 nmoles) or PBS with 1.6 mM CaCl₂ and 1.6 mM MgCl₂ (0.1 ml) by intravenous injection at least
20 1 h prior to an aerosol challenge containing either PBS or 6% OVA for 25 min/day. On day 18, 24 h after the last aerosol challenge, bilateral bronchoalveolar lavage (BAL) (2 aliquots of 1 ml PBS plus 0.6 mM EDTA) was performed. BAL fluids were centrifuged (2000 g, 10 min) and cells were resuspended in HBSS, enumerated by hemocytometer, and concentrated onto
25 microscope slides by cytocentrifuge (STATspin) (265 g). Cells were stained with a Wright-Giemsa stain (Sigma Chemical Co.) to determine leukocyte differentials (after counting > 200 cells).

[040] Figure 7. Depicts regulation of allergic airway inflammation by 15-epi,16-para-fluoro-phenoxy-LXA₄-methyl ester in comparison to a commercially
30 available drug (Montelukast) used to treat chronic asthma. Montelukast is a leukotriene antagonist and is the subject of US Patent No. 5,565,473, the contents

of which are incorporated herein in their entirety. Figure 7 provides that the LXA₄ methyl ester is statistically more effective than Montelukast in terms of lymphocyte reduction.

[041] As a consequence of these studies, LXA₄ analogs inhibit both airway
5 hyper-responsiveness and inflammation in response to allergen sensitization and aerosol challenges in murine models of asthma. These models correlate with and are acceptable models for validation in humans and other mammals.

[042] While the results of recent clinical trials in human asthma have challenged
10 long held views on the relationship between airway inflammation and bronchial hyper-responsiveness^{20,21}, the present invention provides that LXa prevented both of these key asthma phenotypes in an experimental model of asthma, displaying a multi-pronged impact *in vivo* that in summation appears to give beneficial airway responses.

[043] When this longer acting stable analog mimetic of endogenous LXA₄ was
15 administered prior to OVA aerosol, airway hyper-responsiveness to methacholine as well as several measures of inflammation were markedly reduced. Indeed, pharmacologic levels of LXA₄ had direct effects on airway smooth muscle responses to challenge since native LXA₄ itself given to human asthmatics inhibits LTC₄-stimulated airway hyper-responsiveness²² as well as blocks LTD₄-
20 initiated constriction of airway smooth muscle *in vitro*²³. In addition to LXA₄ analog actions on its cognate high affinity receptor (ALX), LXA₄ and the present analog (constructed on the aspirin-triggered 15-epimer LXA₄ biotemplate; denoted here as LXa) compete for binding with high affinity at LTD₄ recognition sites and compete at recombinant cysLT1 receptors expressed in Chinese hamster
25 ovary cells *in vitro*¹². The LX-mediated decrements in allergic mediators documented are also be responsible, in part, for the observed protection from the development of airway hyper-responsiveness to methacholine. In this regard, recombinant IL-4 and IL-13 induce airway hyper-responsiveness within hours without inflammatory cell recruitment or mucus production²⁴, IL-13 decreases
30 human airway smooth muscle β -adrenergic responsiveness *in vitro*²⁵, and IL-5 and eotaxin lead to increased production of CysLT^{26, 27}, which are potent

bronchoconstrictors²⁸. Taken together, these findings show that endogenous LX's produced within the local microenvironment might temporally regulate and reduce airway hyper-responsiveness via multiple sites of action *in vivo* to counterregulate pathways currently considered important in the genesis of this
5 asthma phenotype, namely via inhibition of IL-5, IL-13 and CysLT-mediated actions on leukocytes, epithelia and smooth muscle.

[044] Pulmonary eosinophilia was also sharply reduced in mice given LXA. Eosinophil recruitment to the lung in asthma is primarily a consequence of T_H2 lymphocyte activation¹, which was reduced by LXA as evidenced by lower levels
10 of T_H2 cytokines in BAL fluid and decreased number of lymphocytes in both BAL fluid and lung tissue. LXA₄ may inhibit lymphocyte recruitment by competition for ALX binding with the urokinase plasminogen activator receptor, a potent lymphocyte chemoattractant²⁹. In addition, LXA₄ directly inhibits eosinophil chemotaxis³⁰, and several of its stable analog mimetics block eotaxin
15 formation *in vivo* in parasite animal models ³¹, promoting resolution of allergen-mediated pleural inflammation³². While endogenous LX production paralyzes splenic dendritic cell responsiveness to IL-12 and IL-12 production in the setting of microbial immunity³³, LXA administration during allergen challenge did not significantly alter IL-12 levels in BAL fluids, suggesting distinct sites of LXA₄
20 action in allergic inflammation down-stream from dendritic cells. In aggregate, the present invention provides potent inhibition by LXA and lipoxin analogs for both T_H2 lymphocyte and eosinophil recruitment *in vivo*, processes that characterize asthma pathobiology. Moreover, the capacity for LXA and lipoxin analogs to inhibit these inflammatory responses raises the likelihood that
25 endogenous LX production serves in health as a pivotal regulatory event in airway and allergic inflammation.

[045] LX's were generated endogenously during allergen induced airway inflammation. The amounts of LXA₄ recovered were similar to the levels of LTB₄, yet amounts of both of these eicosanoids were substantially lower than
30 either CysLT's or PGE₂ within BAL of allergen challenged mice. Spatial and temporal analyses during an acute inflammatory response indicate that maximal

LX levels recovered are delayed in onset compared to either LT's or PG's and concurrent with resolution, namely reduction of the exudate rather than initiation of inflammation⁹. Thus, the local levels of endogenously generated LX's after allergen challenge determined in this model reflect not only the interval of sample acquisition, but also the cell origins of the mediators present in BAL and other spatial relationships to cellular generators within the lung.

[046] LXA₄ interacts with ALX to mediate leukocyte-selective effects that promote resolution^{7, 8}. In the present invention, allergic pulmonary inflammation was dramatically inhibited by expression of human ALX on murine leukocytes leading to marked decreases in serum total IgE, leukocyte tissue infiltration, and cytokine and lipid mediator formation. Of note, parallel decreases in bronchial hyper-responsiveness to methacholine were not observed in the ALX-tg mice. An uncoupling or dissociation of airway reactivity from inflammation per se has been uncovered in recent clinical trials^{20, 21} and may have resulted from restricted expression of the hALX transgene to cells expressing the CD11b promoter, namely, leukocytes and not resident tissues of the airways. Of interest, ALX-tg and Non-tg mice given LXa had similar reductions in IL-13 and CysLT. These findings suggest that the mechanisms for LXa-mediated inhibition of bronchial hyper-responsiveness following i.v. administration were distinct and likely secondary to the recently established direct interactions between LXa and recombinant CysLT1 receptors¹². These present invention also provides that an increased ligand, as given pharmacologically via stable mimetic (LXa), or increased receptor (ALX) expression can prevent allergic pulmonary inflammation. Elucidation of endogenous regulators of LX pathways further provide insight into its role in inflammatory lung disease. Along these lines, whole blood from human aspirin-intolerant asthmatic individuals gives reduced LX biosynthetic capacity relative to aspirin-tolerant asthmatic or healthy individuals³⁴, which it is believed accounts, in part, for a more protracted and severe clinical course in patients with aspirin-intolerant asthma.

[047] The present invention provides direct protective and regulatory roles for LX mimetics (lipoxin analogs of the invention) in airway hyper-responsiveness

and asthmatic inflammation. In light of their ability to inhibit both of these key asthma phenotypes *in vivo*, LX mimetics represent a new treatment and therapeutic approach for asthma. Rather than inhibiting the actions of a single class of airway mediators to control asthma, LX (lipoxins) and their stable
5 analogs promote resolution of inflammation via multiple mechanisms, including inhibition of leukocyte (eosinophil and lymphocyte) recruitment and activation; cytokine and chemokine production; and biosynthesis of pro-inflammatory lipid mediators, as demonstrated here, as well as by stimulating the non-phlogistic clearance of apoptotic leukocytes³⁵, blocking edema formation³², and inhibiting
10 neutrophil trafficking and functional responses¹⁶. Of note, LX stable analogs inhibit neutrophil responses *in vivo* with similar potency as corticosteroids¹³. Together, the present invention provides the profile of *in vivo* actions of LXa indicate that LX mimetics and related compounds could provide novel therapeutic approaches to the treatment of airway hyper-responsiveness and
15 pulmonary inflammation in select populations of human asthmatics.

[048] The present invention has surprisingly uncovered the ability for LX's and their analogs described herein to potently inhibit allergen-mediated pulmonary inflammation and airway hyper-responsiveness, establishing novel multi-pronged protective actions for these compounds. Moreover, these results indicate that the
20 LX system, including LXs and ALX, plays pivotal and previously unappreciated roles in regulating allergy and pulmonary inflammation of interest in asthma and related pulmonary disorders.

[049] The lipoxin analogs of the invention are useful in the treatment and prevention of airway inflammation, asthma and related disorders of the
25 respiratory tract and lung, such as chronic bronchitis, bronchiectasis, eosinophilic lung diseases (including parasitic infection, idiopathic eosinophilic pneumonias and Churg-Strauss vasculitis), allergic bronchopulmonary aspergillosis, allergic inflammation of the respiratory tract (including rhinitis and sinusitis), bronchiolitis, bronchiolitis obliterans, bronchiolitis obliterans with organizing
30 pneumonia, eosinophilic granuloma, Wegener's granulomatosis, sarcoidosis, hypersensitivity pneumonitis, idiopathic pulmonary fibrosis, pulmonary

manifestations of connective tissue diseases, acute or chronic lung injury, adult respiratory distress syndrome, plus several other non-infectious, inflammatory disorders of the lung.

[050] Additionally, lipoxin and the lipoxin analogs of the invention are useful in the inhibition and/or treatment and/or prevention of tissue injury, preservation of material for lung transplantation, lung injury associated with parasite infection(s), chronic obstructive pulmonary disease (COPD) (lipoxin and lipoxin analogs reduce lung damage from infiltrating inflammatory cells in this disease), aspirin-intolerant asthma, airway destruction and loss of function due to chronic inflammation, lung injury as a result of septic shock, and lung injury as a result of operating room-induced pumped lung syndrome (known as second-organ reperfusion injury of the lung).

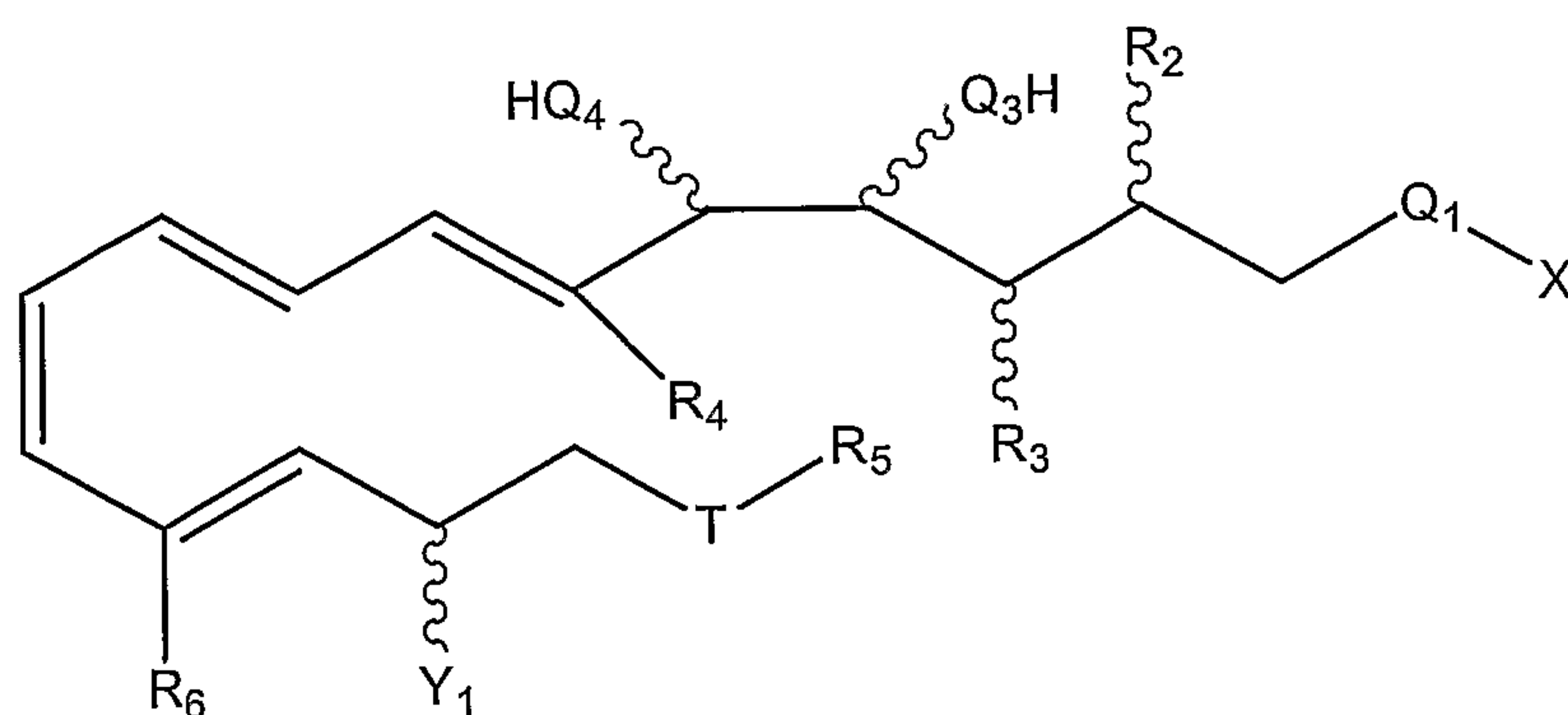
[051] For example, the lipoxin analogs of the invention are useful in the inhibition, treatment and/or prevention of: eosinophil-mediated inflammation of the lung or tissues; neutrophil-mediated inflammation of the lung; lymphocyte-mediated inflammation of the lung; cytokine and chemokine production, including interleukin-5, interleukin-13 and eotaxin; lipid mediator generation, including prostaglandin E₂ and cysteinyl leukotrienes; airway hyper-responsiveness; and airway and vascular inflammation.

[052] The lipoxins are lipoxygenase interaction products or are based on such products. By definition, the lipoxins include a conjugated tetraene containing structure and three alcohol groups as defining features in this class of compounds. The lipoxygenase/lipoxygenase interactions are brought about during cell-cell interactions, and are exemplified by cells carrying 5-lipoxygenase that could interact with either 15-lipoxygenase bearing cells, or 12-lipoxygenase bearing cells in vivo to generate lipoxins. The stereochemistry at the C15 position are of the S configuration. More recently it was discovered that cyclooxygenase, which is inhibited by aspirin and non-steroidal anti-inflammatory drugs, and is also involved in the generation of prostanoids, can play a role in the generation of novel class of tetraene containing compounds, which have been isolated structurally elucidated, namely 15 epimeric form of lipoxins.

[053] Suprisingly, with the discovery of cyclooxygenase-2 (an isoform of cyclooxygenase), that has an unusual enzymatic clef, it was discovered that acetylation of the enzyme by aspirin not only inhibits the generation of prostaglandins, but leads to continued activity of the COX-2 in its acetylated form to convert arachidonic acid to generate 15RHETE. 15RHETE is the biosynthetic pre-cursor that is converted by neutrophils to 15-epi lipoxins. Again, these compounds are cell cell interaction products that are generated by a unique biosynthetic route, namely the acetylation of cyclooxygenase-2 by aspirin and its conversion of arachidonic acid to endogenous mediators that mimic the action of natural lipoxins.

[054] These are what have been termed "aspirin triggered 15-epi lipoxins" or lipoxins that carry the alcohol group in the C15 position and the R configuration. The distinction between these two biochemical pathways, namely LO-LO interaction versus cyclooxygenase-2 and LO interaction provides some inferences as to the local mediator roles of these compounds and where there may be effective as endogenous biological agents to regulate/down regulate pro-inflammatory processes in vivo. The design and discovery of analogs based on these structures, at present appear to have similar biological actions and serve as mimetics of the natural lipoxins. However, in the not to distant future, it may become apparent that the aspirin COX-2 triggered pathway may have a unique profile of actions, namely actions that are specific for 15-epi lipoxins versus natural lipoxins. Setting a barrier between these two geniuses should not only reflect the chemical structures, but also their biosynthetic origins and potential role in human and mammalian biology and pharmacology.

[055] In one embodiment, lipoxin analogs useful in the invention have the formula (I)



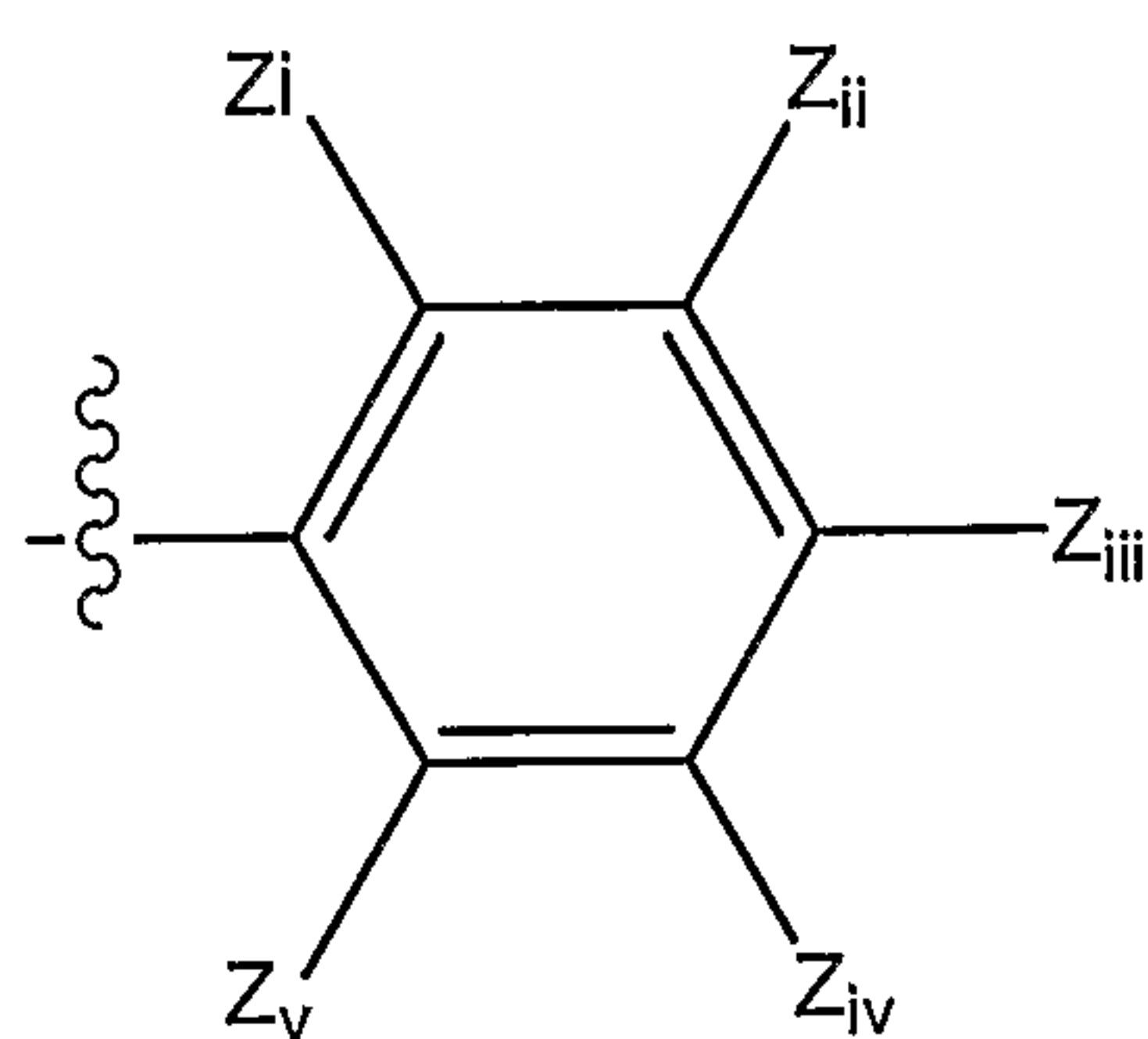
wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

5

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl

10



15

wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched, and hydroxyl;

- 5 (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms, inclusive;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is absent;

10

wherein Q_3 and Q_4 are each independently O , S or NH ;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

- (a) H ;
- 15 (b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;
- (d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be straight chain or branched; or
- 20 (e) $\text{R}_a\text{Q}_2\text{R}_b$ wherein Q_2 is $-\text{O}-$ or $-\text{S}-$; wherein R_a is alkylene of 0 to 6 carbon atoms, inclusive, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which may be straight chain or

branched, provided when R_b is 0, then R_b is a hydrogen atom;

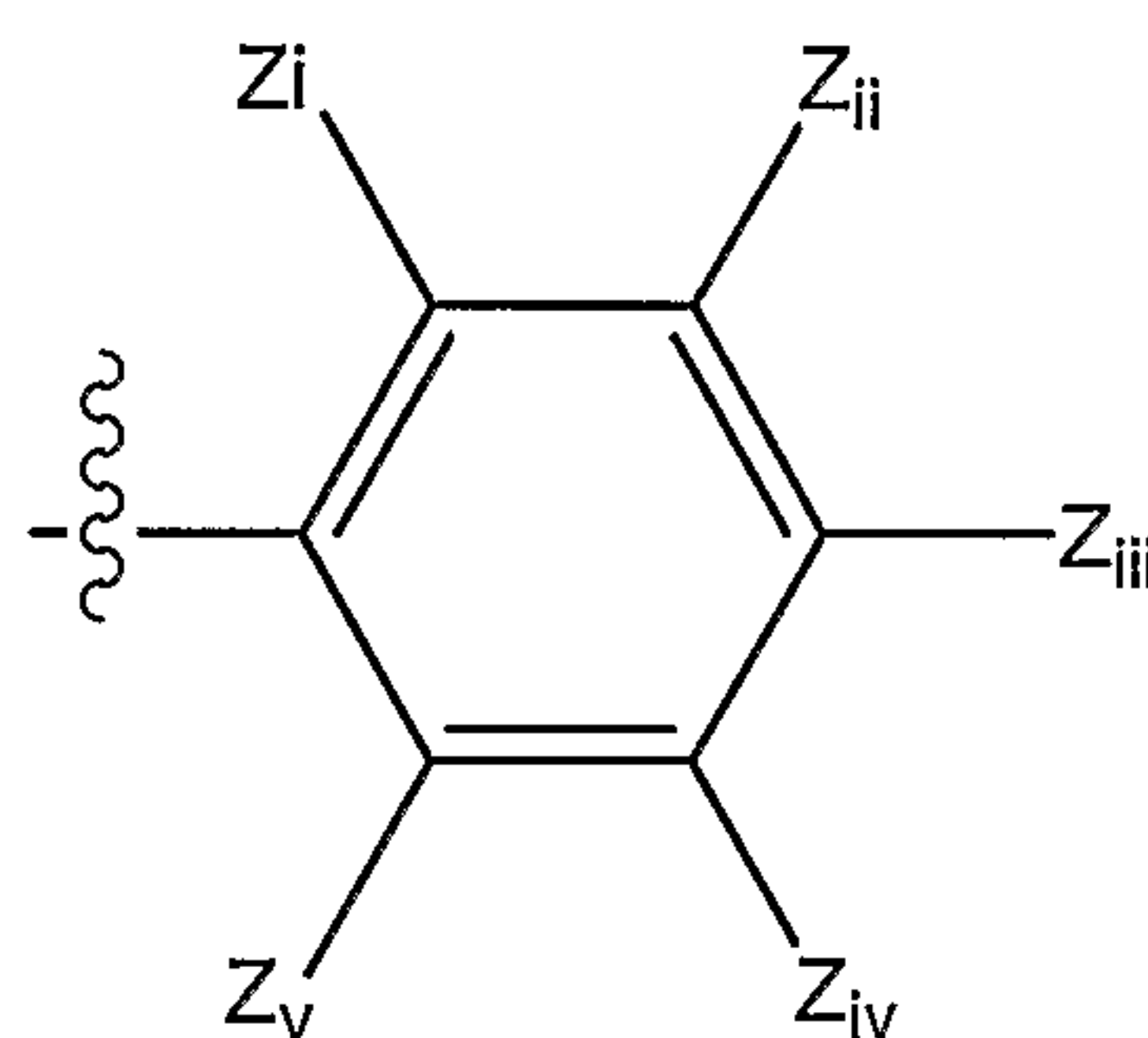
wherein R_4 is

5

- (a) H;
- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 is

10



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from —
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, —
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group;

15

wherein Y_1 is -OH, methyl, -SH, an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched, an alkoxy of 1 to 4 carbon atoms, inclusive, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano, nitro or a halogen;

5

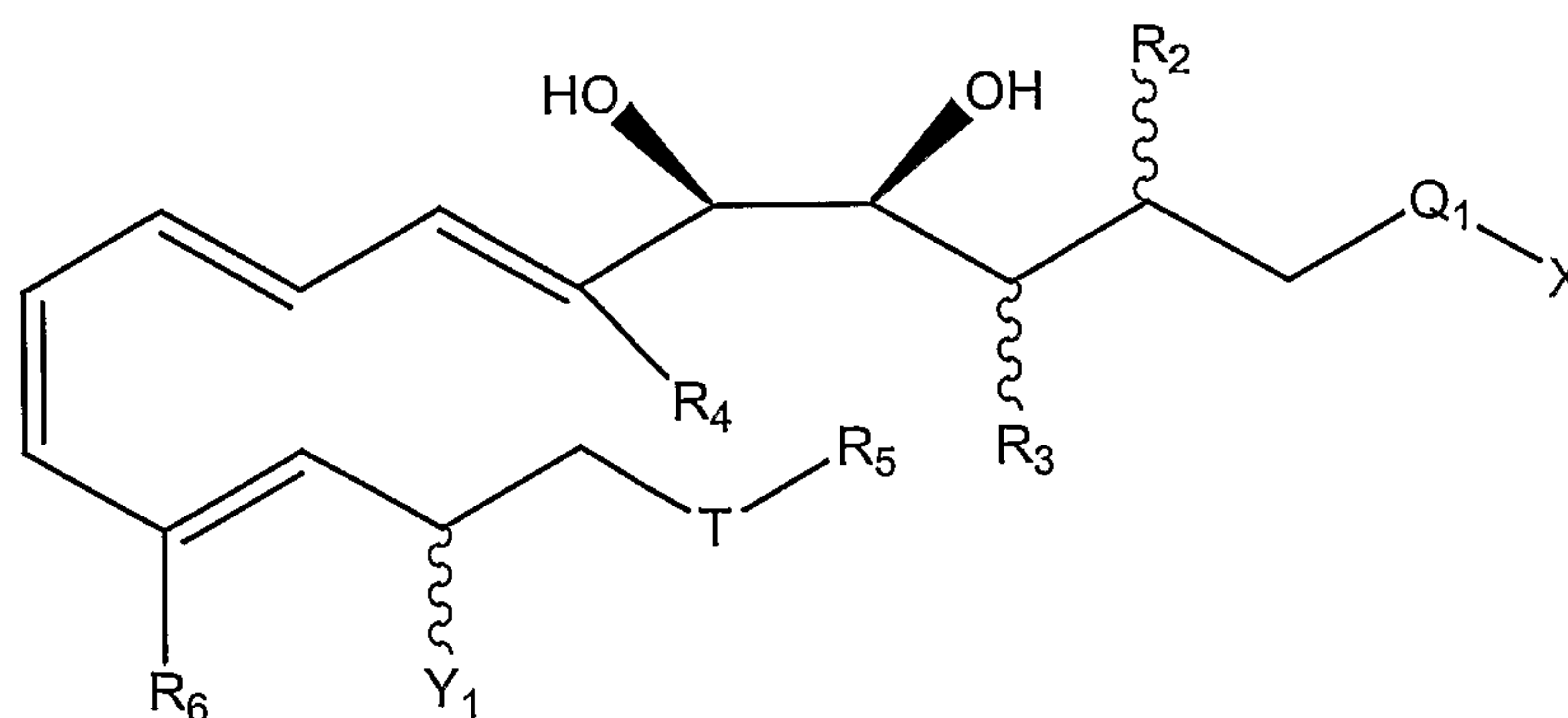
wherein R_6 is

- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

10

wherein T is O or S, and pharmaceutically acceptable salts thereof.

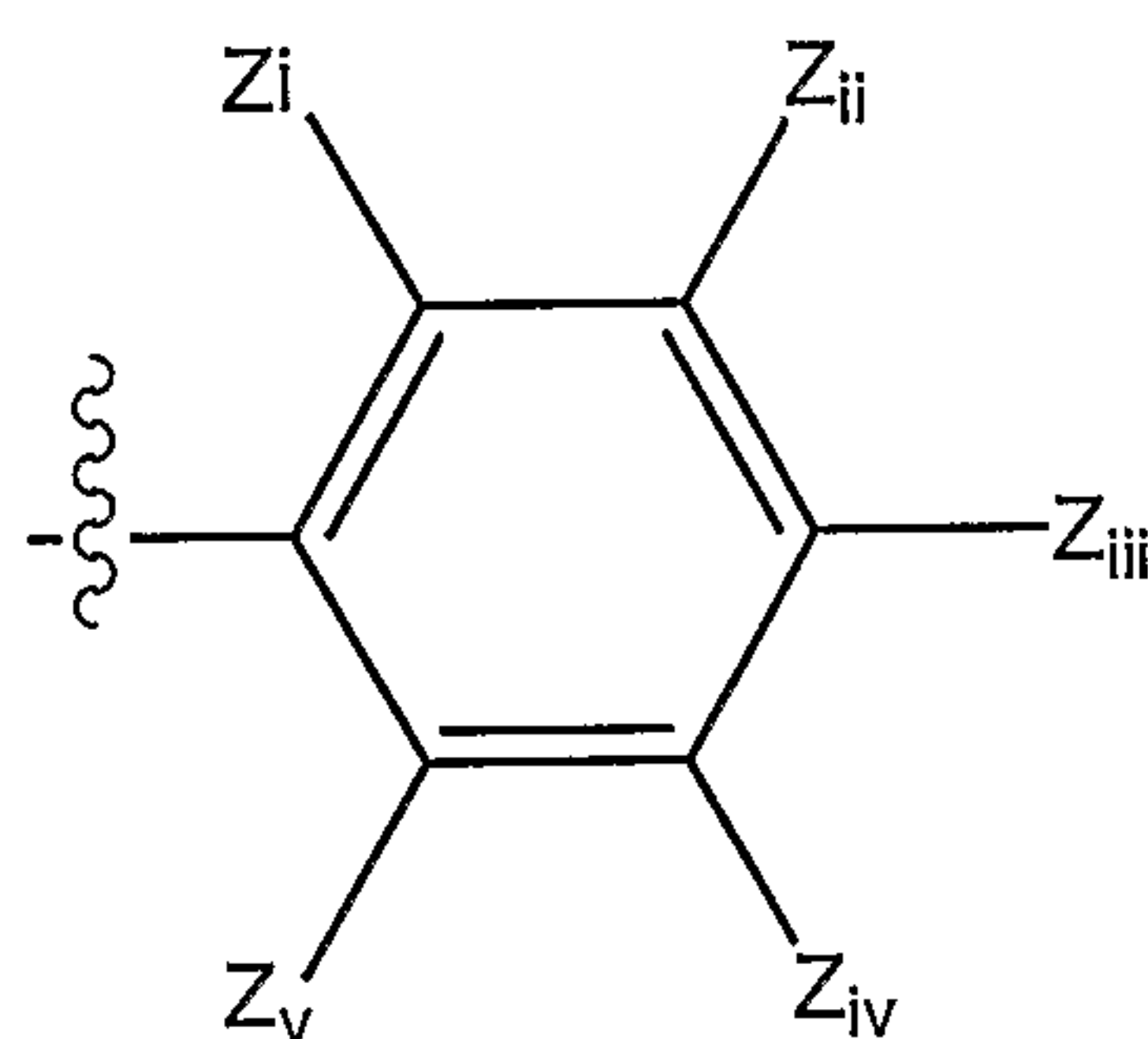
[056] In another embodiment, compounds useful in the invention have the
15 formula (II)



wherein X is R_1 , OR_1 , or SR_1 ;

wherein R_1 is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from —
 NO_2 , $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, —
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or

- (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms, inclusive;

wherein Q_1 is (C=O), SO_2 or (CN), provided when Q_1 is CN, then X is absent;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

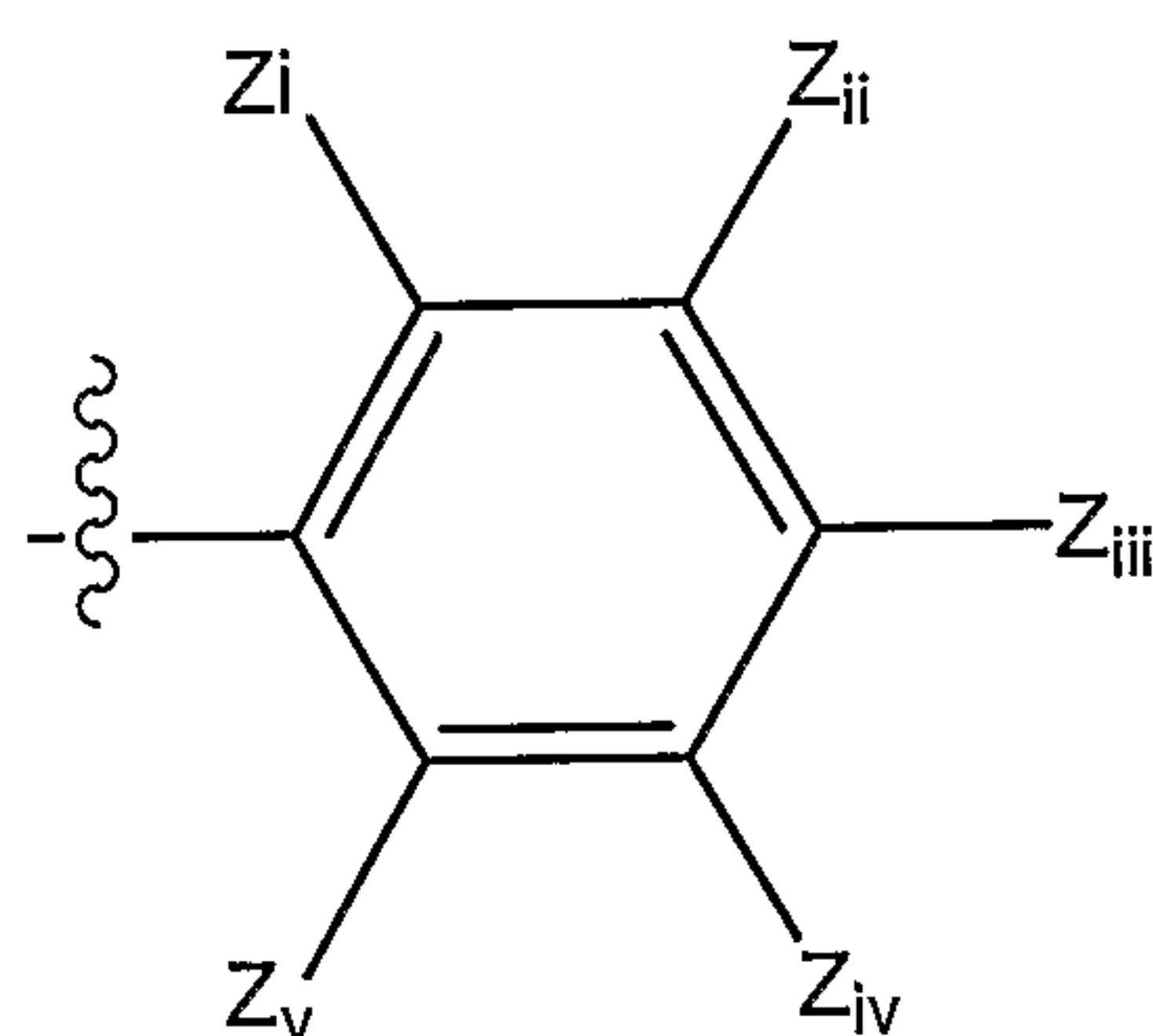
- (a) H;
- (b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;
- (d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be straight chain or branched; or
- (e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene of 0 to 6 carbon atoms, inclusive, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

- (a) H;

- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 is



5

wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-NO_2$, $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched, and hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

10

wherein Y_1 is $-OH$, methyl, $-SH$, an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched, an alkoxy of 1 to 4 carbon atoms, inclusive, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano, nitro or a halogen;

15

wherein R_6 is

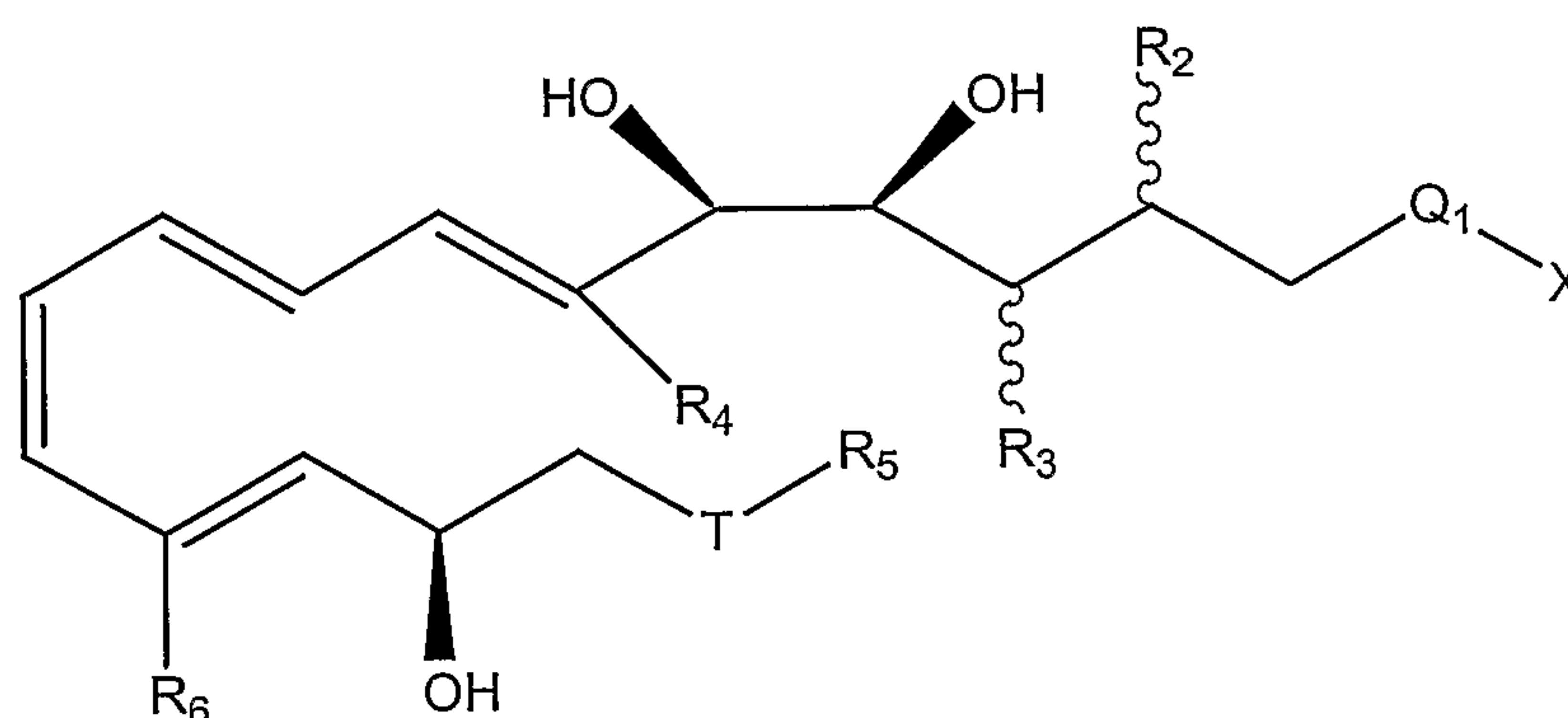
- (a) H;

- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof.

5

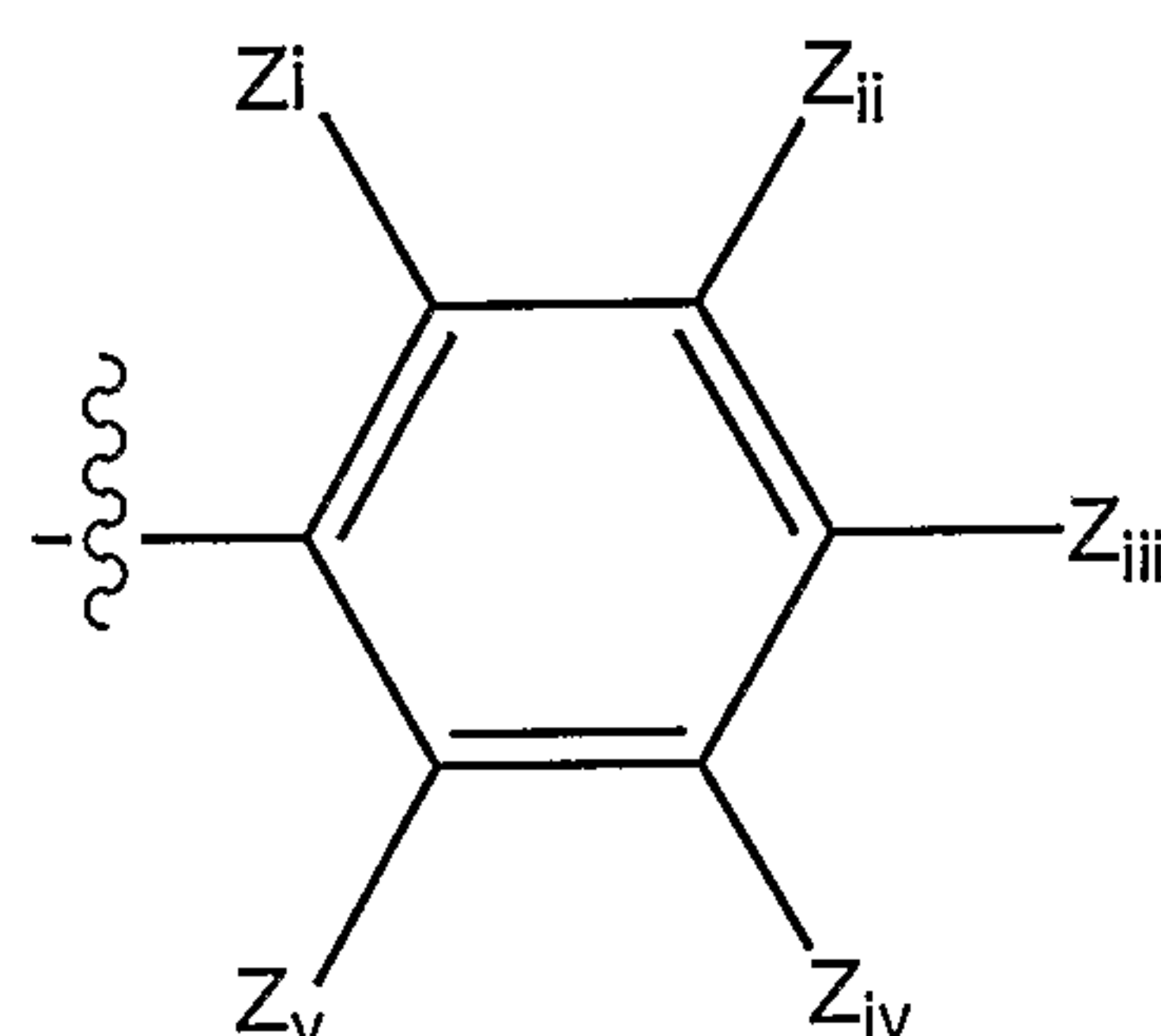
[057] The invention is also directed to useful lipoxin compounds having the formula (III)



10 wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- 15
- (i) a hydrogen atom;
 - (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
 - (iii) a cycloalkyl of 3 to 10 carbon atoms;
 - (iv) an aralkyl of 7 to 12 carbon atoms;
 - (v) phenyl;
 - (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

(vii) a detectable label molecule; or

(viii) a straight or branched chain alkenyl of 2 to 8

carbon atoms, inclusive;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is
 absent;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

(a) H ;

(b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched;

(c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;

(d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be straight chain or branched; or

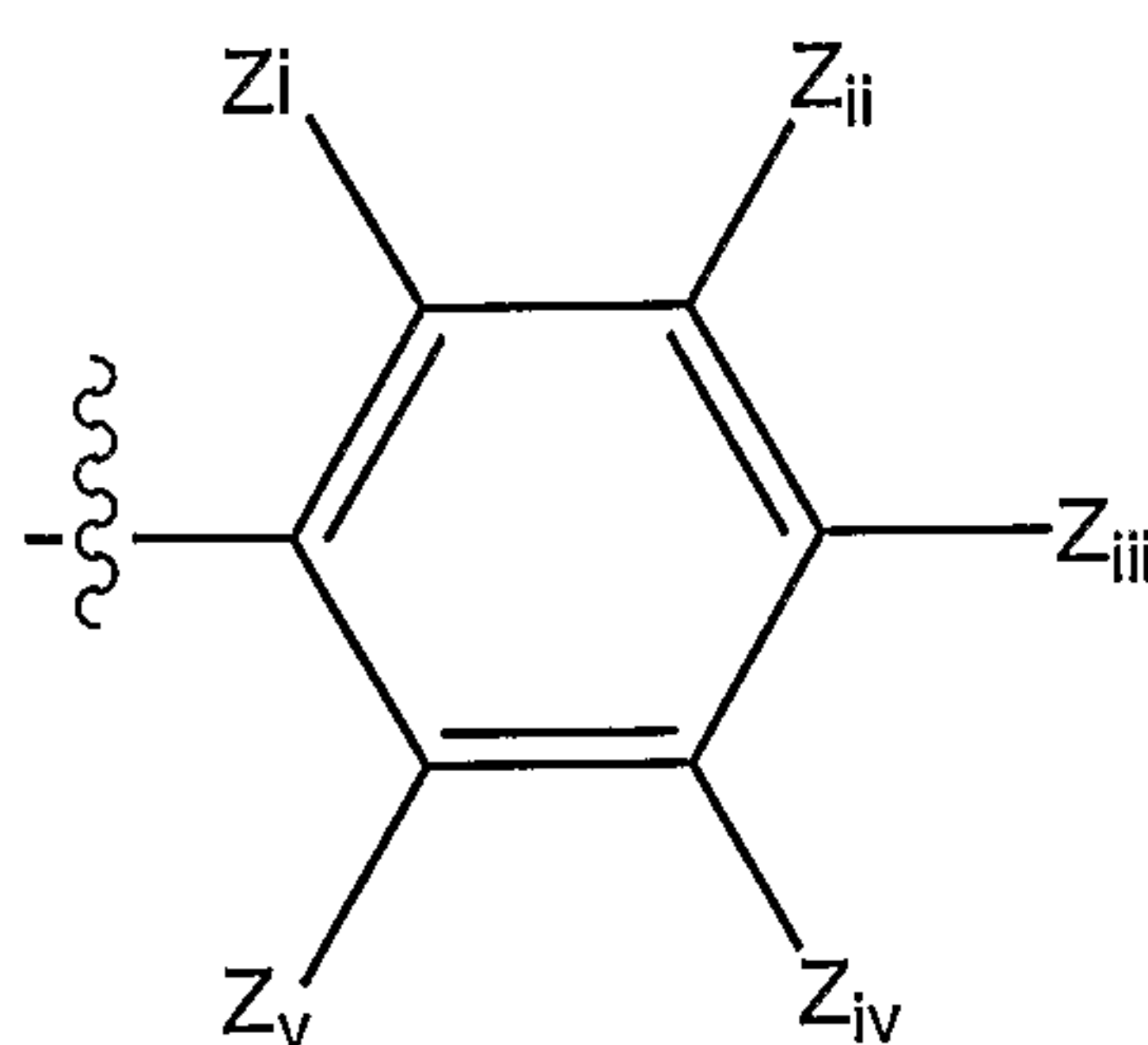
(e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene of 0 to 6 carbon atoms, inclusive, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

(a) H;

(b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group;

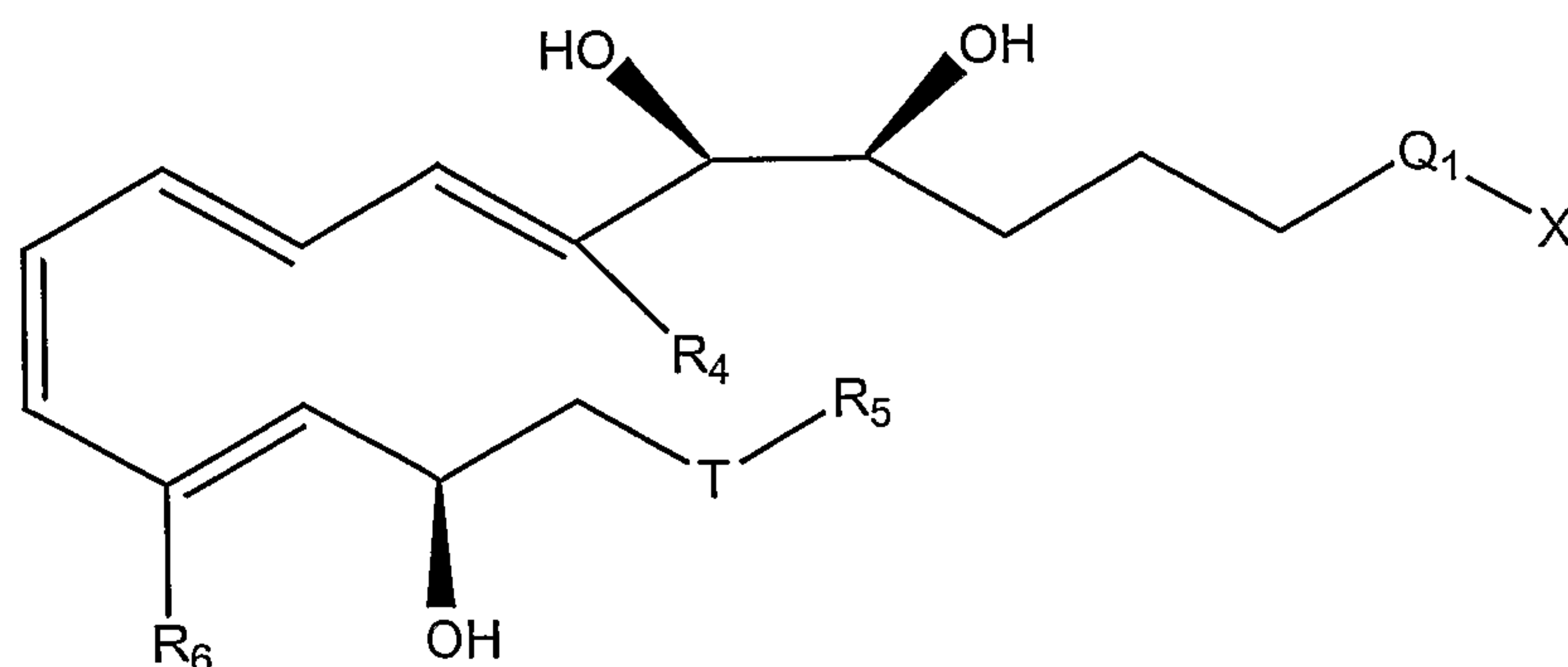
wherein R_6 is

(a) H;

(b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain
 or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof.

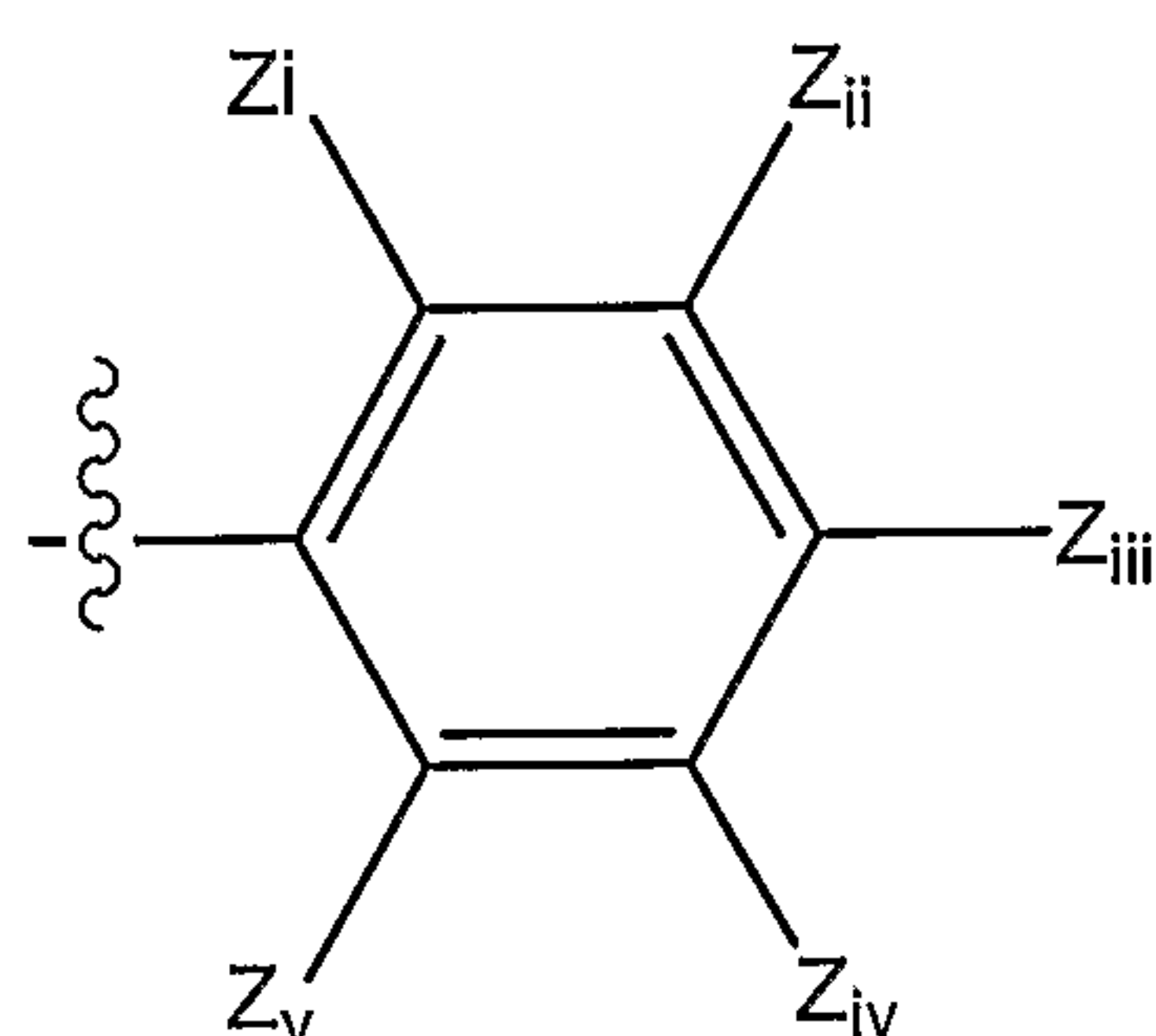
[058] The invention is further directed to useful lipoxin compounds having the
 formula (IV)



wherein X is R_1 , OR_1 , or SR_1 ;

wherein R_1 is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8
carbon atoms, inclusive;

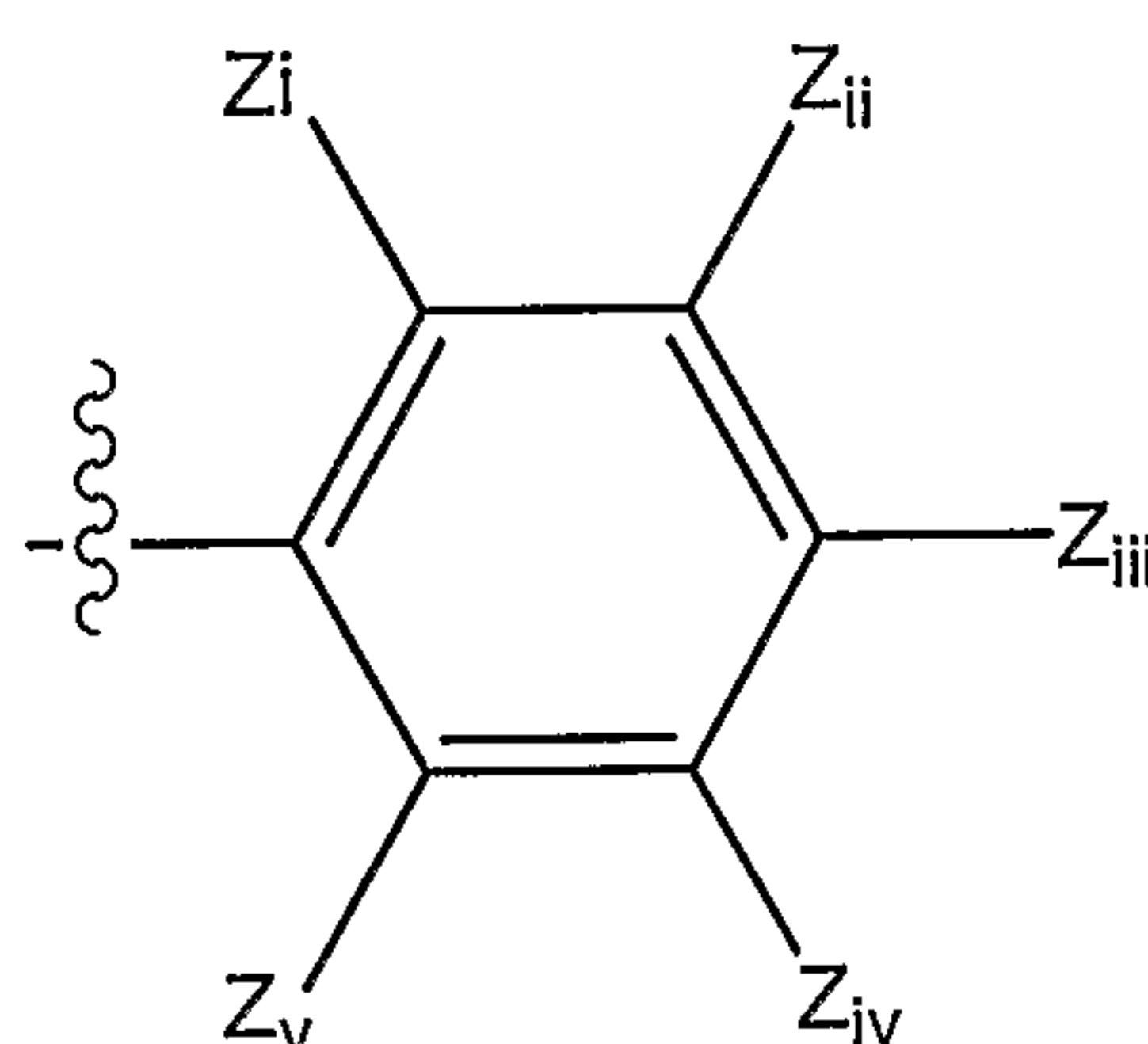
wherein Q_1 is (C=O), SO_2 or (CN), provided when Q_1 is CN, then X is absent;

5 wherein R_4 is

- (a) H;
- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

10

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-NO_2$, $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched, and hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

15

wherein R_6 is

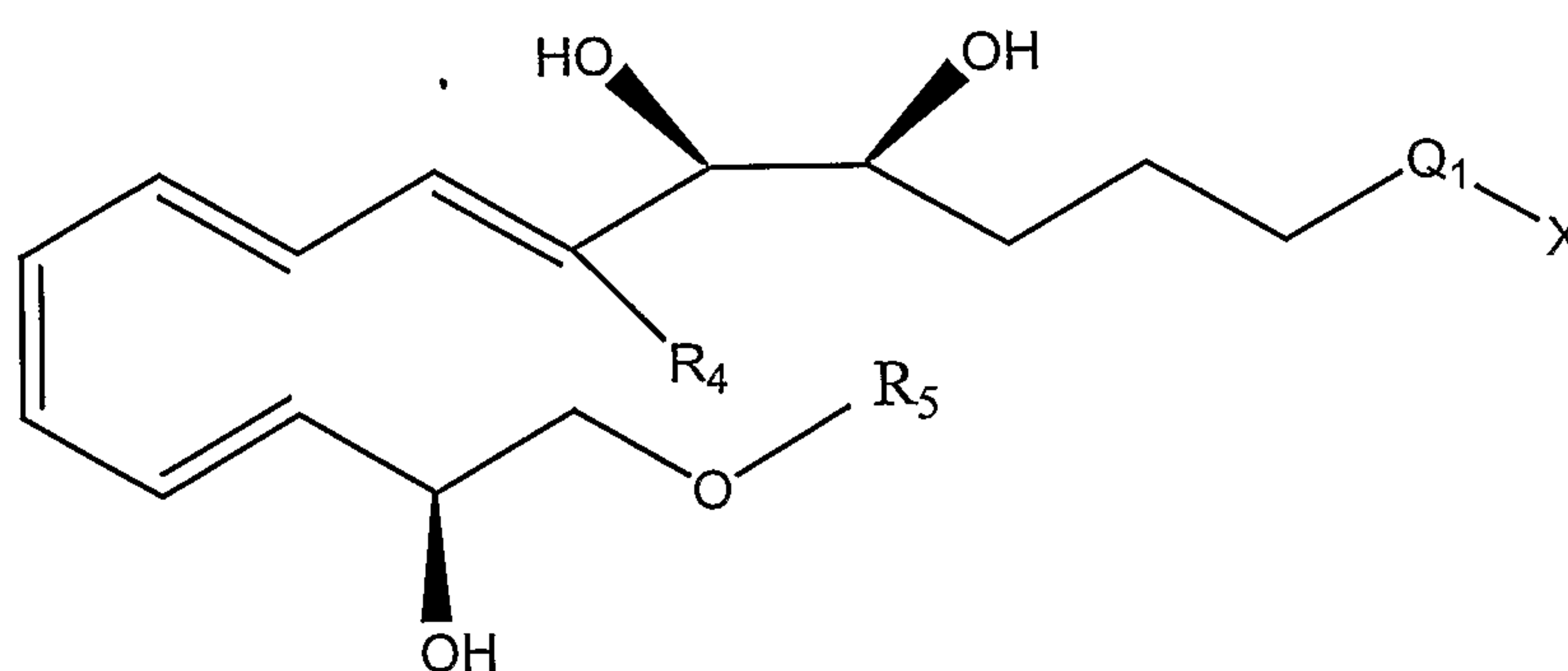
- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

5

wherein T is O or S, and pharmaceutically acceptable salts thereof.

[059] The invention is further directed to useful lipoxin compounds having the formula (V)

10



wherein X is R₁, OR₁, or SR₁;

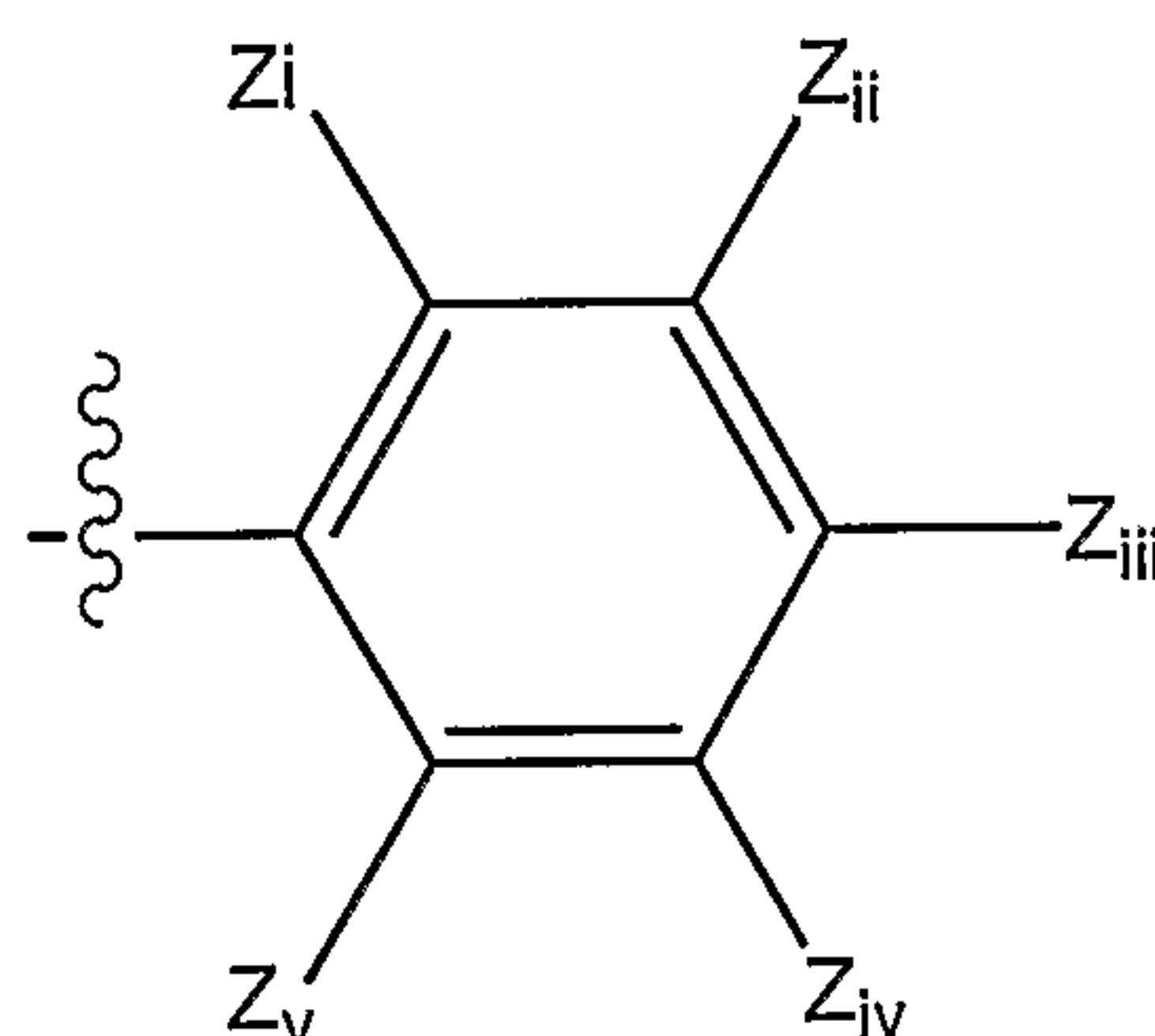
wherein R₁ is

15

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;

20

(vi) substituted phenyl



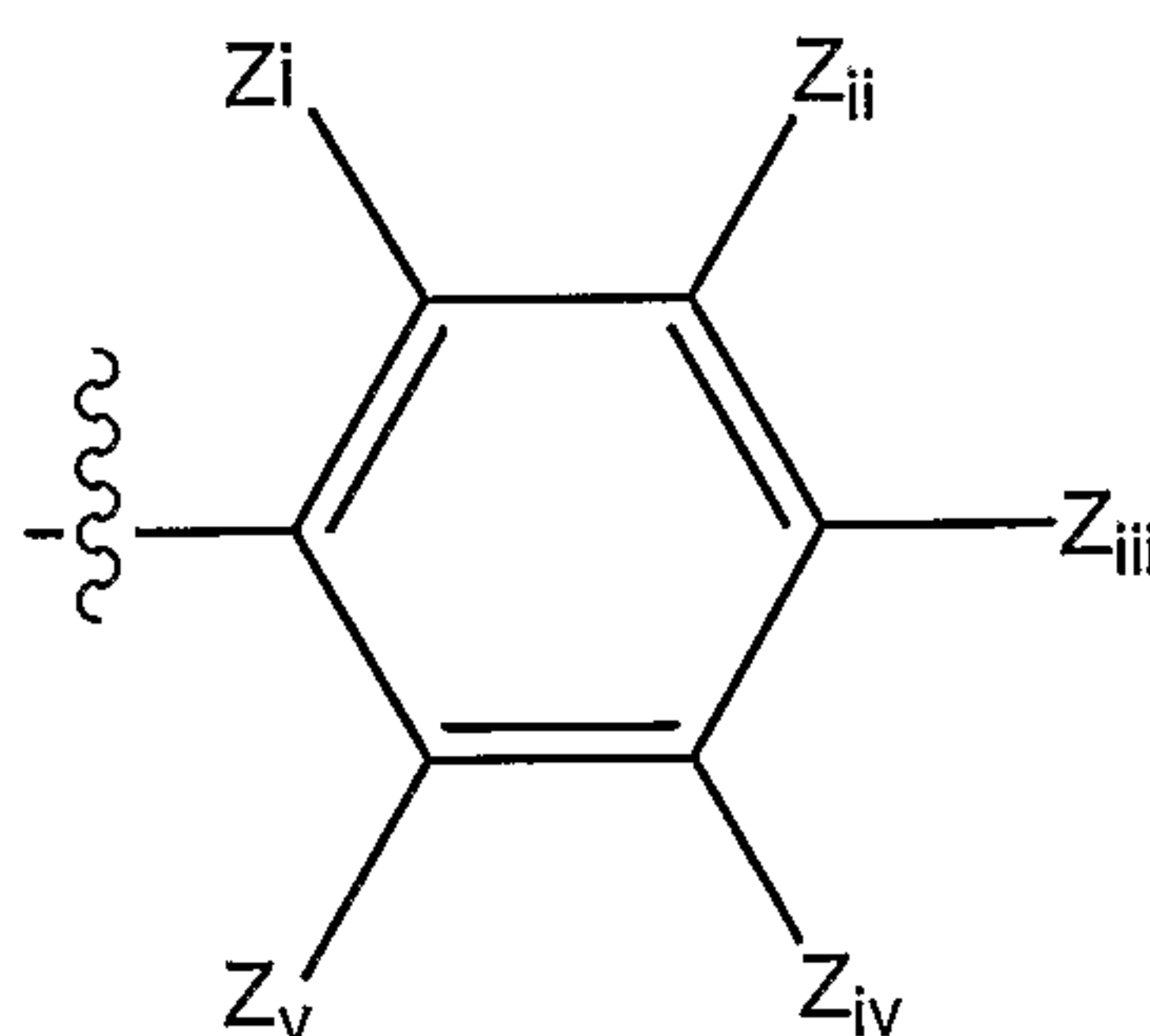
wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 5 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

(vii) a detectable label molecule; or
 10 (viii) a straight or branched chain alkenyl of 2 to 8
 carbon atoms, inclusive;

wherein R_4 is

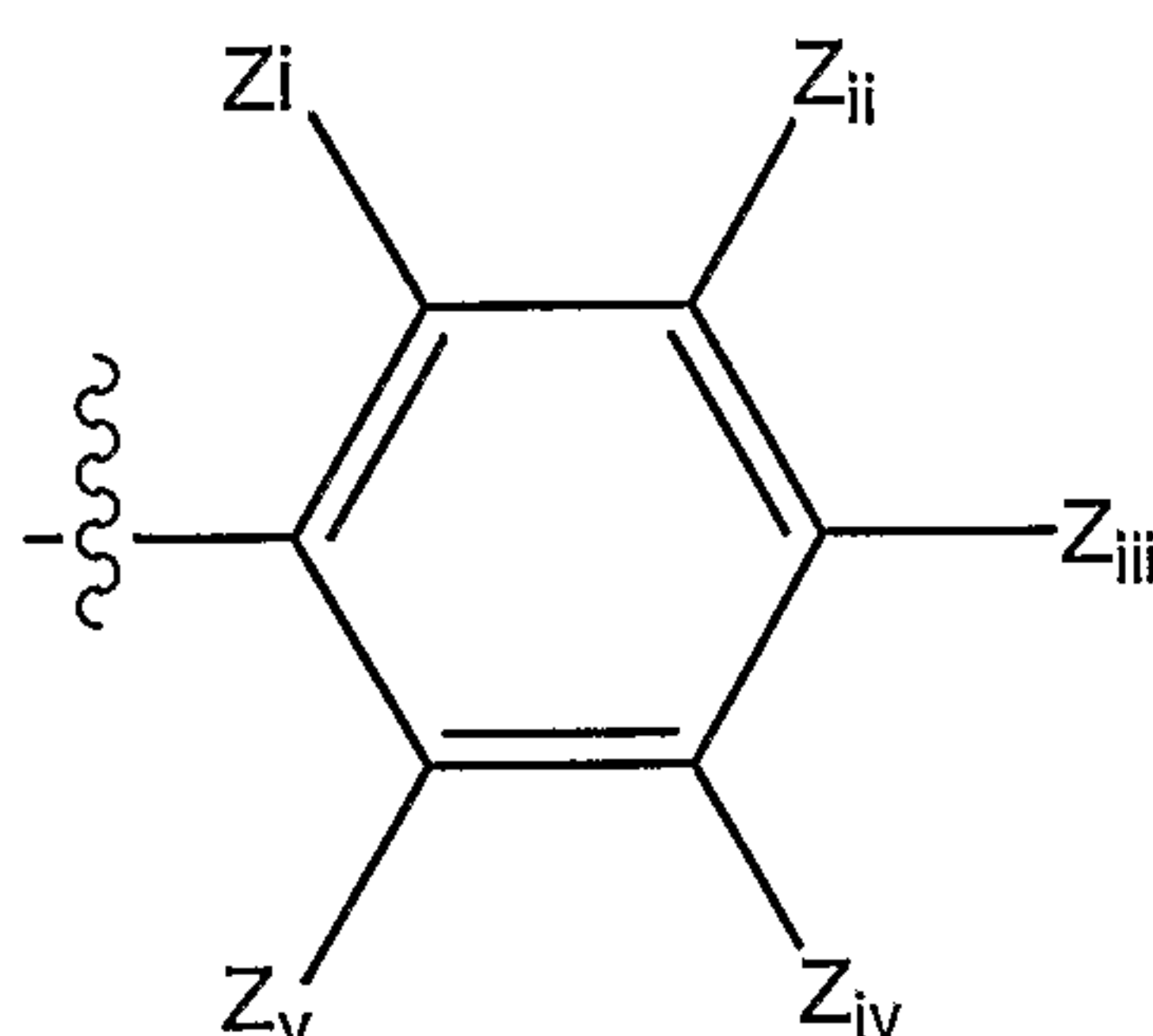
(a) H;
 15 (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a
 straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 5 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group; and
 pharmaceutically acceptable salts thereof.

10 **[060]** In certain embodiments, X is OR_1 wherein R_1 is a hydrogen atom, an
 alkyl group of 1 to 4 carbon atoms or a pharmaceutically acceptable salt, Q_1 is
 $\text{C}=\text{O}$, R_2 and R_3 , if present, are hydrogen atoms, R_4 is a hydrogen atom or methyl,
 Q_3 and Q_4 , if present, are both O, R_6 , if present, is a hydrogen atom, Y_1 , if
 present, is OH, T is O and R_5 is a substituted phenyl, e.g.,



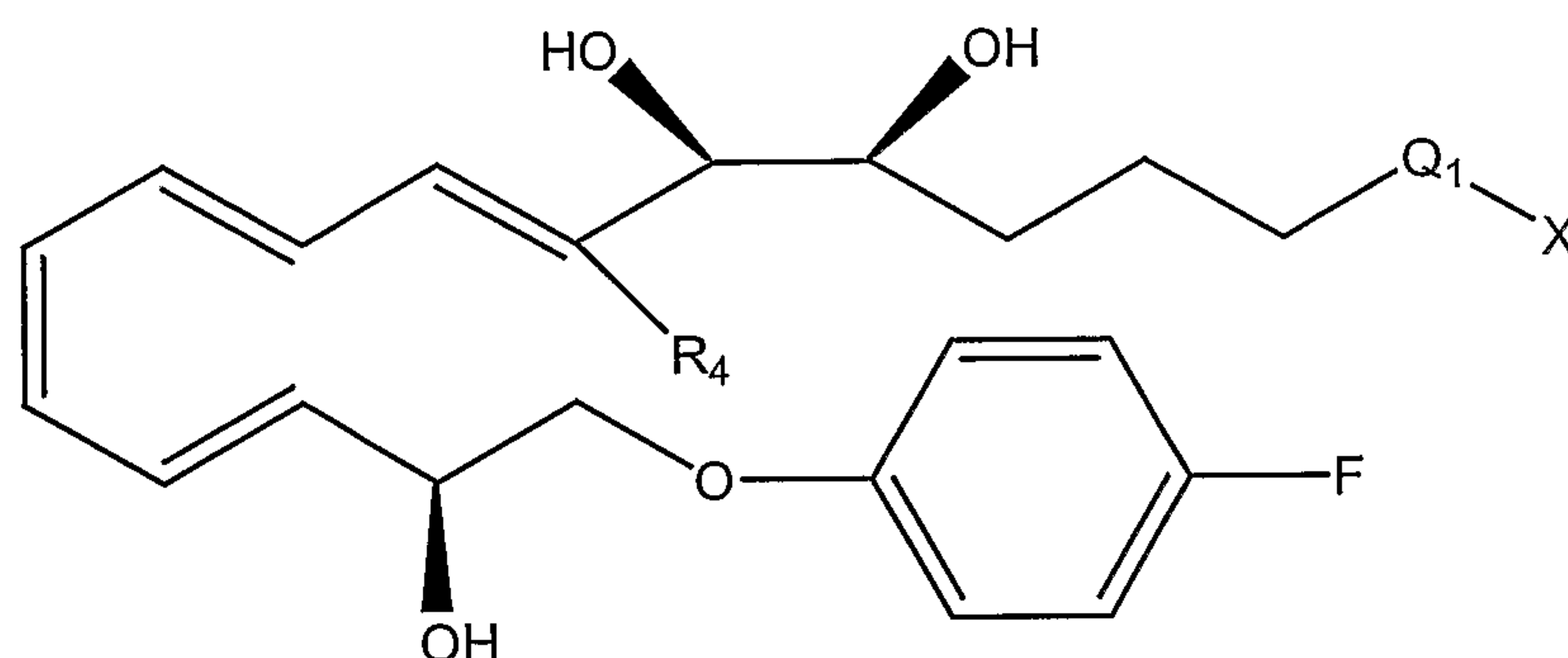
15

wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$,

wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or
 branched, and hydroxyl. In certain embodiments for R_5 , para-fluorophenyl

5 and/or unsubstituted phenyl are optimal, e.g., 15-epi-16-(para-fluoro)-phenoxy-
 LXA_4 , 16-(para-fluoro)-phenoxy- LXA_4 , 15-epi-16-phenoxy- LXA_4 or 16-
 phenoxy- LXA_4 . The compounds encompassed by U.S. Patent 5,441,951 are
 excluded from certain aspects of the present invention.

[061] In still another aspect, the present invention is directed to pharmaceutical
 10 compositions including compounds having the above-described formulae and a
 pharmaceutically acceptable carrier. In one embodiment, the compound is



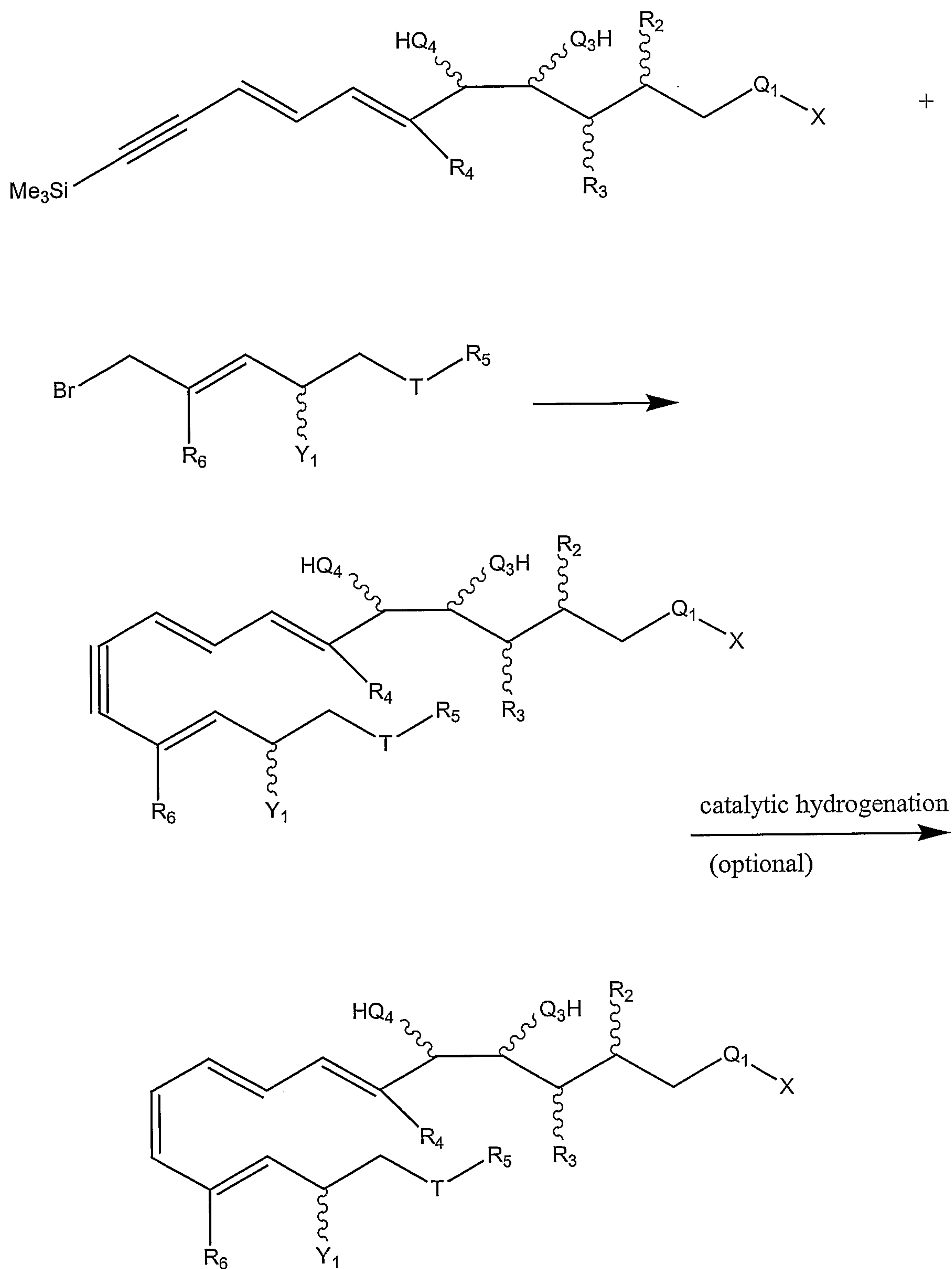
[062] In certain embodiments, Y_1 is a hydroxyl and the carbon bearing the
 15 hydroxyl can have an R or S configuration. In most embodiments, the chiral
 carbon bearing the hydroxyl group, e.g., Y_1 , is designated as a 15-epi-lipoxin as
 is known in the art.

[063] In certain embodiments the chirality of the carbons bearing the R_2 , R_3 , Q_3
 and Q_4 groups can each independently be either R or S. In certain embodiments,
 20 Q_3 and Q_4 have the chiralities shown in structures II, III, IV or V.

[064] In certain embodiments, R_4 is a hydrogen. In other embodiments, R_6 is a
 hydrogen.

[065] Additionally, R₅ can be a substituted or unsubstituted, branched or unbranched alkyl group having between 1 and about 6 carbon atoms, preferably between 1 and 4 carbon atoms, most preferably between 1 and 3, and preferably one or two carbon atoms. The carbon atoms can have substituents which include
5 halogen atoms, hydroxyl groups, or ether groups.

[066] The compounds useful in the present invention can be prepared by the following synthetic scheme:



wherein X, Q₁, Q₃, Q₄, R₂, R₃, R₄, R₅, R₆, Y₁ and T are as defined above.

Suitable methods known in the art to can be used to produce each fragment. For

example, the acetylenic fragment can be prepared by the methods discussed in

Nicolaou, K.C. et al. (1991) *Angew. Chem. Int. Ed. Engl.* 30:1100; Nicolaou,

5 K.C. et al. (1989) *J. Org. Chem.* 54:5527; Webber, S.E. et al. (1988) *Adv. Exp.*

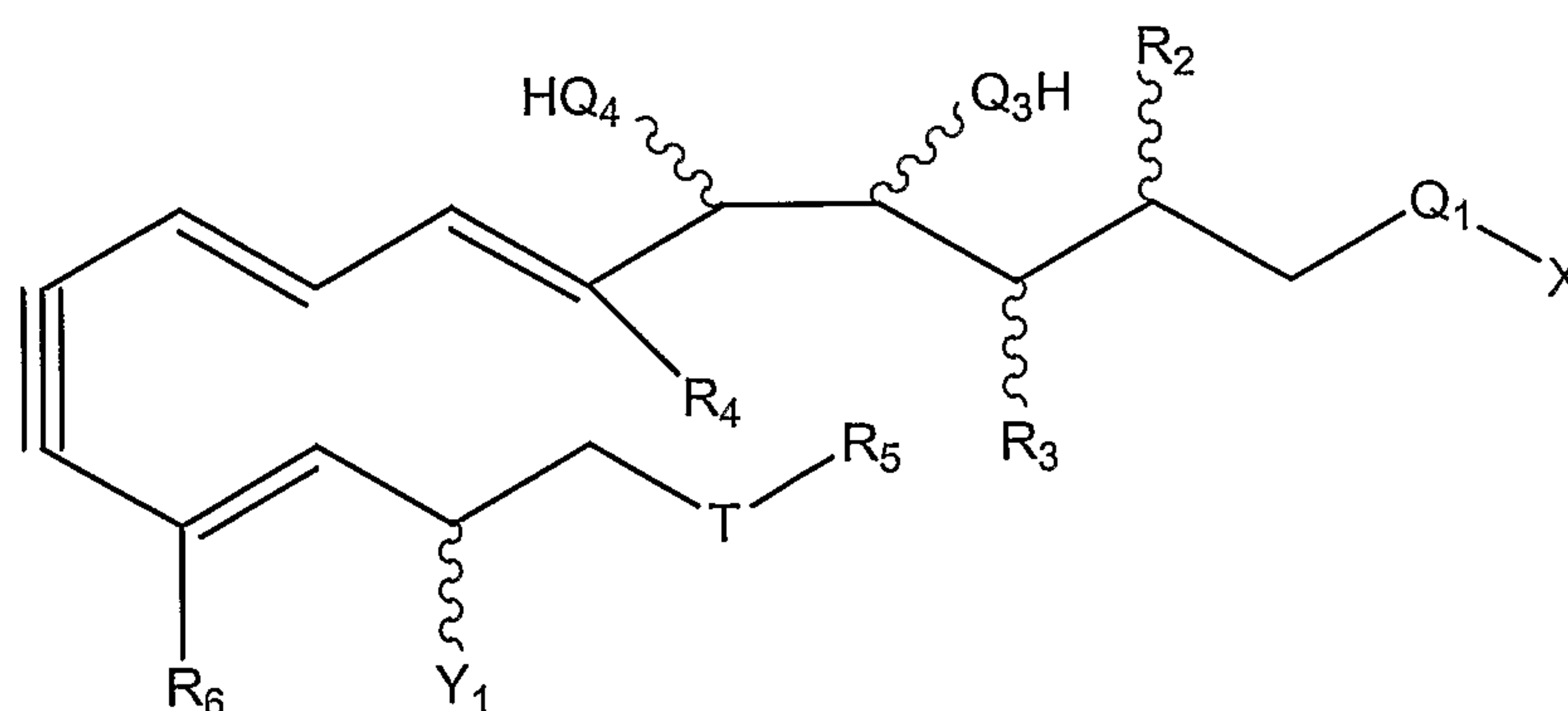
Med. Biol. 229:61; and U.S. Patent 5,441,951. The second fragment can be

prepared by the methods of Raduchel, B. and Vorbruggen, H. (1985) *Adv.*

Prostaglandin Thromboxane Leukotriene Res. 14:263.

[067] In another embodiment, compounds useful in the invention have the

10 formula (VI)



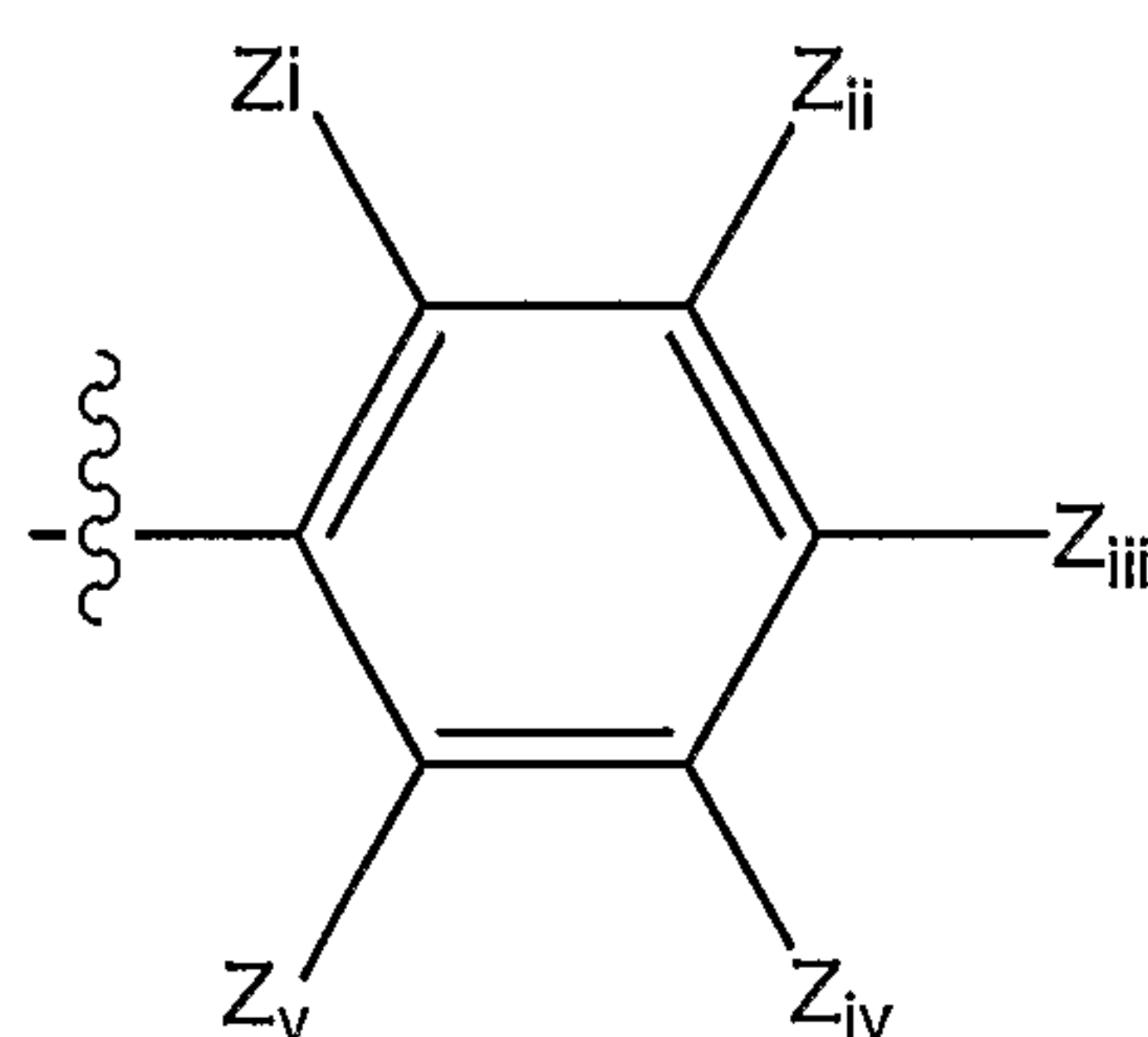
wherein X is R₁, OR₁, or SR₁;

15 wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;

- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl

5



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, -
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

10

- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8
 carbon atoms, inclusive;

15

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is
 absent;

wherein Q_3 and Q_4 are each independently O , S or NH ;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

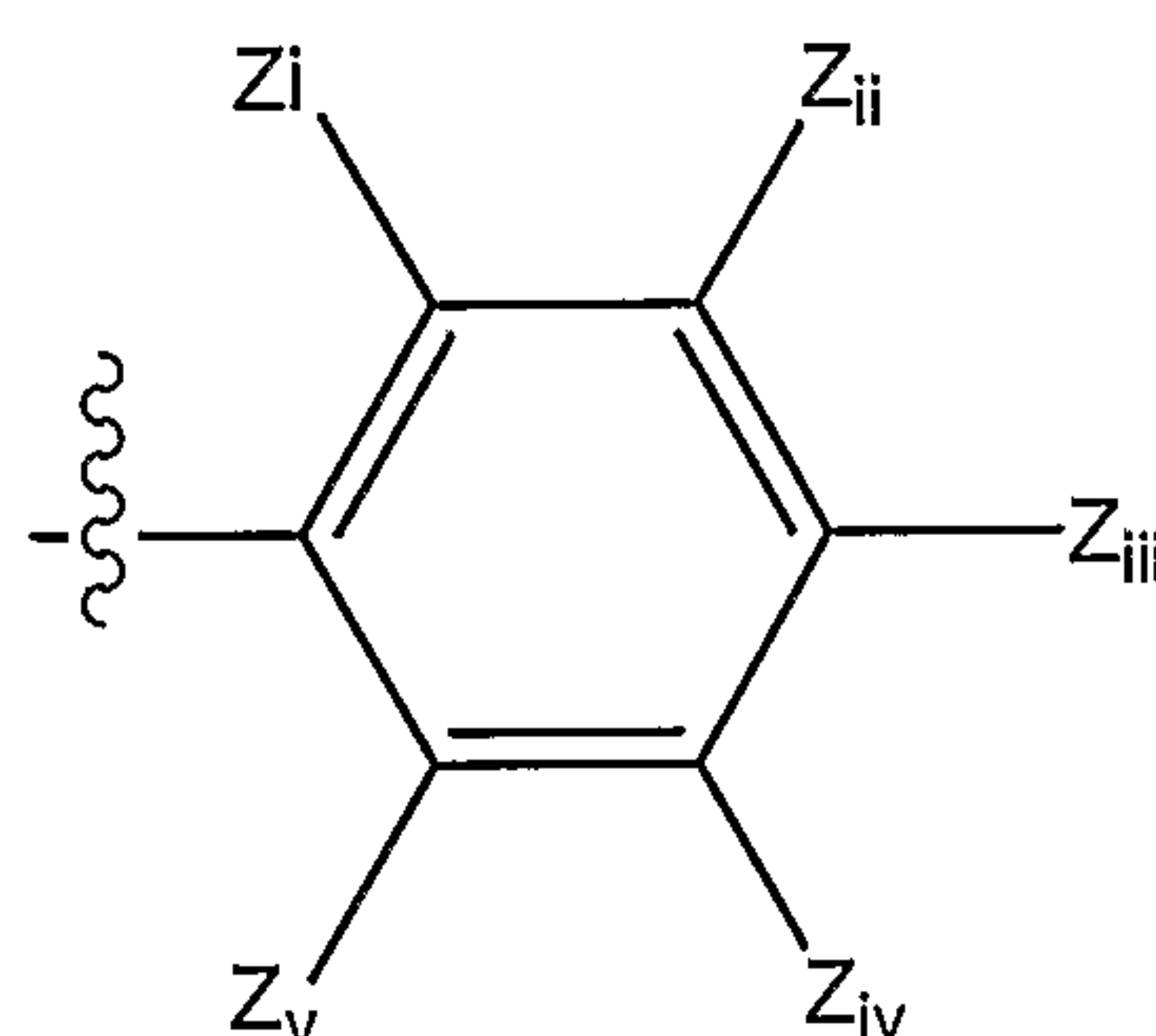
- 5
- (a) H;
 - (b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a
straight chain or branched;
 - (c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;
 - (d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be
straight chain or branched; or
 - (e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene
10 of 0 to 6 carbon atoms, inclusive, which may be straight
chain or branched and wherein R_b is alkyl of 0 to 8
carbon atoms, inclusive, which may be straight chain or
branched, provided when R_b is 0, then R_b is a hydrogen
atom;

15 wherein R_4 is

- (a) H;
- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a
straight chain or branched;

20

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 5 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group;

wherein Y_1 is $-\text{OH}$, methyl, $-\text{SH}$, an alkyl of 2 to 4 carbon atoms,
 10 inclusive, straight chain or branched, an alkoxy of 1 to 4 carbon atoms,
 inclusive, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano,
 nitro or a halogen;

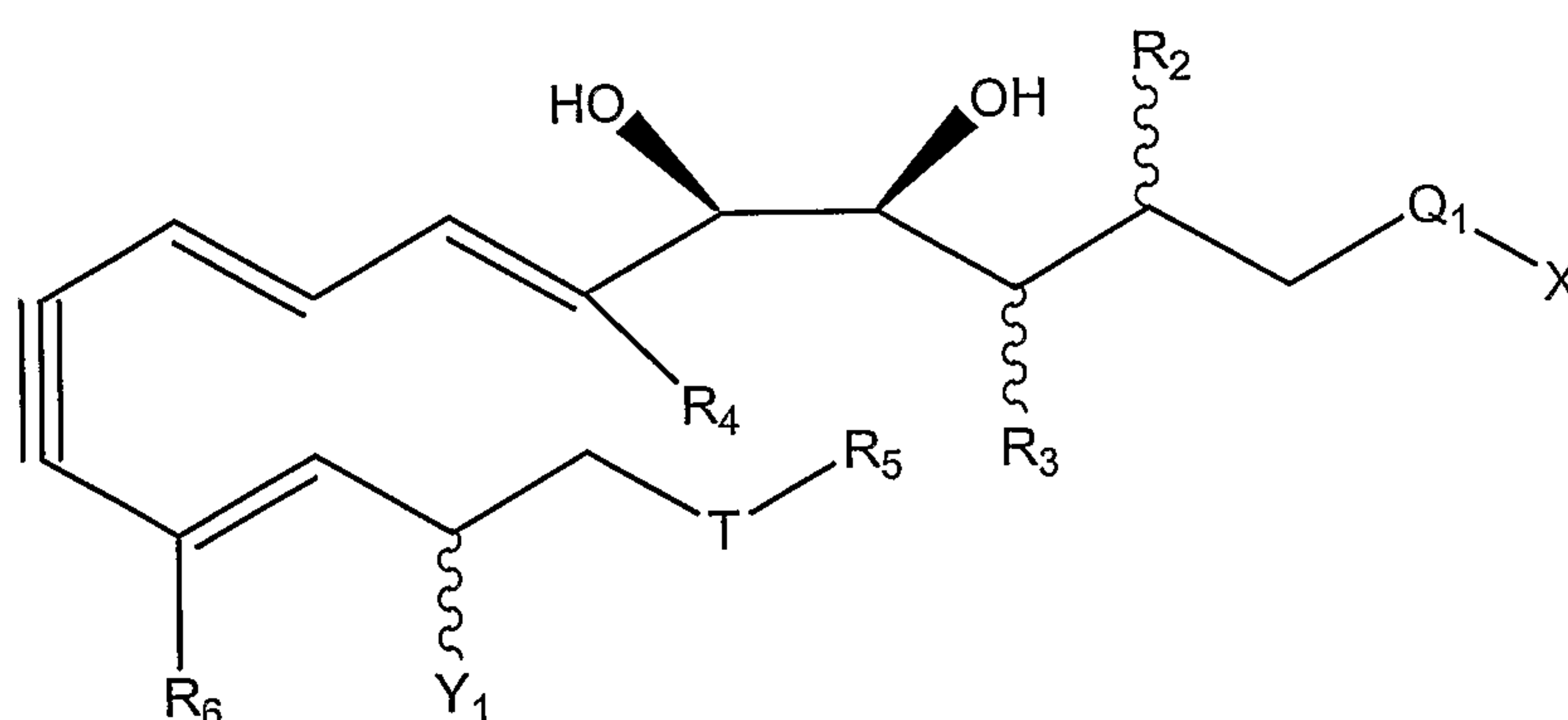
wherein R_6 is

- 15
- (a) H;
 - (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain
or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof.

[068] It should be understood that the depictions of the acetylenic portion of the lipoxin compounds described throughout this specification are drawn for convenience and not as actual representation of bond angles.

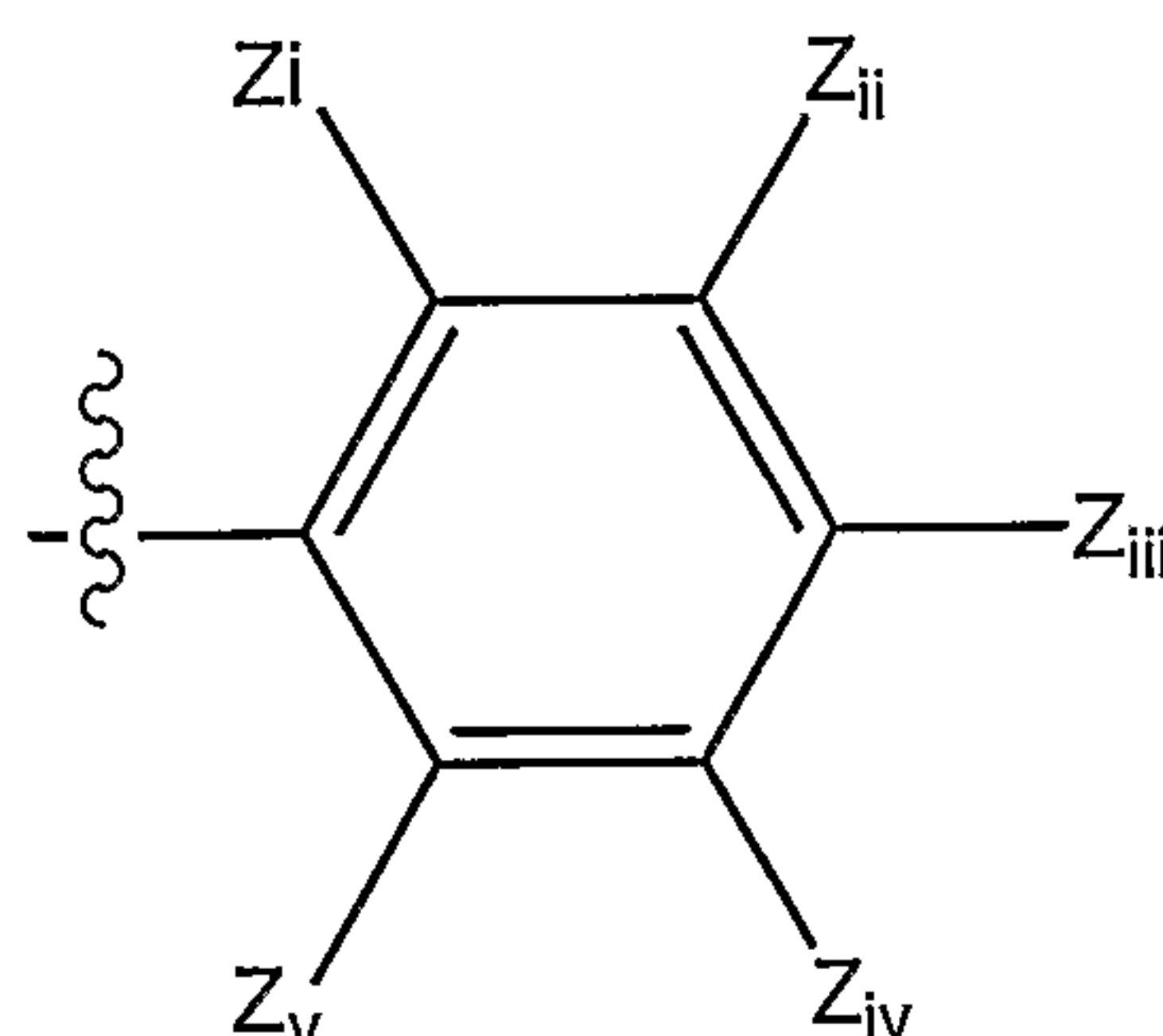
5 [069] In another embodiment, compounds useful in the invention have the formula (VII)



wherein X is R₁, OR₁, or SR₁;

10 wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
- 15 (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl;

(vii) a detectable label molecule; or

(viii) a straight or branched chain alkenyl of 2 to 8

carbon atoms, inclusive;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is
 absent;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

(a) H ;

(b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched;

(c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;

(d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be straight chain or branched; or

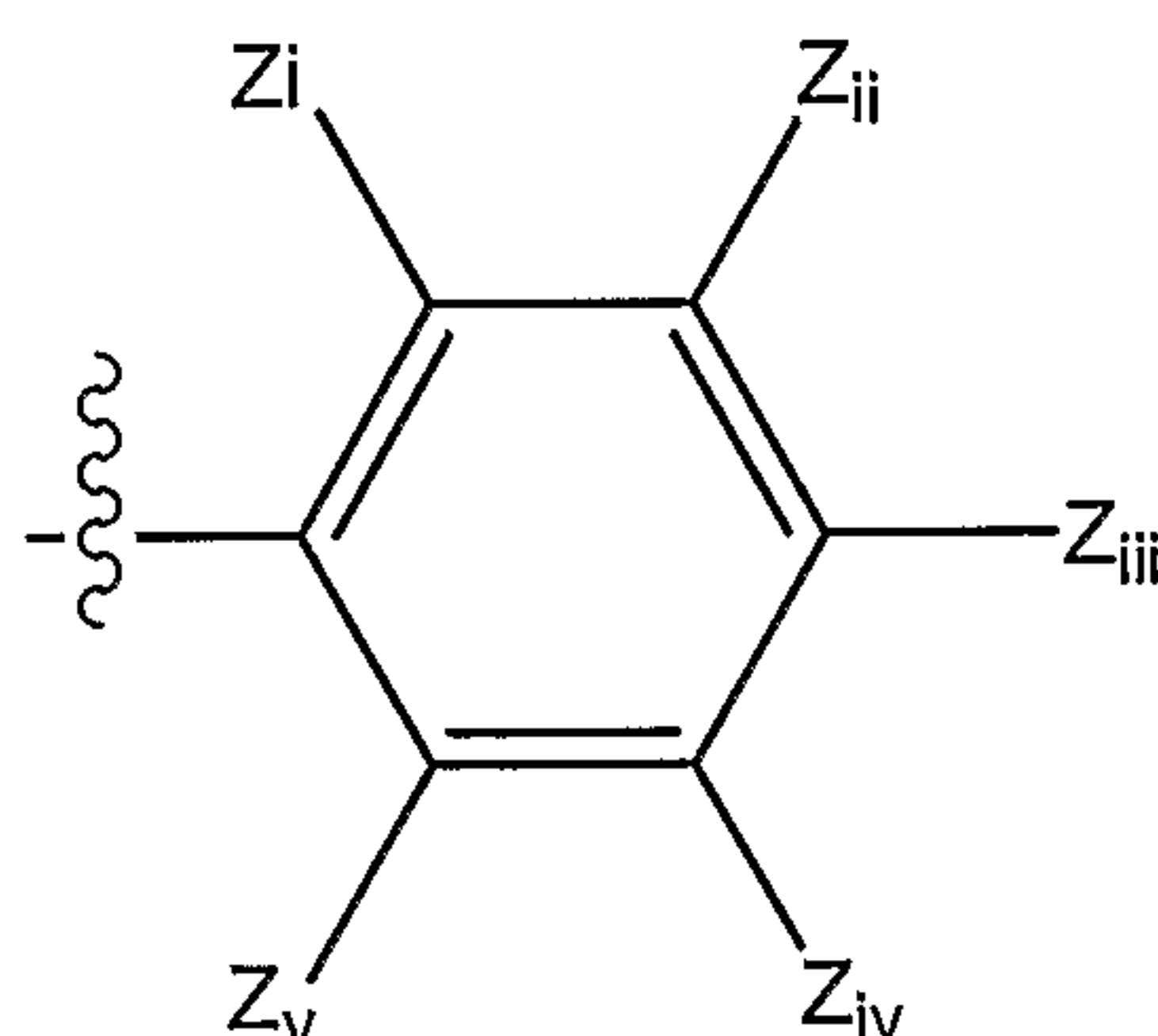
(e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene of 0 to 6 carbon atoms, inclusive, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

(a) H;

(b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-NO_2$, $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a

straight chain or branched, and hydroxyl or a substituted or
unsubstituted, branched or unbranched alkyl group;

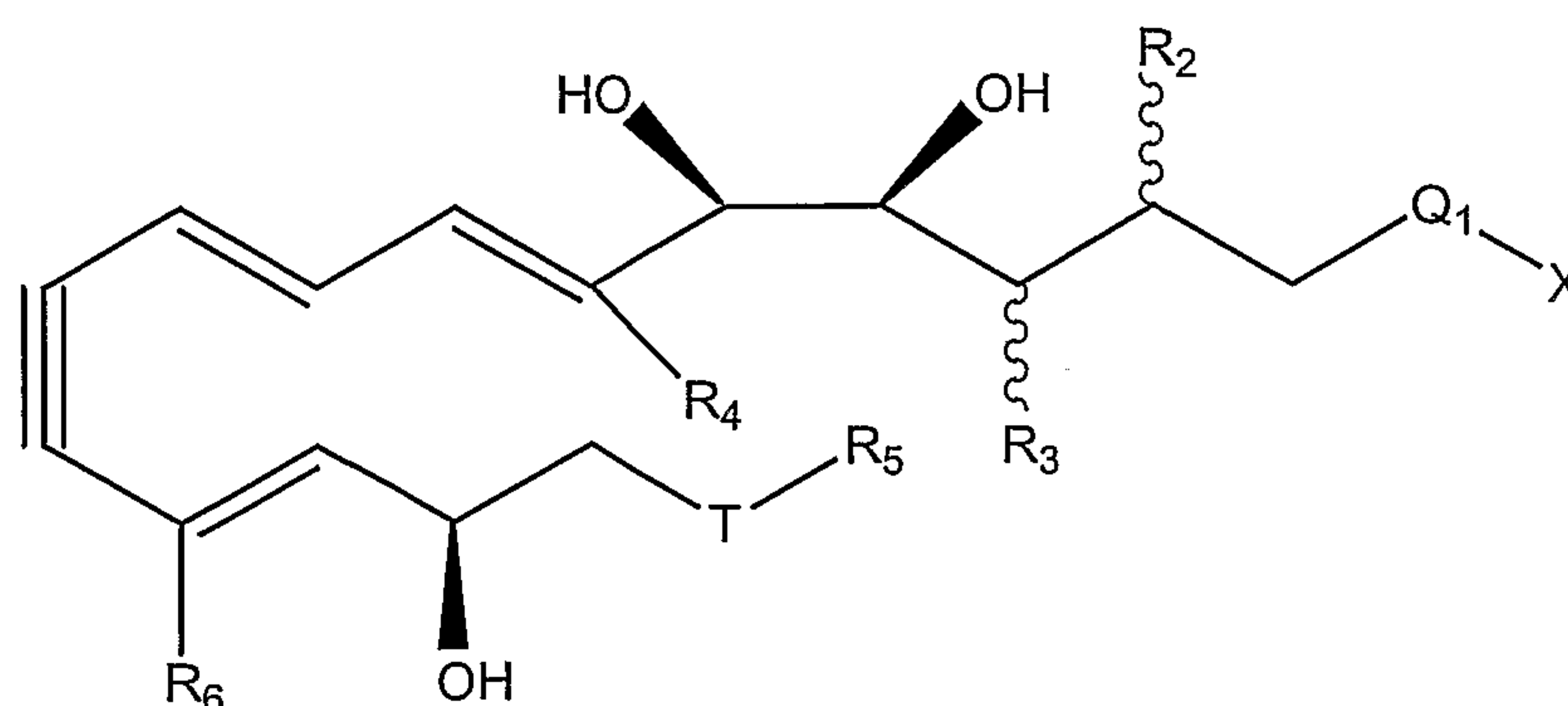
wherein Y_1 is -OH, methyl, -SH, an alkyl of 2 to 4 carbon atoms,
inclusive, straight chain or branched, an alkoxy of 1 to 4 carbon atoms,
inclusive, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano,
nitro or a halogen;

wherein R_6 is

- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain
or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof.

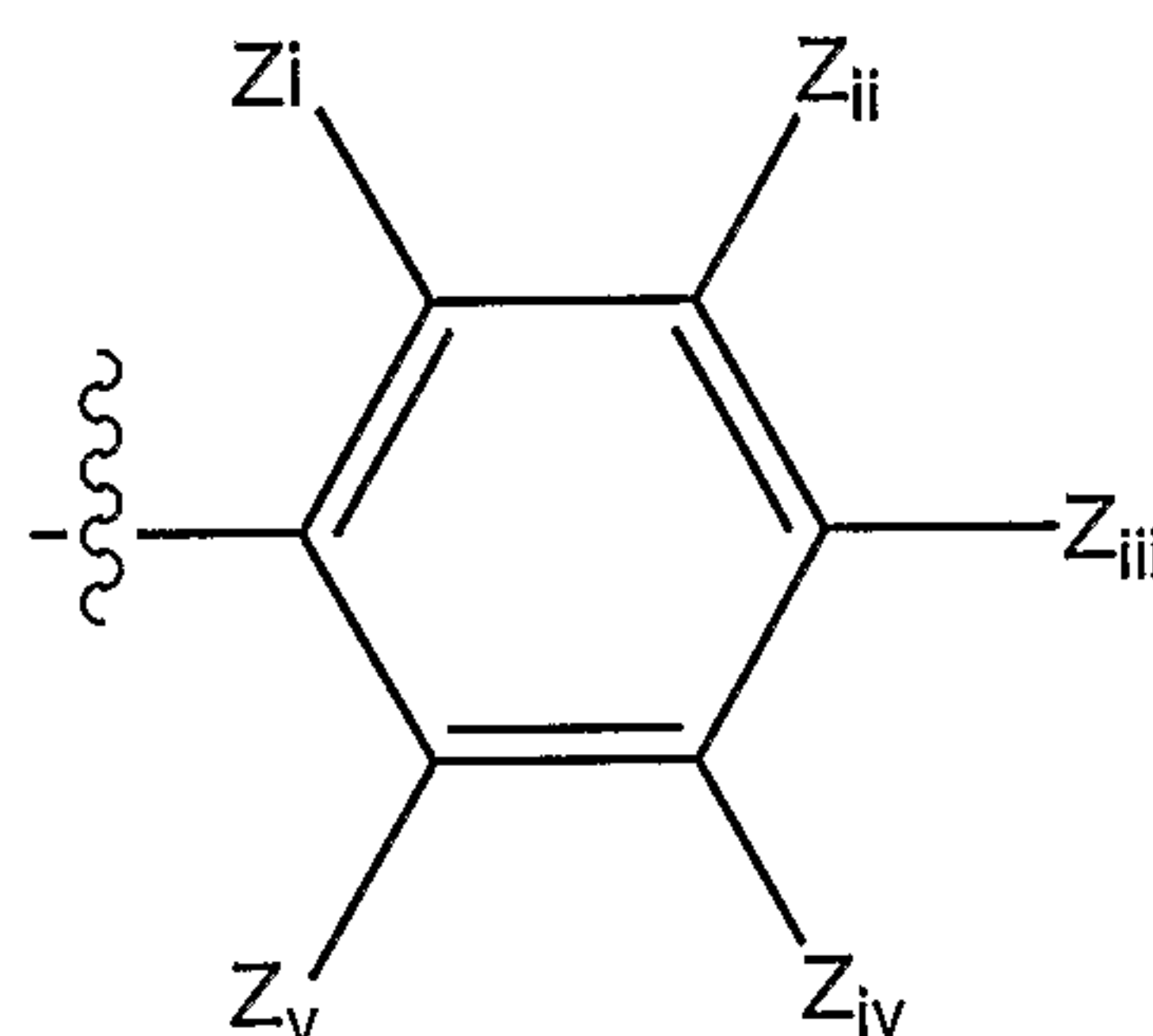
[070] The invention is also directed to useful lipoxin compounds having the
formula (VIII)



wherein X is R_1 , OR_1 , or SR_1 ;

wherein R_1 is

- 5
- (i) a hydrogen atom;
 - (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
 - (iii) a cycloalkyl of 3 to 10 carbon atoms;
 - (iv) an aralkyl of 7 to 12 carbon atoms;
 - (v) phenyl;
 - 10 (vi) substituted phenyl



15

wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or

- (viii) a straight or branched chain alkenyl of 2 to 8
carbon atoms, inclusive;

wherein Q_1 is (C=O), SO_2 or (CN), provided when Q_1 is CN, then X is
absent;

wherein one of R_2 and R_3 is a hydrogen atom and the other is

- (a) H;
- (b) an alkyl of 1 to 8 carbon atoms, inclusive, which may be a
straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms, inclusive;
- (d) an alkenyl of 2 to 8 carbon atoms, inclusive, which may be
straight chain or branched; or
- (e) $R_a Q_2 R_b$ wherein Q_2 is $-O-$ or $-S-$; wherein R_a is alkylene
of 0 to 6 carbon atoms, inclusive, which may be straight
chain or branched and wherein R_b is alkyl of 0 to 8
carbon atoms, inclusive, which may be straight chain or
branched, provided when R_b is 0, then R_b is a hydrogen
atom;

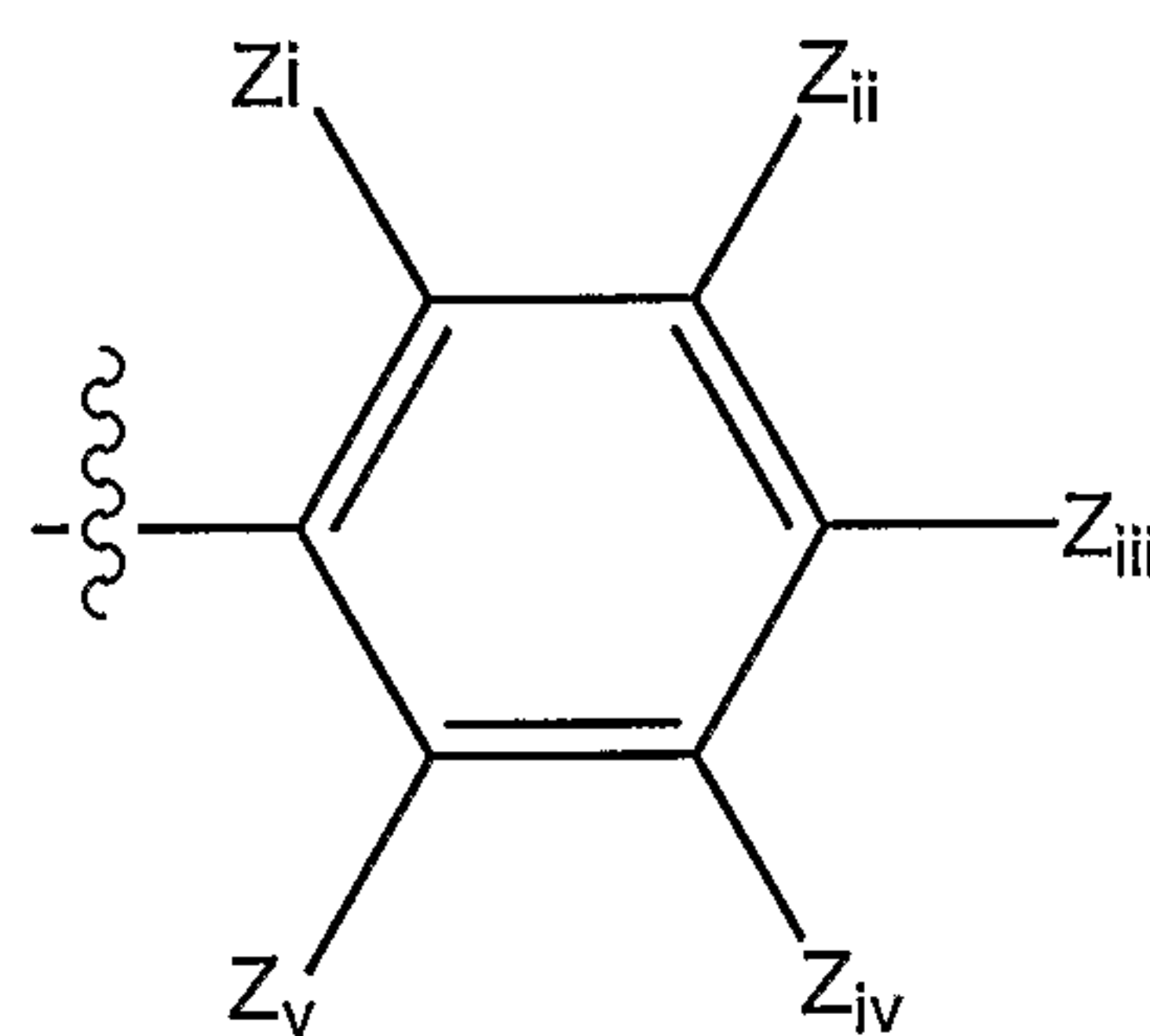
wherein R_4 is

- (a) H;

- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

wherein R_5 is

5



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group;

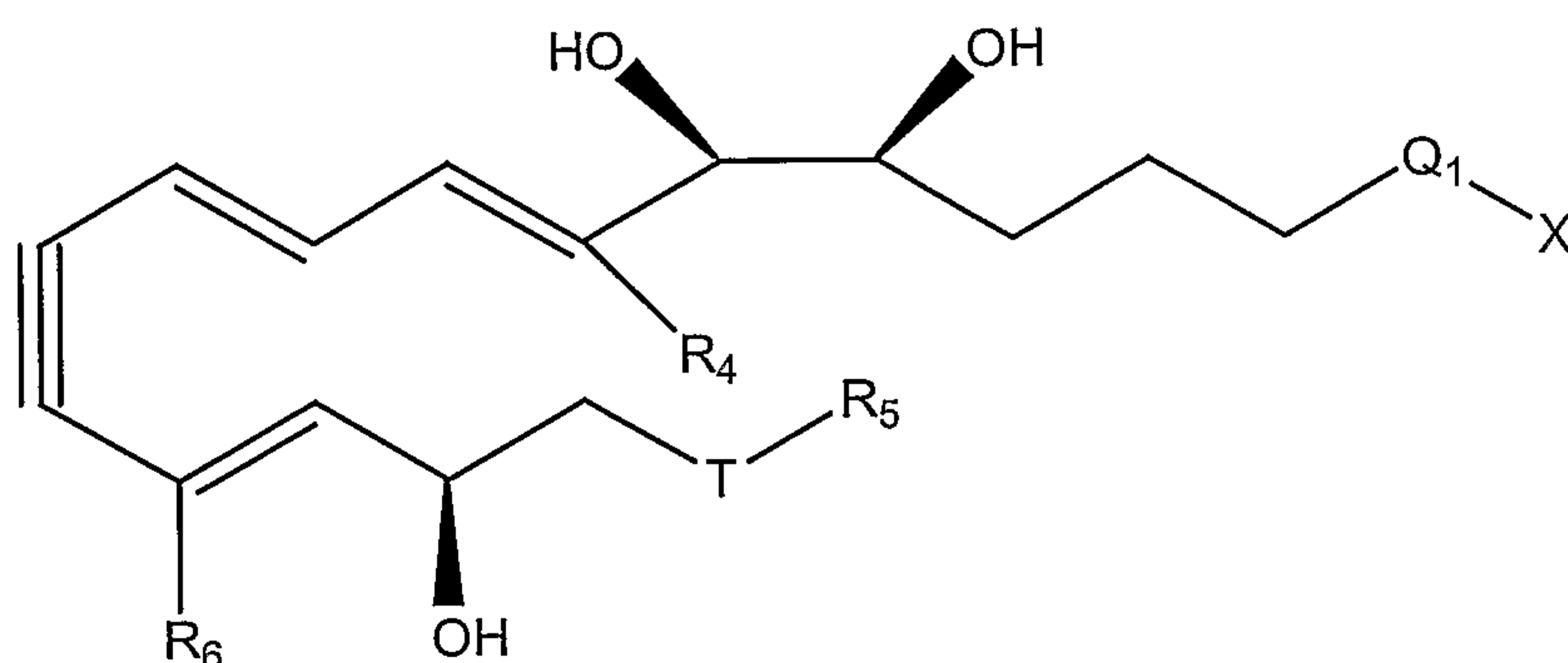
wherein R_6 is

15

- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof.

[071] The invention is further directed to useful lipoxin compounds having the formula (IX)

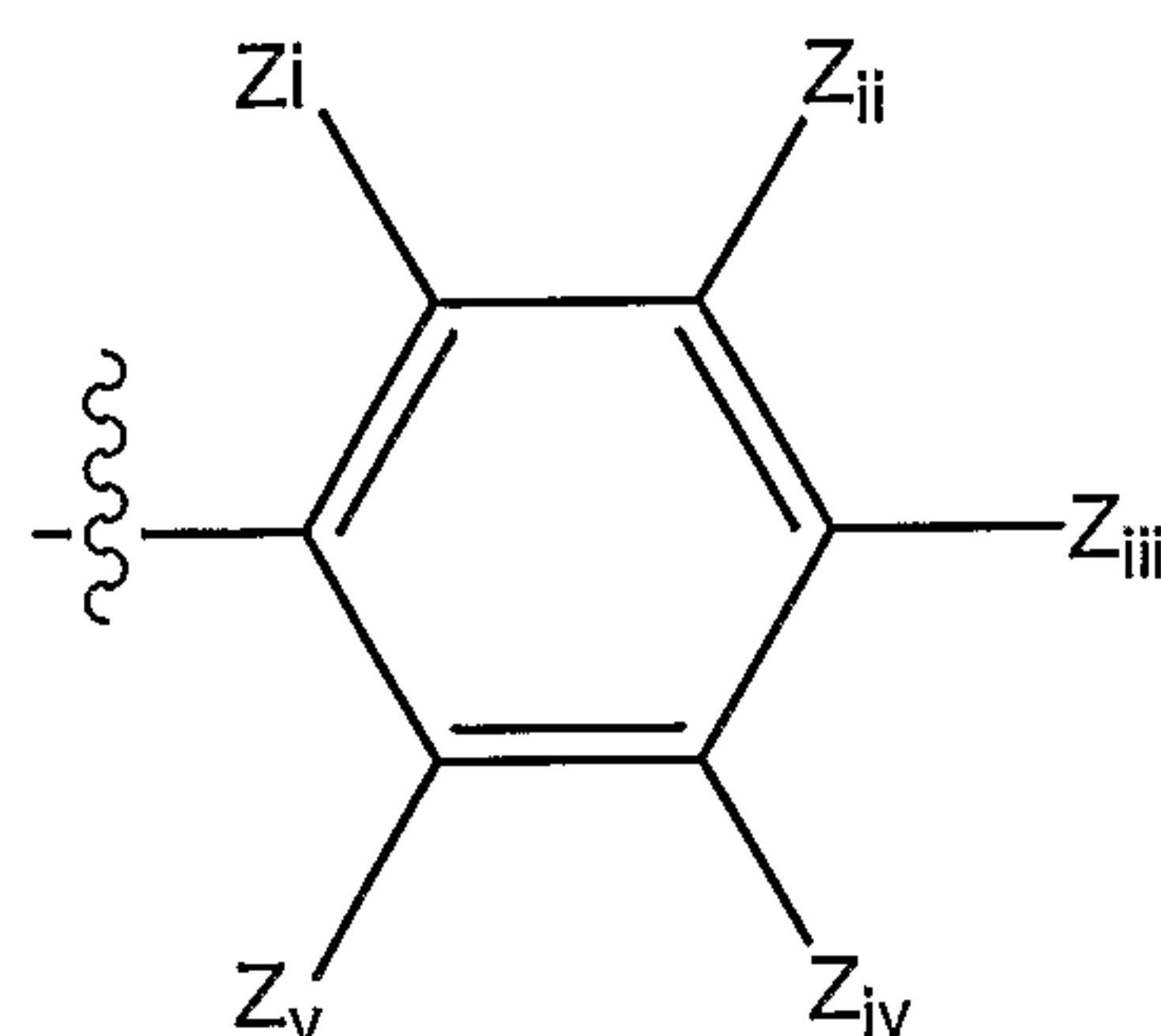


5

wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- | | | |
|----|-------|---|
| 10 | (i) | a hydrogen atom; |
| | (ii) | an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched; |
| | (iii) | a cycloalkyl of 3 to 10 carbon atoms; |
| | (iv) | an aralkyl of 7 to 12 carbon atoms; |
| 15 | (v) | phenyl; |
| | (vi) | substituted phenyl |



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, –
 OR_x , wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a
 5 straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8
 carbon atoms, inclusive;

10

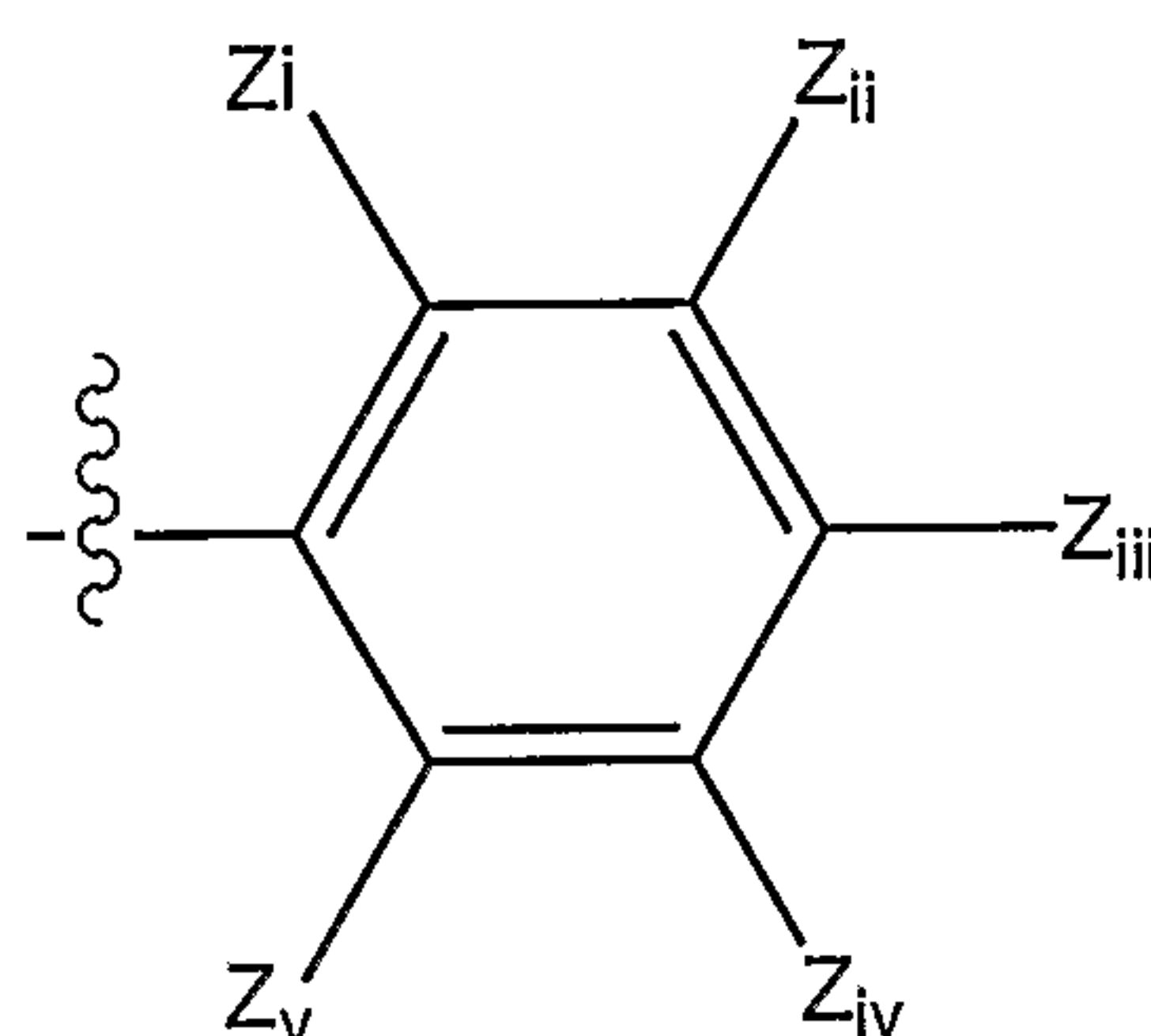
wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is
 absent;

wherein R_4 is

15

- (a) H ;
- (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a
 straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$,
 5 $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is 1 to
 8 carbon atoms, inclusive, which may be a straight chain or branched, and
 hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

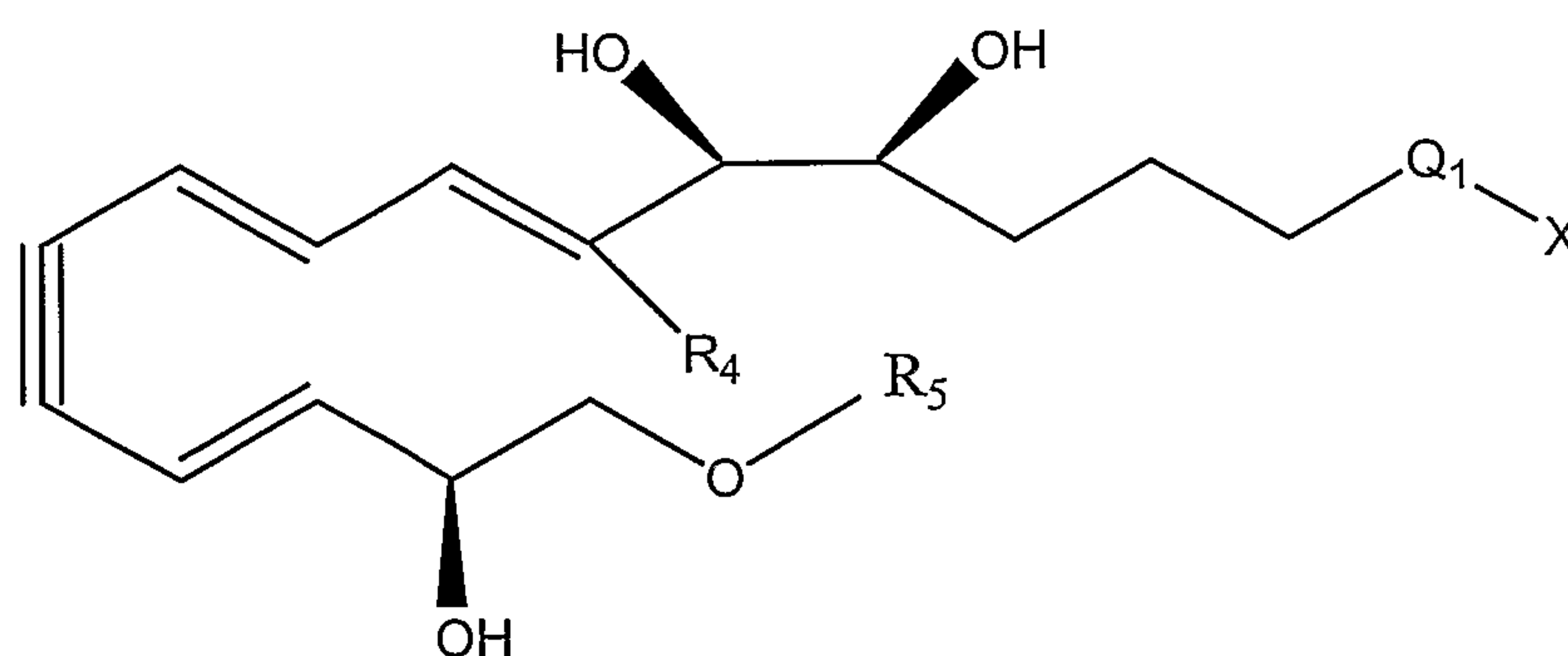
wherein R_6 is

10

- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain
or branched;

15 wherein T is O or S, and pharmaceutically acceptable salts thereof.

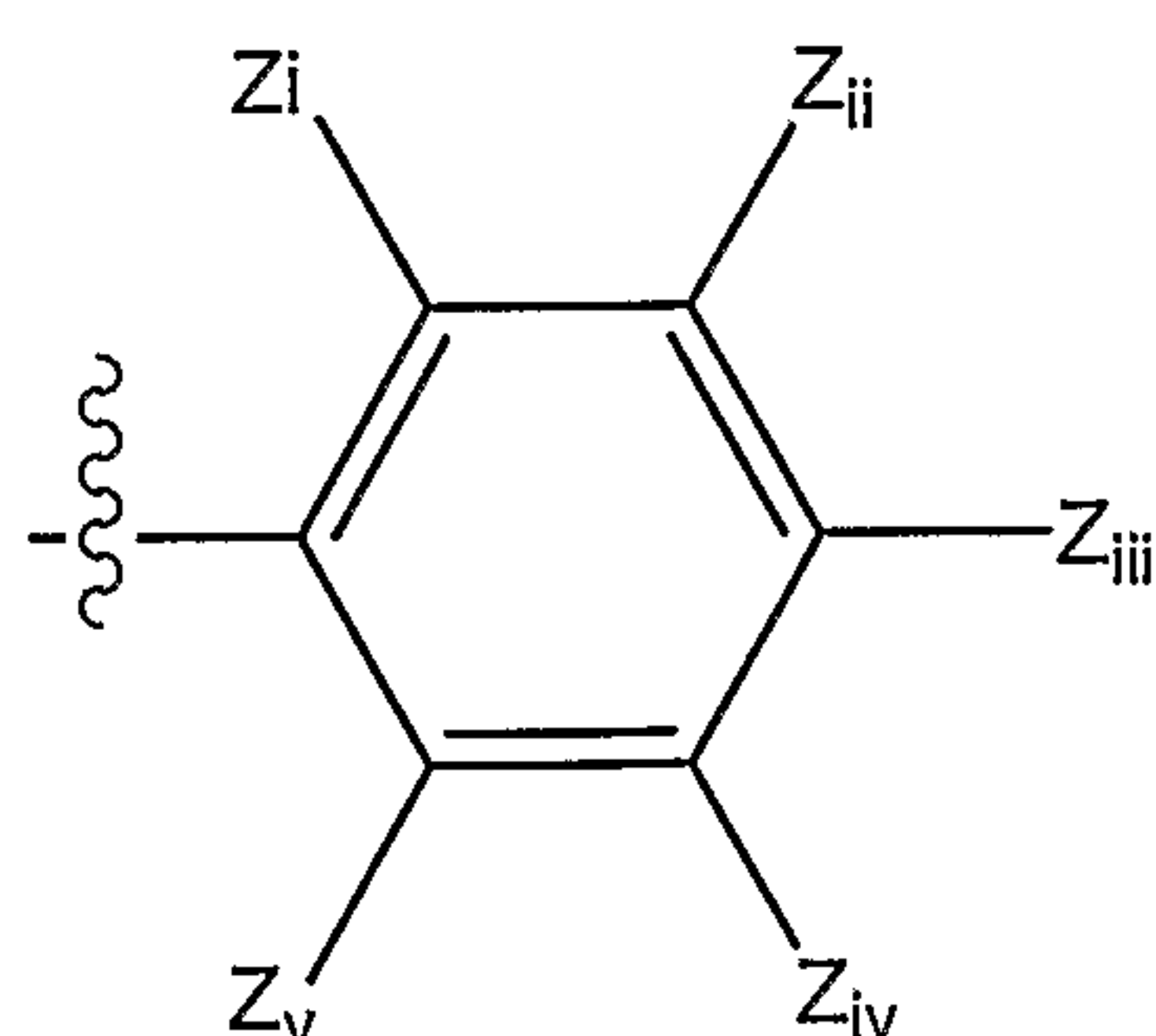
[072] The invention is further directed to useful lipoxin compounds having the
 formula (X)



wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- 5
- (i) a hydrogen atom;
 - (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
which may be straight chain or branched;
 - (iii) a cycloalkyl of 3 to 10 carbon atoms;
 - (iv) an aralkyl of 7 to 12 carbon atoms;
 - 10 (v) phenyl;
 - (vi) substituted phenyl



wherein Z_i, Z_{ii}, Z_{iii}, Z_{iv} and Z_v are each independently selected from –

15 NO₂, –CN, –C(=O)–R₁, –SO₃H, a hydrogen atom, halogen, methyl, –

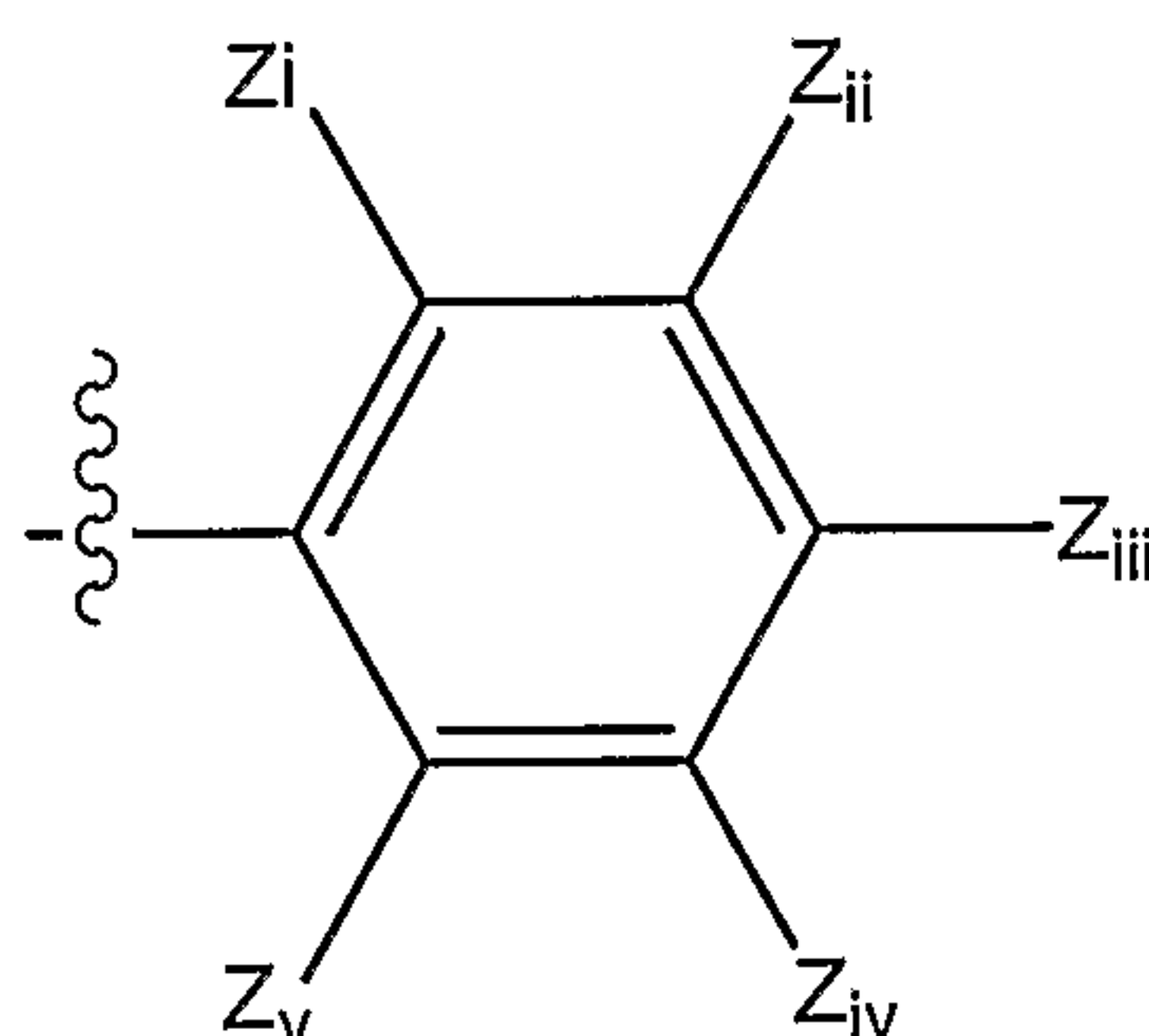
OR_x, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or
- 5 (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms, inclusive;

wherein R₄ is

- (a) H;
- 10 (b) an alkyl of 1 to 6 carbon atoms, inclusive, which may be a straight chain or branched;

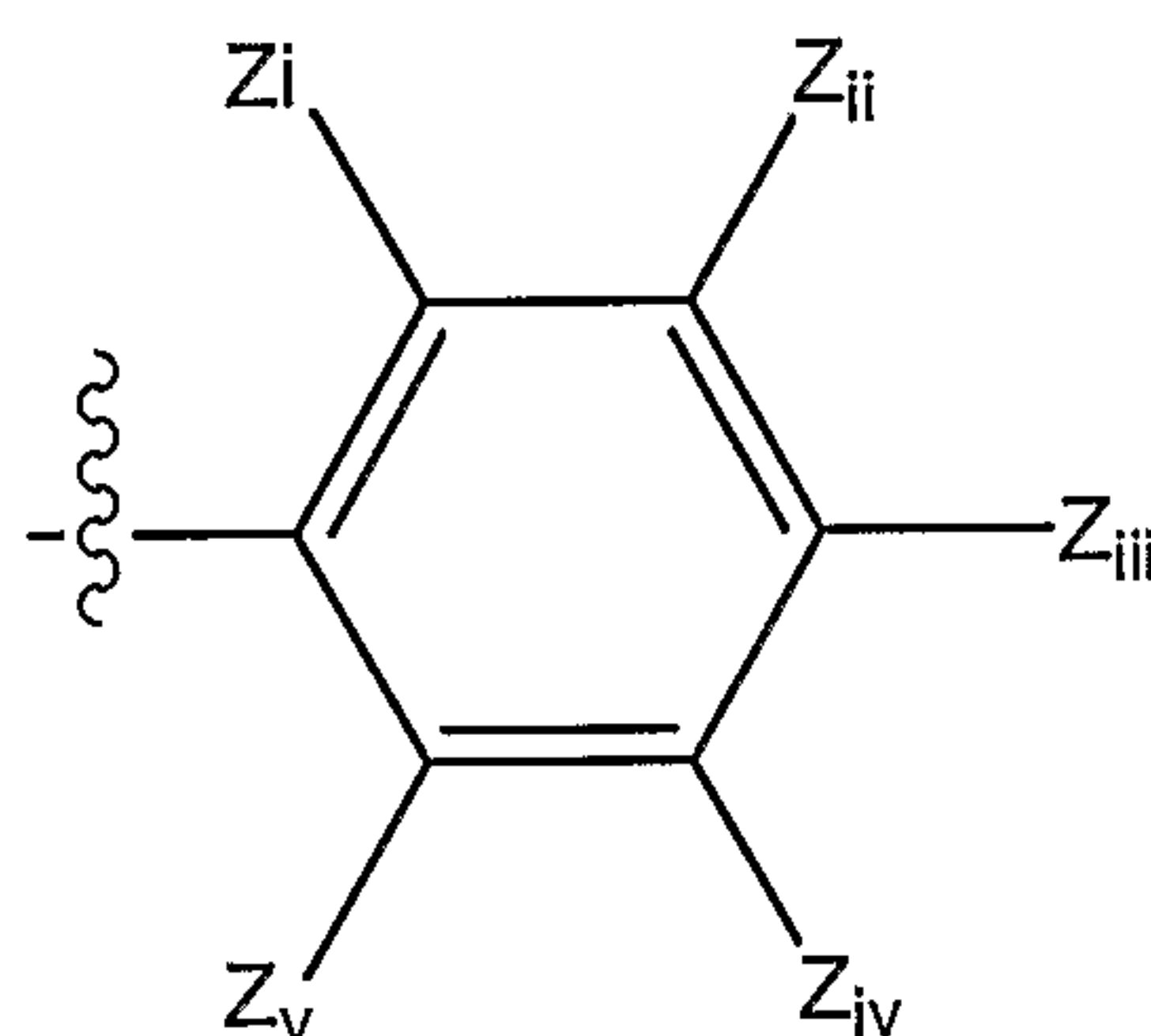
wherein R₅ is



15 wherein Zi, Zii, Ziii, Ziv and Zv are each independently selected from –NO₂, –CN, –C(=O)–R₁, –SO₃H, a hydrogen atom, halogen, methyl, –OR_x, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a

straight chain or branched, and hydroxyl or a substituted or
 unsubstituted, branched or unbranched alkyl group; and
 pharmaceutically acceptable salts thereof.

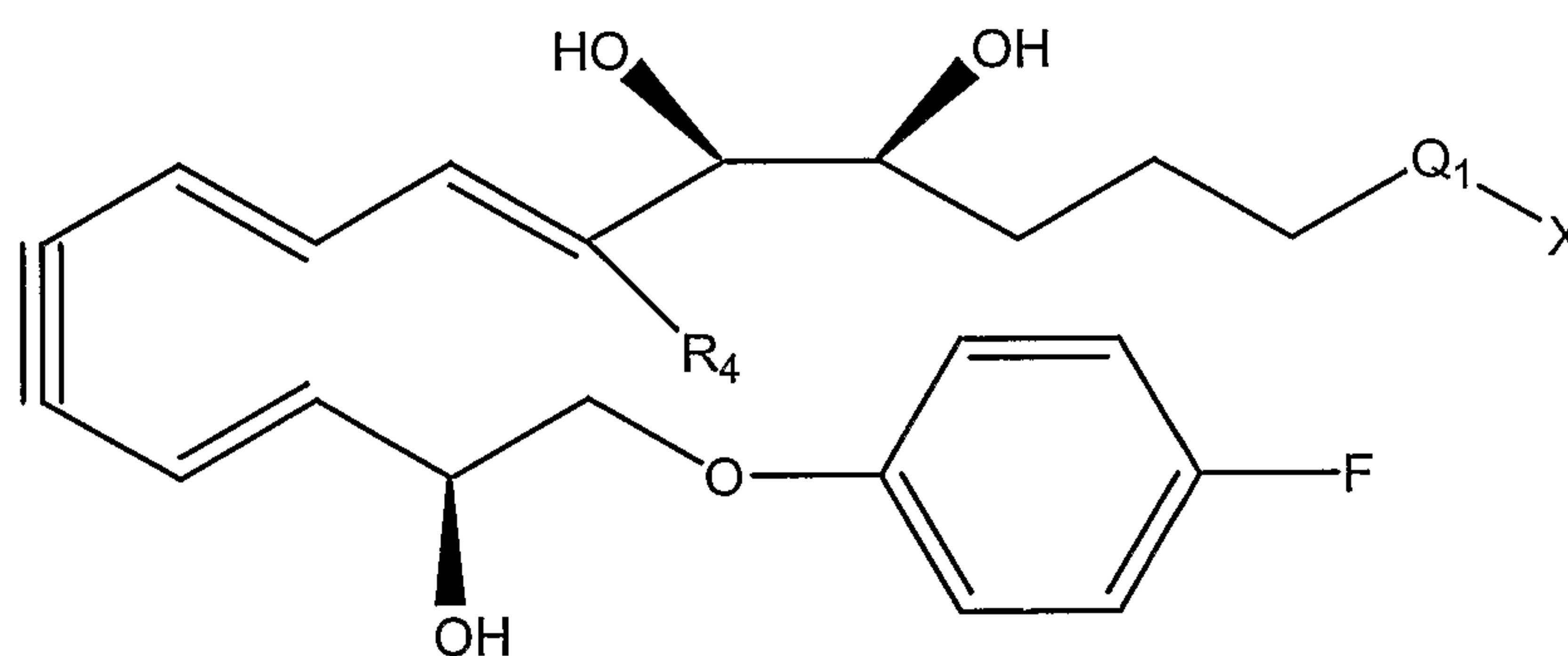
[073] In certain embodiments, X is OR_1 wherein R_1 is a hydrogen atom, an
 5 alkyl group of 1 to 4 carbon atoms or a pharmaceutically acceptable salt, Q_1 is
 $C=O$, R_2 and R_3 , if present, are hydrogen atoms, R_4 is a hydrogen atom or methyl,
 Q_3 and Q_4 , if present, are both O, R_6 , if present, is a hydrogen atom, Y_1 , if
 present, is OH, T is O and R_5 is a substituted phenyl, e.g.,



10

wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from —
 NO_2 , $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$,
 wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain or
 15 branched, and hydroxyl. In certain embodiments for R_5 , para-fluorophenyl
 and/or unsubstituted phenyl acetylenic lipoxins are optimal, e.g., 15-epi-16-(para-
 fluoro)-phenoxy-acetylenic LXA_4 , 16-(para-fluoro)-phenoxy-acetylenic LXA_4 ,
 15-epi-16-phenoxy-acetylenic LXA_4 or 16-phenoxy-acetylenic LXA_4 . The
 compounds encompassed by U.S. Patent 5,650,435 are excluded from certain
 20 aspects of the present invention.

[074] In still another aspect, the present invention is directed to pharmaceutical compositions including compounds having the above-described formulae and a pharmaceutically acceptable carrier. In one embodiment, an acetylenic lipoxin is

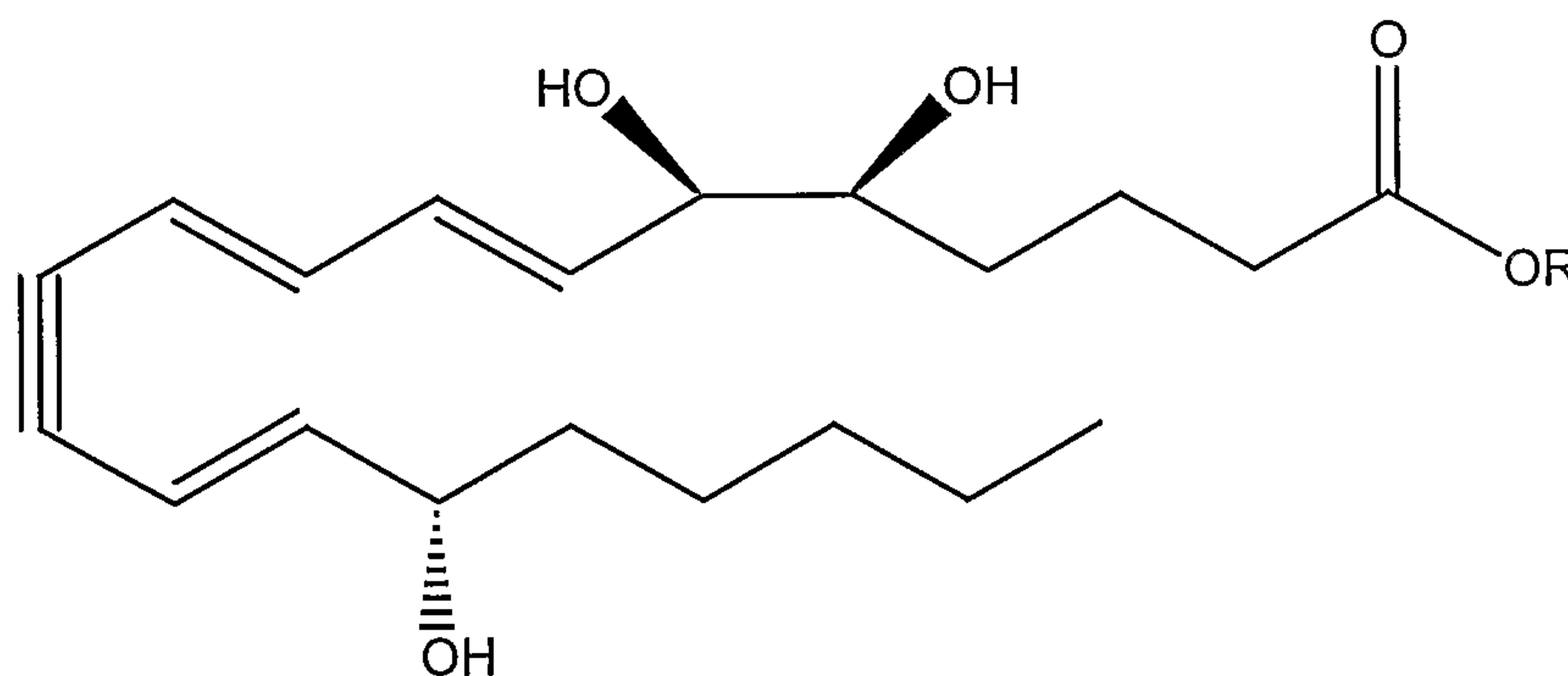


5

[075] In certain embodiments, Y_1 is a hydroxyl and the carbon bearing the hydroxyl can have an R or S configuration. In most embodiments, the chiral carbon bearing the hydroxyl group, e.g., Y_1 , is designated as a 15-epi-lipoxin as is known in the art.

10

[076] In another embodiment, the lipoxin acetylenic analog is (XI)



wherein R is a lower alkyl group, i.e., ethyl, methyl, a hydrogen atom, or a pharmaceutically acceptable salt. Lipoxin acetylenic analog XI had excellent anti-inflammatory properties as evidenced in Figs. 5 and 6.

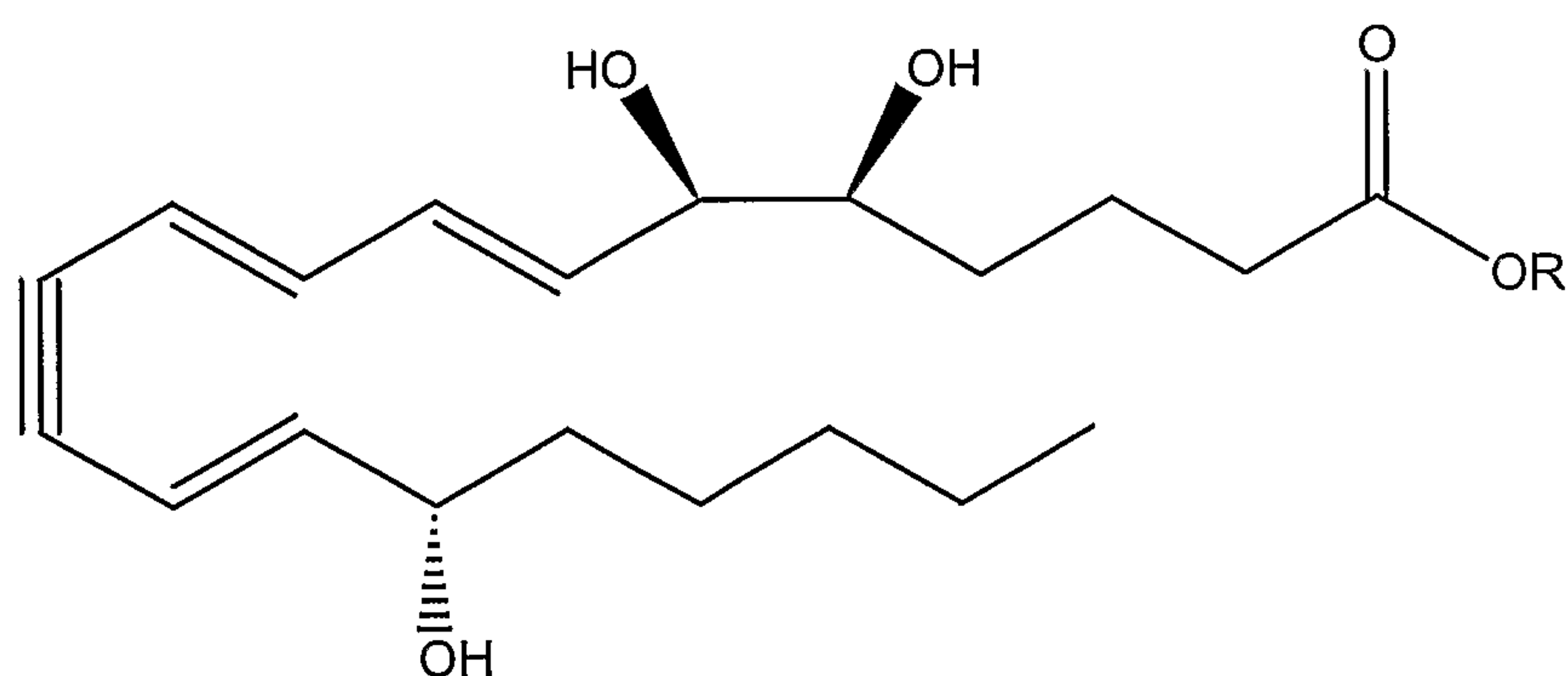
15

[077] In certain embodiments the chirality of the carbons bearing the R_2 , R_3 , Q_3 and Q_4 groups can each independently be either R or S. In certain embodiments, Q_3 and Q_4 have the chiralities shown in structures II, III, IV or V.

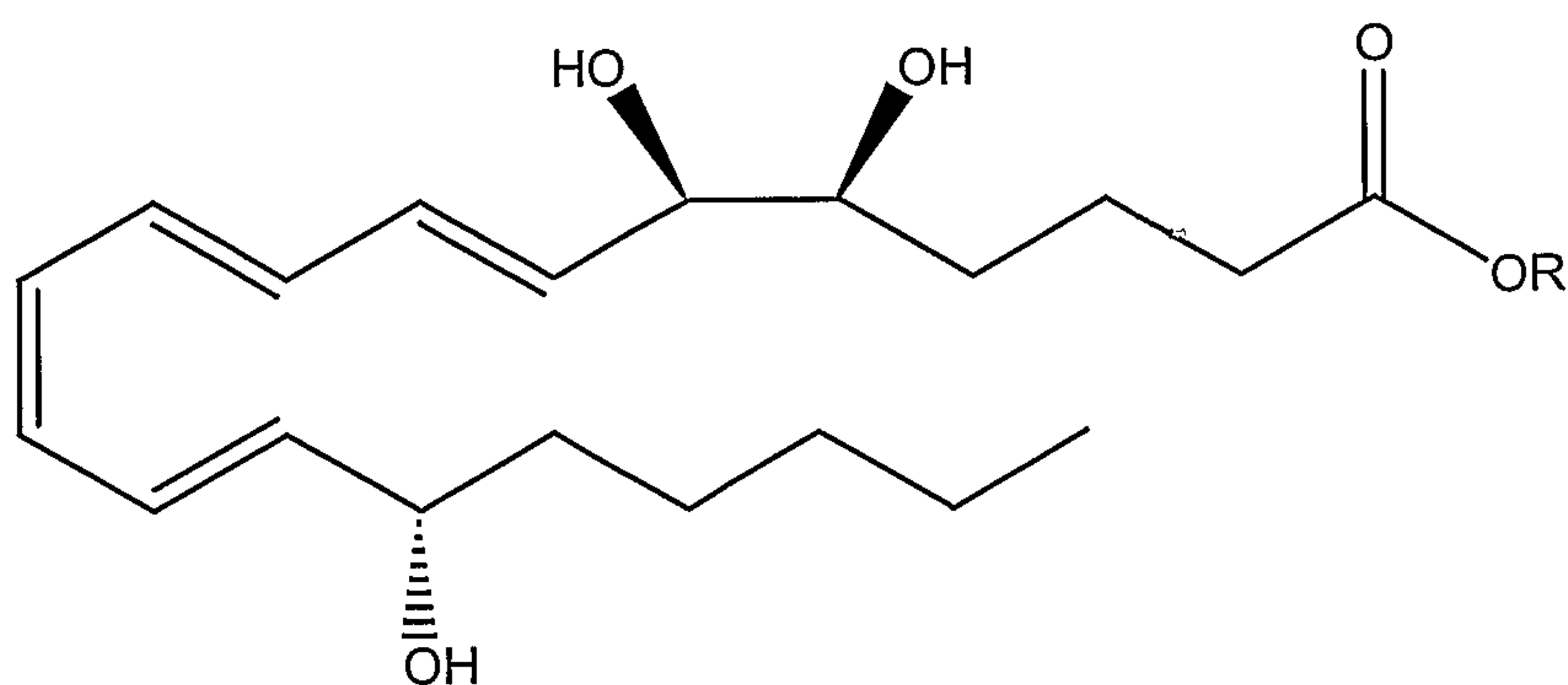
[078] In other embodiments, R_4 is a hydrogen. In still other embodiments, R_6 is a hydrogen.

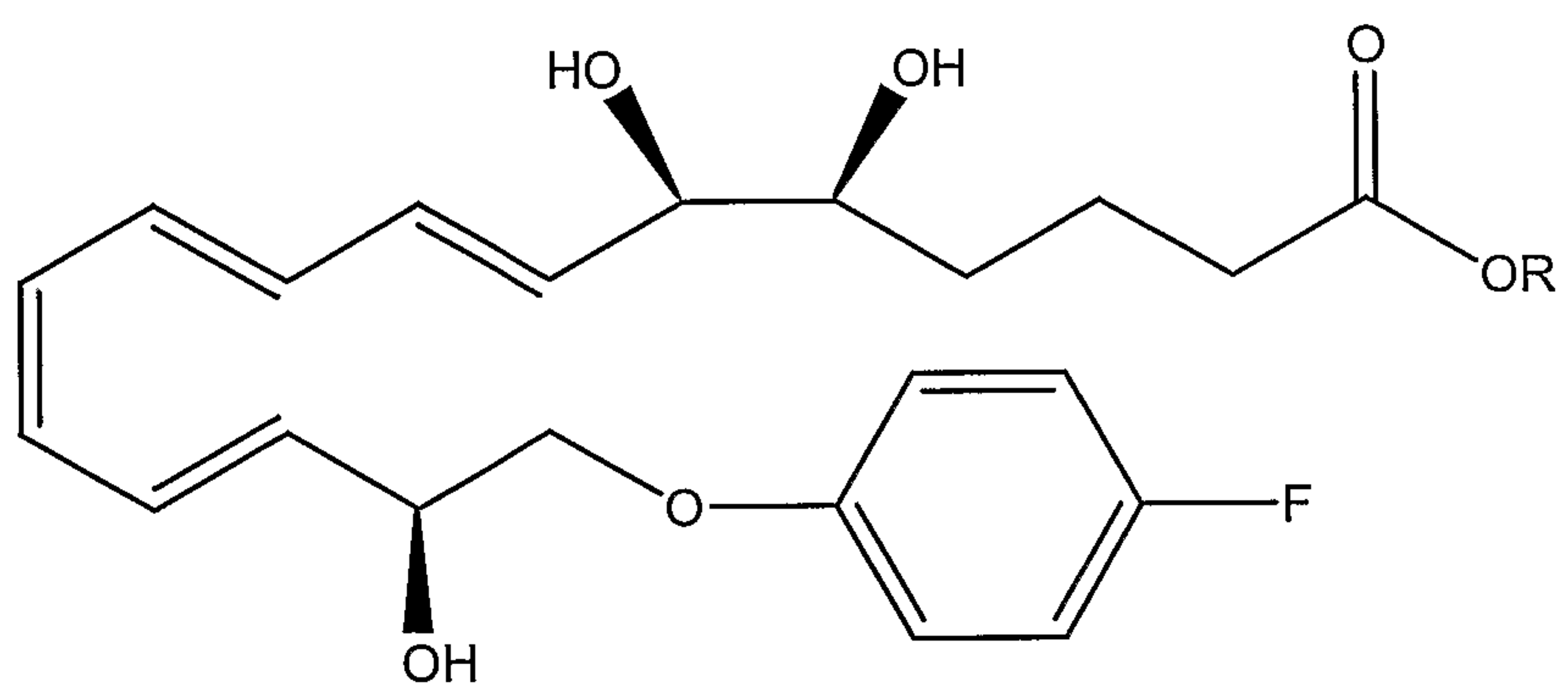
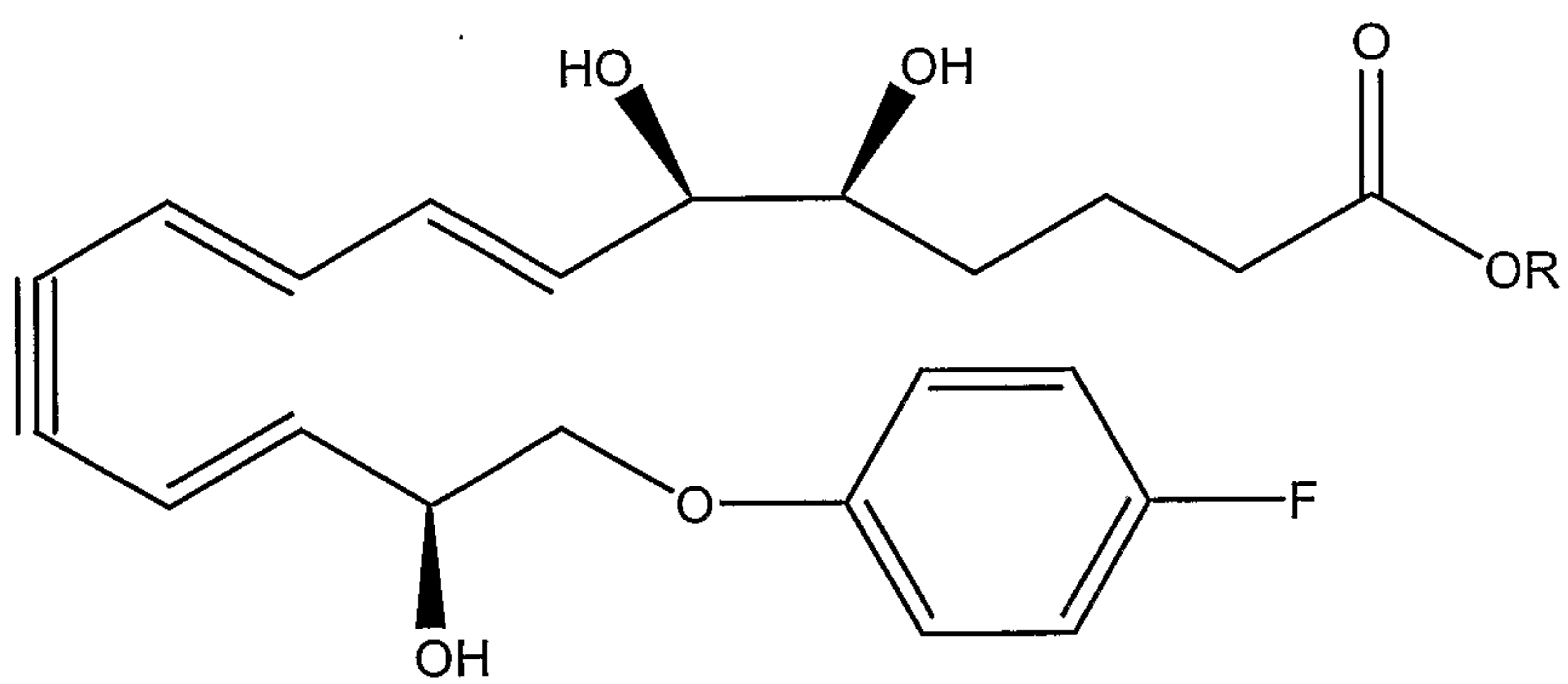
[079] Additionally, R_5 can be a substituted or unsubstituted, branched or unbranched alkyl group having between 1 and about 6 carbon atoms, preferably between 1 and 4 carbon atoms, most preferably between 1 and 3, and preferably one or two carbon atoms. The carbon atoms can have substituents which include halogen atoms, hydroxyl groups, or ether groups.

[080] In particular, the lipoxins and lipoxin analogs useful for the above identified respiratory ailments include, for example:

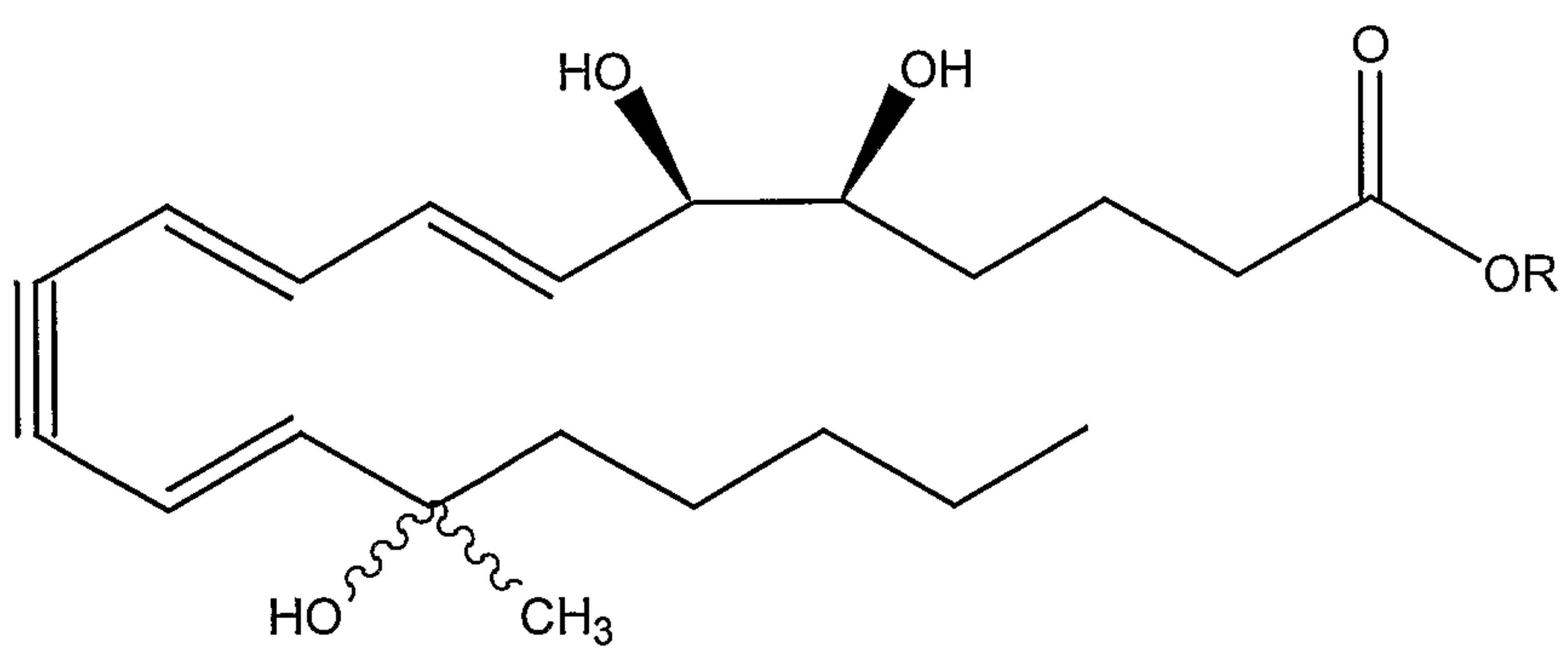


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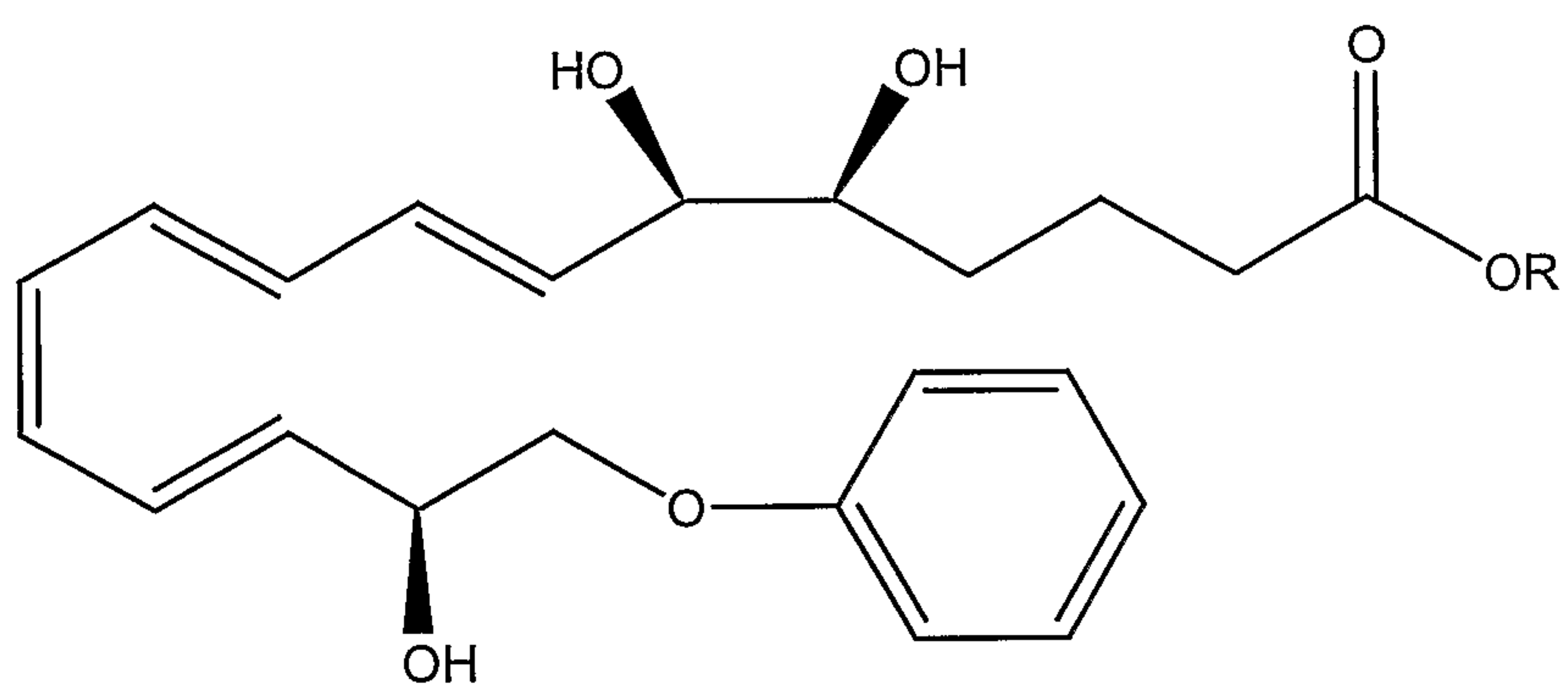
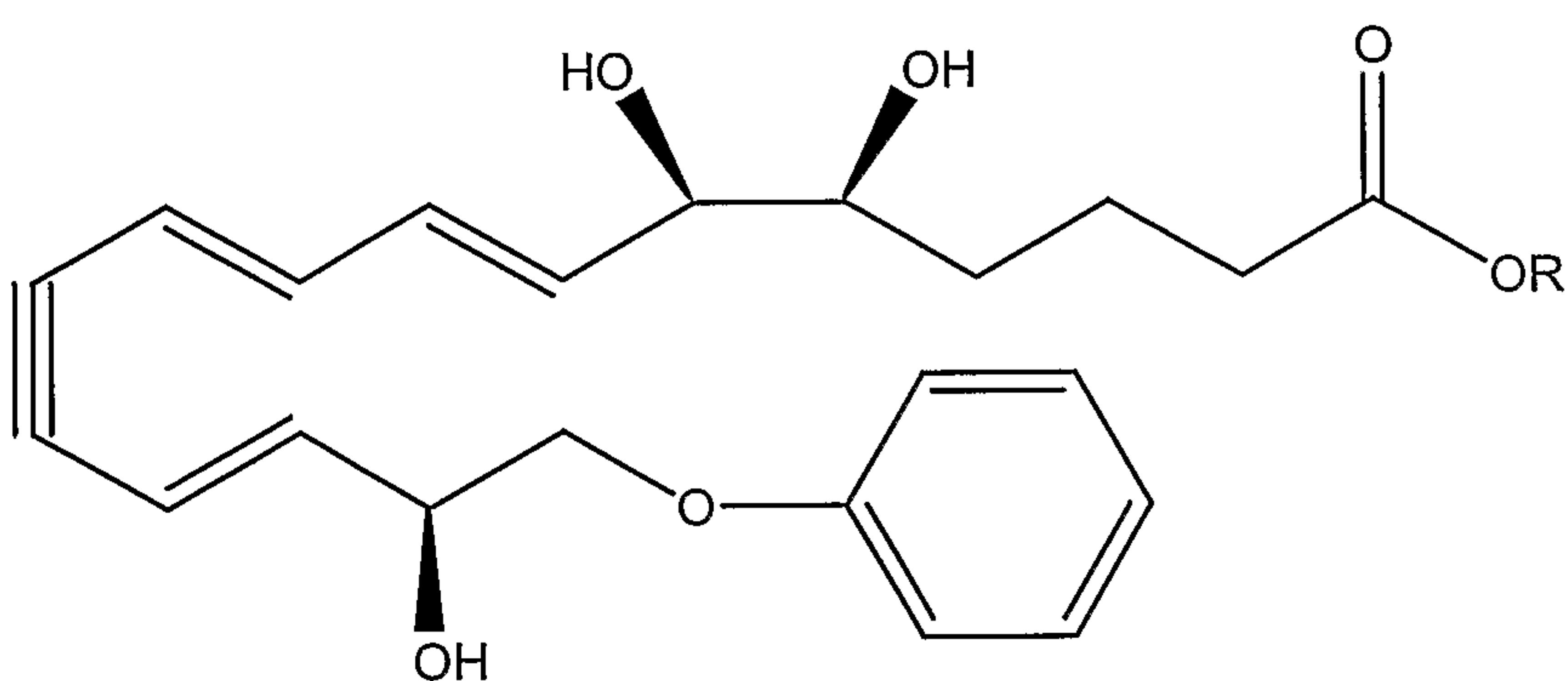
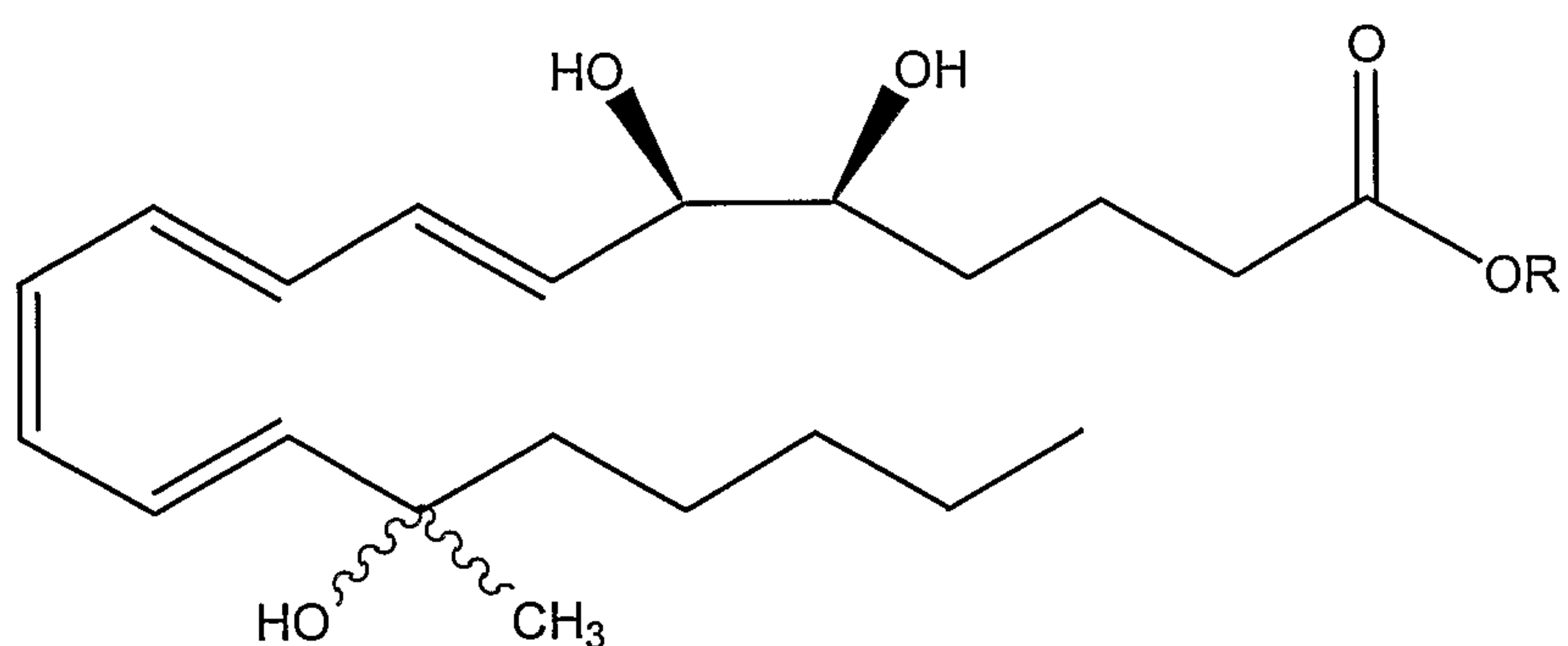




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wherein R is a branched or unbranched alkyl, alkylene, or alkynyl group having 1 to 20 carbon atoms, e.g., a methyl or ethyl group, a hydrogen atom, or a pharmaceutically acceptable salt.

10 **[081]** A “lipoxin analog” shall mean a compound which has an “active region” that functions like the active region of a “natural lipoxin”, but which has a

“metabolic transformation region” that differs from natural lipoxin. Lipoxin analogs include compounds which are structurally similar to a natural lipoxin, compounds which share the same receptor recognition site, compounds which share the same or similar lipoxin metabolic transformation region as lipoxin, and
5 compounds which are art-recognized as being analogs of lipoxin. Lipoxin analogs include lipoxin analog metabolites. The compounds disclosed herein may contain one or more centers of asymmetry. Where asymmetric carbon atoms are present, more than one stereoisomer is possible, and all possible isomeric forms are intended to be included within the structural representations shown.
10 Optically active (R) and (S) isomers may be resolved using conventional techniques known to the ordinarily skilled artisan. The present invention is intended to include the possible diastereoisomers as well as the racemic and optically resolved isomers.

[082] The terms “corresponding lipoxin” and “natural lipoxin” refer to a
15 naturally-occurring lipoxin or lipoxin metabolite. Where an analog has activity for a lipoxin-specific receptor, the corresponding or natural lipoxin is the normal ligand for that receptor. For example, where an analog is a LXA₄ specific receptor on differentiated HL-60 cells, the corresponding lipoxin is LXA₄. Where an analog has activity as an antagonist to another compound (such as
20 leukotriene C₄ and/or leukotriene D₄), which is antagonized by a naturally-occurring lipoxin, that natural lipoxin is the corresponding lipoxin.

[083] “Active region” shall mean the region of a natural lipoxin or lipoxin analog, which is associated with in vivo cellular interactions. The active region may bind the “recognition site” of a cellular lipoxin receptor or a macromolecule
25 or complex of macromolecules, including an enzyme and its cofactor. Lipoxin A₄ analogs of the invention have an active region comprising C₅—C₁₅ of natural lipoxin A₄. Lipoxin B₄ analogs have an active region comprising C₅-C₁₄ of natural lipoxin B₄.

[084] The term “recognition site” or receptor is art-recognized and is intended
30 to refer generally to a functional macromolecule or complex of macromolecules with which certain groups of cellular messengers, such as hormones,

leukotrienes, and lipoxins, must first interact before the biochemical and physiological responses to those messengers are initiated. As used in this application, a receptor may be isolated, on an intact or permeabilized cell, or in tissue, including an organ. A receptor may be from or in a living subject, or it
5 may be cloned. A receptor may normally exist or it may be induced by a disease state, by an injury, or by artificial means. A compound of this invention may bind reversibly, irreversibly, competitively, noncompetitively, or uncompetitively with respect to the natural substrate of a recognition site.

[085] The term “metabolic transformation region” is intended to refer generally
10 to that portion of a lipoxin, a lipoxin metabolite, or lipoxin analog including a lipoxin analog metabolite, upon which an enzyme or an enzyme and its cofactor attempts to perform one or more metabolic transformations which that enzyme or enzyme and cofactor normally transform on lipoxins. The metabolic transformation region may or may not be susceptible to the transformation. A
15 nonlimiting example of a metabolic transformation region of a lipoxin is a portion of LXA₄ that includes the C-13,14 double bond or the C-15 hydroxyl group, or both.

[086] The term “detectable label molecule” is meant to include fluorescent, phosphorescent, and radiolabeled molecules used to trace, track, or identify the
20 compound or receptor recognition site to which the detectable label molecule is bound. The label molecule may be detected by any of the several methods known in the art.

[087] The term “labeled lipoxin analog” is further understood to encompass compounds which are labeled with radioactive isotopes, such as but not limited to
25 tritium (³H), deuterium (²H), carbon (¹⁴C), or otherwise labeled (e.g. fluorescently). The compounds of this invention may be labeled or derivatized, for example, for kinetic binding experiments, for further elucidating metabolic pathways and enzymatic mechanisms, or for characterization by methods known in the art of analytical chemistry.

30 **[088]** The term “inhibits metabolism” means the blocking or reduction of activity of an enzyme which metabolizes a native lipoxin. The blockage or

reduction may occur by covalent bonding, by irreversible binding, by reversible binding which has a practical effect of irreversible binding, or by any other means which prevents the enzyme from operating in its usual manner on another lipoxin analog, including a lipoxin analog metabolite, a lipoxin, or a lipoxin
5 metabolite.

[089] The term “resists metabolism” is meant to include failing to undergo one or more of the metabolic degradative transformations by at least one of the enzymes which metabolize lipoxins. Two nonlimiting examples of LXA₄ analog that resists metabolism are 1) a structure which can not be oxidized to the 15-oxo
10 form, and 2) a structure which may be oxidized to the 15-oxo form, but is not susceptible to enzymatic reduction to the 13,14-dihydro form.

[090] The term “more slowly undergoes metabolism” means having slower reaction kinetics, or requiring more time for the completion of the series of metabolic transformations by one or more of the enzymes which metabolize
15 lipoxin. A nonlimiting example of a LXA₄ analog which more slowly undergoes metabolism is a structure which has a higher transition state energy for C-15 dehydrogenation than does LXA₄ because the analog is sterically hindered at the C-16.

[091] The term “tissue” is intended to include intact cells, blood, blood
20 preparations such as plasma and serum, bones, joints, muscles, smooth muscles, and organs.

[092] The term “halogen” is meant to include fluorine, chlorine, bromine and iodine, or fluoro, chloro, bromo, and iodo. In certain aspects, the compounds of the invention do not include halogenated compounds, e.g., fluorinated
25 compounds.

[093] The term “subject” is intended to include living organisms susceptible to conditions or diseases caused or contributed to by inflammation, asthma, asthma related disorders of the respiratory tract and lung, or disorders of the respiratory tract and lung as generally disclosed, but not limited to, throughout this
30 specification. Examples of subjects include humans, dogs, cats, cows, goats, and mice. The term subject is further intended to include transgenic species.

[094] When the compounds of the present invention are administered as pharmaceuticals, to humans and mammals, they can be given per se or as a pharmaceutical composition containing, for example, 0.1 to 99.5% (more preferably, 0.5 to 90%) of active ingredient in combination with a
5 pharmaceutically acceptable carrier.

[095] The phrase "pharmaceutically acceptable carrier" as used herein means a pharmaceutically acceptable material, composition or vehicle, such as a liquid or solid filler, diluent, excipient, solvent or encapsulating material, involved in carrying or transporting a compound(s) of the present invention within or to the
10 subject such that it can perform its intended function. Typically, such compounds are carried or transported from one organ, or portion of the body, to another organ, or portion of the body. Each carrier must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not injurious to the patient. Some examples of materials which can serve as
15 pharmaceutically acceptable carriers include: sugars, such as lactose, glucose and sucrose; starches, such as corn starch and potato starch; cellulose, and its derivatives, such as sodium carboxymethyl cellulose, ethyl cellulose and cellulose acetate; powdered tragacanth; malt; gelatin; talc; excipients, such as cocoa butter and suppository waxes; oils, such as peanut oil, cottonseed oil,
20 safflower oil, sesame oil, olive oil, corn oil and soybean oil; glycols, such as propylene glycol; polyols, such as glycerin, sorbitol, mannitol and polyethylene glycol; esters, such as ethyl oleate and ethyl laurate; agar; buffering agents, such as magnesium hydroxide and aluminum hydroxide; alginic acid; pyrogen-free water; isotonic saline; Ringer's solution; ethyl alcohol; phosphate buffer
25 solutions; and other non-toxic compatible substances employed in pharmaceutical formulations.

[096] In certain embodiment, the compounds of the present invention may contain one or more acidic functional groups and, thus, are capable of forming pharmaceutically acceptable salts with pharmaceutically acceptable bases. The
30 term "pharmaceutically acceptable salts, esters, amides, and prodrugs" as used herein refers to those carboxylate salts, amino acid addition salts, esters, amides,

and prodrugs of the compounds of the present invention which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of patients without undue toxicity, irritation, allergic response, and the like, commensurate with a reasonable benefit/risk ratio, and effective for their intended use of the compounds of the invention. The term "salts" refers to the relatively non-toxic, inorganic and organic acid addition salts of compounds of the present invention. These salts can be prepared *in situ* during the final isolation and purification of the compounds or by separately reacting the purified compound in its free base form with a suitable organic or inorganic acid and isolating the salt thus formed. These may include cations based on the alkali and alkaline earth metals, such as sodium, lithium, potassium, calcium, magnesium and the like, as well as non-toxic ammonium, quaternary ammonium, and amine cations including, but not limited to ammonium, tetramethylammonium, tetraethylammonium, methylamine, dimethylamine, trimethylamine, triethylamine, ethylamine, and the like. (See, for example, Berge S. M., et al., "Pharmaceutical Salts," J. Pharm. Sci., 1977;66:1-19).

[097] The term "pharmaceutically acceptable esters" refers to the relatively non-toxic, esterified products of the compounds of the present invention. These esters can be prepared *in situ* during the final isolation and purification of the compounds, or by separately reacting the purified compound in its free acid form or hydroxyl with a suitable esterifying agent. Carboxylic acids can be converted into esters *via* treatment with an alcohol in the presence of a catalyst. The term is further intended to include lower hydrocarbon groups capable of being solvated under physiological conditions, e.g., alkyl esters, methyl, ethyl and propyl esters. In certain embodiments, the ester is not a methyl ester (See, for example, Berge et al., *supra*).

[098] Wetting agents, emulsifiers and lubricants, such as sodium lauryl sulfate and magnesium stearate, as well as coloring agents, release agents, coating agents, sweetening, flavoring and perfuming agents, preservatives and antioxidants can also be present in the compositions.

[099] Examples of pharmaceutically acceptable antioxidants include: water soluble antioxidants, such as ascorbic acid, cysteine hydrochloride, sodium bisulfate, sodium metabisulfite, sodium sulfite and the like; oil-soluble antioxidants, such as ascorbyl palmitate, butylated hydroxyanisole (BHA),
5 butylated hydroxytoluene (BHT), lecithin, propyl gallate, alpha-tocopherol, and the like; and metal chelating agents, such as citric acid, ethylenediamine tetraacetic acid (EDTA), sorbitol, tartaric acid, phosphoric acid, and the like.

[0100] Formulations of the present invention include those suitable for intravenous, oral, nasal, topical, transdermal, buccal, sublingual, rectal, vaginal
10 and/or parenteral administration. The formulations may conveniently be presented in unit dosage form and may be prepared by any methods well known in the art of pharmacy. The amount of active ingredient which can be combined with a carrier material to produce a single dosage form will generally be that amount of the compound which produces a therapeutic effect. Generally, out of
15 one hundred per cent, this amount will range from about 1 per cent to about ninety-nine percent of active ingredient, preferably from about 5 per cent to about 70 per cent, most preferably from about 10 per cent to about 30 per cent.

[0101] Methods of preparing these formulations or compositions include the step of bringing into association a compound of the present invention with the carrier
20 and, optionally, one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association a compound of the present invention with liquid carriers, or finely divided solid carriers, or both, and then, if necessary, shaping the product.

[0102] Formulations of the invention suitable for oral administration may be in
25 the form of capsules, cachets, pills, tablets, lozenges (using a flavored basis, usually sucrose and acacia or tragacanth), powders, granules, or as a solution or a suspension in an aqueous or non-aqueous liquid, or as an oil-in-water or water-in-oil liquid emulsion, or as an elixir or syrup, or as pastilles (using an inert base, such as gelatin and glycerin, or sucrose and acacia) and/or as mouth washes and
30 the like, each containing a predetermined amount of a compound of the present

invention as an active ingredient. A compound of the present invention may also be administered as a bolus, electuary or paste.

[0103] In solid dosage forms of the invention for oral administration (capsules, tablets, pills, dragees, powders, granules and the like), the active ingredient is mixed with one or more pharmaceutically acceptable carriers, such as sodium citrate or dicalcium phosphate, and/or any of the following: fillers or extenders, such as starches, lactose, sucrose, glucose, mannitol, and/or silicic acid; binders, such as, for example, carboxymethylcellulose, alginates, gelatin, polyvinyl pyrrolidone, sucrose and/or acacia; humectants, such as glycerol; disintegrating agents, such as agar-agar, calcium carbonate, potato or tapioca starch, alginic acid, certain silicates, and sodium carbonate; solution retarding agents, such as paraffin; absorption accelerators, such as quaternary ammonium compounds; wetting agents, such as, for example, cetyl alcohol and glycerol monostearate; absorbents, such as kaolin and bentonite clay; lubricants, such as talc, calcium stearate, magnesium stearate, solid polyethylene glycols, sodium lauryl sulfate, and mixtures thereof; and coloring agents. In the case of capsules, tablets and pills, the pharmaceutical compositions may also comprise buffering agents. Solid compositions of a similar type may also be employed as fillers in soft and hard-filled gelatin capsules using such excipients as lactose or milk sugars, as well as high molecular weight polyethylene glycols and the like.

[0104] A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared using binder (for example, gelatin or hydroxypropylmethyl cellulose), lubricant, inert diluent, preservative, disintegrant (for example, sodium starch glycolate or cross-linked sodium carboxymethyl cellulose), surface-active or dispersing agent. Molded tablets may be made by molding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent.

[0105] The tablets, and other solid dosage forms of the pharmaceutical compositions of the present invention, such as dragees, capsules, pills and granules, may optionally be scored or prepared with coatings and shells, such as enteric coatings and other coatings well known in the pharmaceutical-formulating

art. They may also be formulated so as to provide slow or controlled release of the active ingredient therein using, for example, hydroxypropylmethyl cellulose in varying proportions to provide the desired release profile, other polymer matrices, liposomes and/or microspheres. They may be sterilized by, for example, filtration through a bacteria-retaining filter, or by incorporating sterilizing agents in the form of sterile solid compositions which can be dissolved in sterile water, or some other sterile injectable medium immediately before use. These compositions may also optionally contain opacifying agents and may be of a composition that they release the active ingredient(s) only, or preferentially, in a certain portion of the gastrointestinal tract, optionally, in a delayed manner. Examples of embedding compositions which can be used include polymeric substances and waxes. The active ingredient can also be in micro-encapsulated form, if appropriate, with one or more of the above-described excipients.

[0106] Liquid dosage forms for oral administration of the compounds of the invention include pharmaceutically acceptable emulsions, microemulsions, solutions, suspensions, syrups and elixirs. In addition to the active ingredient, the liquid dosage forms may contain inert diluents commonly used in the art, such as, for example, water or other solvents, solubilizing agents and emulsifiers, such as ethyl alcohol, isopropyl alcohol, ethyl carbonate, ethyl acetate, benzyl alcohol, benzyl benzoate, propylene glycol, 1,3-butylene glycol, oils (in particular, cottonseed, groundnut, corn, germ, olive, castor and sesame oils), glycerol, tetrahydrofuryl alcohol, polyethylene glycols and fatty acid esters of sorbitan, and mixtures thereof.

[0107] Besides inert diluents, the oral compositions can also include adjuvants such as wetting agents, emulsifying and suspending agents, sweetening, flavoring, coloring, perfuming and preservative agents.

[0108] Suspensions, in addition to the active compounds, may contain suspending agents as, for example, ethoxylated isostearyl alcohols, polyoxyethylene sorbitol and sorbitan esters, microcrystalline cellulose, aluminum metahydroxide, bentonite, agar-agar and tragacanth, and mixtures thereof.

[0109] Formulations of the pharmaceutical compositions of the invention for rectal or vaginal administration may be presented as a suppository, which may be prepared by mixing one or more compounds of the invention with one or more suitable nonirritating excipients or carriers comprising, for example, cocoa butter, polyethylene glycol, a suppository wax or a salicylate, and which is solid at room temperature, but liquid at body temperature and, therefore, will melt in the rectum or vaginal cavity and release the active compound.

[0110] Formulations of the present invention which are suitable for vaginal administration also include pessaries, tampons, creams, gels, pastes, foams or spray formulations containing such carriers as are known in the art to be appropriate.

[0111] Dosage forms for the topical or transdermal administration of a compound of this invention include powders, sprays, ointments, pastes, creams, lotions, gels, solutions, patches and inhalants. The active compound may be mixed under sterile conditions with a pharmaceutically acceptable carrier, and with any preservatives, buffers, or propellants which may be required.

[0112] The ointments, pastes, creams and gels may contain, in addition to an active compound of this invention, excipients, such as animal and vegetable fats, oils, waxes, paraffins, starch, tragacanth, cellulose derivatives, polyethylene glycols, silicones, bentonites, silicic acid, talc and zinc oxide, or mixtures thereof.

[0113] Powders and sprays can contain, in addition to a compound of this invention, excipients such as lactose, talc, silicic acid, aluminum hydroxide, calcium silicates and polyamide powder, or mixtures of these substances. Sprays can additionally contain customary propellants, such as chlorofluorohydrocarbons and volatile unsubstituted hydrocarbons, such as butane and propane.

[0114] Transdermal patches have the added advantage of providing controlled delivery of a compound of the present invention to the body. Such dosage forms can be made by dissolving or dispersing the compound in the proper medium. Absorption enhancers can also be used to increase the flux of the compound across the skin. The rate of such flux can be controlled by either providing a rate

controlling membrane or dispersing the active compound in a polymer matrix or gel.

[0115] Ophthalmic formulations, eye ointments, powders, solutions and the like, are also contemplated as being within the scope of this invention.

5 **[0116]** Pharmaceutical compositions of this invention suitable for parenteral administration comprise one or more compounds of the invention in combination with one or more pharmaceutically acceptable sterile isotonic aqueous or nonaqueous solutions, dispersions, suspensions or emulsions, or sterile powders which may be reconstituted into sterile injectable solutions or dispersions just
10 prior to use, which may contain antioxidants, buffers, bacteriostats, solutes which render the formulation isotonic with the blood of the intended recipient or suspending or thickening agents.

[0117] Examples of suitable aqueous and nonaqueous carriers which may be employed in the pharmaceutical compositions of the invention include water,
15 ethanol, polyols (such as glycerol, propylene glycol, polyethylene glycol, and the like), and suitable mixtures thereof, vegetable oils, such as olive oil, and injectable organic esters, such as ethyl oleate. Proper fluidity can be maintained, for example, by the use of coating materials, such as lecithin, by the maintenance of the required particle size in the case of dispersions, and by the use of
20 surfactants.

[0118] These compositions may also contain adjuvants such as preservatives, wetting agents, emulsifying agents and dispersing agents. Prevention of the action of microorganisms may be ensured by the inclusion of various antibacterial and antifungal agents, for example, paraben, chlorobutanol, phenol
25 sorbic acid, and the like. It may also be desirable to include isotonic agents, such as sugars, sodium chloride, and the like into the compositions. In addition, prolonged absorption of the injectable pharmaceutical form may be brought about by the inclusion of agents which delay absorption such as aluminum monostearate and gelatin.

30 **[0119]** In some cases, in order to prolong the effect of a drug, it is desirable to slow the absorption of the drug from subcutaneous or intramuscular injection.

This may be accomplished by the use of a liquid suspension of crystalline or amorphous material having poor water solubility. The rate of absorption of the drug then depends upon its rate of dissolution which, in turn, may depend upon crystal size and crystalline form. Alternatively, delayed absorption of a parenterally-administered drug form is accomplished by dissolving or suspending the drug in an oil vehicle.

[0120] Injectable depot forms are made by forming microencapsule matrices of the subject compounds in biodegradable polymers such as polylactide-polyglycolide. Depending on the ratio of drug to polymer, and the nature of the particular polymer employed, the rate of drug release can be controlled. Examples of other biodegradable polymers include poly(orthoesters) and poly(anhydrides). Depot injectable formulations are also prepared by entrapping the drug in liposomes or microemulsions which are compatible with body tissue.

[0121] The preparations of the present invention may be given orally, parenterally, topically, or rectally. They are of course given by forms suitable for each administration route. For example, they are administered in tablets or capsule form, by injection, inhalation, eye lotion, ointment, suppository, etc. administration by injection, infusion or inhalation; topical by lotion or ointment; and rectal by suppositories. Intravenous injection administration is preferred.

[0122] The phrases "parenteral administration" and "administered parenterally" as used herein means modes of administration other than enteral and topical administration, usually by injection, and includes, without limitation, intravenous, intramuscular, intraarterial, intrathecal, intracapsular, intraorbital, intracardiac, intradermal, intraperitoneal, transtracheal, subcutaneous, subcuticular, intraarticular, subcapsular, subarachnoid, intraspinal and intrasternal injection and infusion.

[0123] The phrases "systemic administration," "administered systematically," "peripheral administration" and "administered peripherally" as used herein mean the administration of a compound, drug or other material other than directly into the central nervous system, such that it enters the patient's system and, thus, is

subject to metabolism and other like processes, for example, subcutaneous administration.

[0124] These compounds may be administered to humans and other animals for therapy by any suitable route of administration, including orally, nasally, as by,
5 for example, a spray, rectally, intravaginally, parenterally, intracisternally and topically, as by powders, ointments or drops, including buccally and sublingually.

[0125] Regardless of the route of administration selected, the compounds of the present invention, which may be used in a suitable hydrated form, and/or the pharmaceutical compositions of the present invention, are formulated into
10 pharmaceutically acceptable dosage forms by conventional methods known to those of ordinary skill in the art.

[0126] Actual dosage levels of the active ingredients in the pharmaceutical compositions of this invention may be varied so as to obtain an amount of the active ingredient which is effective to achieve the desired therapeutic response
15 for a particular patient, composition, and mode of administration, without being toxic to the patient.

[0127] The selected dosage level will depend upon a variety of factors including the activity of the particular compound of the present invention employed, or the ester, salt or amide thereof, the route of administration, the time of
20 administration, the rate of excretion of the particular compound being employed, the duration of the treatment, other drugs, compounds and/or materials used in combination with the particular compound employed, the age, sex, weight, condition, general health and prior medical history of the patient being treated, and like factors well known in the medical arts.

[0128] A physician or veterinarian having ordinary skill in the art can readily determine and prescribe the effective amount of the pharmaceutical composition required. For example, the physician or veterinarian could start doses of the compounds of the invention employed in the pharmaceutical composition at levels lower than that required in order to achieve the desired therapeutic effect
25 and gradually increase the dosage until the desired effect is achieved.
30

[0129] In general, a suitable daily dose of a compound of the invention will be that amount of the compound which is the lowest dose effective to produce a therapeutic effect. Such an effective dose will generally depend upon the factors described above. Generally, intravenous and subcutaneous doses of the compounds of this invention for a patient, when used for the indicated analgesic effects, will range from about 0.0001 to about 100 mg per kilogram of body weight per day, more preferably from about 0.01 to about 50 mg per kg per day, and still more preferably from about 0.1 to about 40 mg per kg per day. For example, between about 0.01 microgram and 20 micrograms, between about 20 micrograms and 100 micrograms and between about 10 micrograms and 200 micrograms of the compounds of the invention are administered per 20 grams of subject weight.

[0130] If desired, the effective daily dose of the active compound may be administered as two, three, four, five, six or more sub-doses administered separately at appropriate intervals throughout the day, optionally, in unit dosage forms.

[0131] Delivery of the lipoxin analogs of the present invention to the lung by way of inhalation is an important method of treating a variety of respiratory conditions noted throughout the specification, including such common local conditions as bronchial asthma and chronic obstructive pulmonary disease. The lipoxin analogs can be administered to the lung in the form of an aerosol of particles of respirable size (less than about 10 μm in diameter). The aerosol formulation can be presented as a liquid or a dry powder. In order to assure proper particle size in a liquid aerosol, as a suspension, particles can be prepared in respirable size and then incorporated into the suspension formulation containing a propellant. Alternatively, formulations can be prepared in solution form in order to avoid the concern for proper particle size in the formulation. Solution formulations should be dispensed in a manner that produces particles or droplets of respirable size.

[0132] Once prepared an aerosol formulation is filled into an aerosol canister equipped with a metered dose valve. The formulation is dispensed via an actuator adapted to direct the dose from the valve to the subject.

[0133] Formulations of the invention can be prepared by combining (i) at least one lipoxin analog in an amount sufficient to provide a plurality of therapeutically effective doses; (ii) the water addition in an amount effective to stabilize each of the formulations; (iii) the propellant in an amount sufficient to propel a plurality of doses from an aerosol canister; and (iv) any further optional components e.g. ethanol as a cosolvent; and dispersing the components. The components can be dispersed using a conventional mixer or homogenizer, by shaking, or by ultrasonic energy. Bulk formulation can be transferred to smaller individual aerosol vials by using valve to valve transfer methods, pressure filling or by using conventional cold-fill methods. It is not required that a stabilizer used in a suspension aerosol formulation be soluble in the propellant. Those that are not sufficiently soluble can be coated onto the drug particles in an appropriate amount and the coated particles can then be incorporated in a formulation as described above.

[0134] Aerosol canisters equipped with conventional valves, preferably metered dose valves, can be used to deliver the formulations of the invention. Conventional neoprene and buna valve rubbers used in metered dose valves for delivering conventional CFC formulations can be used with formulations containing HFC-134a or HFC-227. Other suitable materials include nitrile rubber such as DB-218 (American Gasket and Rubber, Schiller Park, Ill.) or an EPDM rubber such as VistalonTM. (Exxon), RoyaleneTM (UniRoyal), bunaEP (Bayer). Also suitable are diaphragms fashioned by extrusion, injection molding or compression molding from a thermoplastic elastomeric material such as FLEXOMERTM GERS 1085 NT polyolefin (Union Carbide).

[0135] Formulations of the invention can be contained in conventional aerosol canisters, coated or uncoated, anodized or unanodized, e.g., those of aluminum, glass, stainless steel, polyethylene terephthalate.

[0136] The formulation(s) of the invention can be delivered to the respiratory tract and/or lung by oral inhalation in order to effect bronchodilation or in order to treat a condition susceptible of treatment by inhalation, e.g., asthma, chronic obstructive pulmonary disease, etc. as described throughout the specification.

[0137] The formulations of the invention can also be delivered by nasal inhalation as known in the art in order to treat or prevent the respiratory conditions mentioned throughout the specification.

5 [0138] While it is possible for a compound of the present invention to be administered alone, it is preferable to administer the compound as a pharmaceutical composition.

[0139] The invention features an article of manufacture that contains packaging material and a lipoxin formulation contained within the packaging material. This formulation contains an at least one lipoxin analog and the packaging material
10 contains a label or package insert indicating that the formulation can be administered to the subject to treat one or more respiratory conditions, in an amount, at a frequency, and for a duration effective to treat or prevent such respiratory condition(s). Such conditions are mentioned throughout the specification.

15 [0140] More specifically, the invention features an article of manufacture that contains packaging material and a lipoxin formulation contained within the packaging material. This formulation contains at least one lipoxin analog. The packaging material contains a label or package insert indicating that the formulation can be administered to the subject to asthma in an amount, at a
20 frequency, and for a duration effective treat or prevent asthma symptoms.

Methods

[0141] *Sensitization and challenge protocols.* Five to seven week old male BALB/c (Charles River Laboratories, Wilmington, Massachusetts) or female hCD11b-hALX FvB transgenic mice were housed in isolation cages under viral
25 antibody-free conditions. After Harvard Medical Area IRB approval (Protocol #02570), mice were sensitized with intraperitoneal injections of ovalbumin (OVA) (Grade III; Sigma Chemical Co., St. Louis, Missouri) (10 µg) plus 1 mg aluminum hydroxide (ALUM) (J.T. Baker Chemical Co.; Phillipsburg, New Jersey) as adjuvant in 0.2 ml PBS on days 0 and 7. On days 14, 15, 16 and 17,

the mice received 10 µg/mouse of an analog (ref. 16) of LXA₄ (15-epi,16-para-fluoro-phenoxy-LXA₄-methyl ester, LXa) or PBS with 1.6 mM CaCl₂ and 1.6 mM MgCl₂ (0.1 ml) by intravenous injection at least 1 h prior to an aerosol challenge containing either PBS or 6% OVA for 25 min/day. The analog was
5 designed to resist rapid enzymatic inactivation and was based on the structure of the aspirin-triggered 15-epi-LXA₄ (ref. 16), which carries its carbon 15 position alcohol in the *R* configuration or epimeric (*R*) to native LXA₄. On day 18, 24 h after the last aerosol challenge, airway responsiveness to intravenous methacholine (33 – 1000 µg/kg) was measured, bilateral bronchoalveolar lavage
10 (BAL) (2 aliquots of 1 ml PBS plus 0.6 mM EDTA) was performed and tissues (whole blood, lungs, mediastinal lymph nodes and spleen) were harvested for histological analysis. Lung resistance (R_L) was measured using a sealed constant mass plethysmograph. The effective dose of methacholine required to increase R_L to 200% of control values was defined as the ED₂₀₀ and used as an index of
15 airway responsiveness (as in refs. 14, 15). For mice studied using a whole body plethysmograph to assess airway responsiveness (Buxco), each mouse was placed into a chamber and box pressure/time waveform was analyzed to yield the indicator of airflow obstruction, enhanced pause (Penh). PBS or methacholine (100 mg/ml) was given by aerosol through an inlet of the chamber for 4.5 min.
20 Readings were initiated at 3 min and continued for 12 min. Peak and one min average Penh values were determined. Serum total IgE levels were determined by ELISA (Crystal Chem, Chicago, Illinois). *In situ* hybridization of murine ALX was performed using an anti-sense oligonucleotide probe (534 bp in length) corresponding to nucleic acids +581 - +1115 (GenBank Accession No.
25 NM008042) with the assistance of the Dana-Farber/ Harvard Cancer Center Pathology Core Facility (under the direction of Dr. Massimo Loda).

[0142] *Allergen-initiated respiratory inflammation.* Measurement of inflammatory mediators was determined in cell-free BAL fluid (2000 g, 10 min) by sensitive and specific ELISA's, in tandem, for interleukin-5 (IL-5), IL-13,
30 eotaxin, tumor necrosis factor (TNFα) (R&D Systems, Minneapolis, Minnesota), LTB₄, cysteinyl LT, PGE₂ (Cayman Chemical Co., Ann Arbor, Michigan), and

*Trade-mark

LXA₄ (Neogen, Lexington, Kentucky). Cells were resuspended in HBSS, enumerated by hemocytometer, and concentrated onto microscope slides by cytocentrifuge (STATspin) (265 g). Cells were stained with a Wright-Giemsa stain (Sigma Chemical Co.) to determine leukocyte differentials (after counting ≥
5 200 cells).

[0143] *Eosinophil tissue infiltration.* Five minutes after topical delivery of LXA (10 µg in 10 µl acetone) or vehicle control, PGE₂ (1 µg, Cayman Chemical Co.) plus LTB₄ (1 µg, Cayman Chemical Co.) were topically applied employing 10 µl acetone that rapidly evaporates to the inside of left and right ears of male mice
10 (hCD11b-hALX FvB transgenic, hCD11b-hBLT FvB transgenic ³⁶ or age and gender-matched non-transgenic littermates; 4-5 months old, ~30g body weight). After 13-16 h, 6 mm diameter skin punch biopsies (Acu-Punch, Fisher Scientific, Pittsburgh, Pennsylvania) were obtained. Samples were sliced finely with scalpels, homogenized in 400 µl of potassium phosphate buffer (pH 6.0)
15 containing 0.5% hexadecyltrimethylammonium bromide, followed by three cycles of sonication and freeze-thaw. The particulate matter was removed by centrifugation (16,000g for 20 min) and resulting supernatants were assayed for eosinophil peroxidase (EPO) activity as in ref. 37. Briefly, 3-(4-hydroxyphenyl) propionic acid (HPA, Aldrich, Milwaukee, Wisconsin) (100 µM) was exposed
20 (10 min, 37°C) to partially purified EPO (~75-150 µg extracted ear skin protein) in the presence of NaBr (100 µM). EPO activity was monitored by mono-bromination of HPA (Br-HPA). To extract Br-HPA, tissue samples were spiked with an internal standard (L-tyrosine(ring)-d₄, Cambridge Isotopes, Andover, Massachusetts), loaded onto C18 SepPak cartridges (Waters, Milford,
25 Massachusetts) for extraction in 0.1% TFA, eluted with MeOH:dH₂O (1:1, v/v) + 0.1% TFA, brought to dryness *in vacuo* (rotoevaporation) and derivatized (overnight) with BSTFA (Pierce, Rockford, Illinois). Trimethyl silylated HPA, Br-HPA and d₄-tyrosine were detected by GC-MS (GC model # 6890, MS model # 5973, Hewlett Packard, San Fernando, California). HPA-TMS had a retention
30 time of 7.01 min with diagnostic molecular ion (M⁺=310) and mass

fragmentation, including m/z 295 [M^+ -CH₃], m/z 192 [M^+ -COOTMS], and a base peak of m/z 179 [M^+ -CH₂COOTMS]. Br-HPA-TMS had a retention time of 9.30 min with diagnostic molecular ion (M^+ =390) and mass fragmentation, including m/z 375 [M^+ -CH₃], m/z 272 [M^+ -COOTMS], and a base peak of m/z 259 [M^+ -CH₂COOTMS]. In addition, the mass spectrum of Br-HPA demonstrated the isotopic pattern of a monobrominated species. EPO activity was quantitated by percent conversion of HPA to Br-HPA, taking into account the recovery of the internal standard (>80%) and normalized for the samples' protein content (determined by BioRad protein reagent) (BioRad, Hercules, California).

10 [0144] TABLE I

Eicosanoid	Levels (pg/ml)	BAL Samples (n)
PGE ₂	1117.7 $\pm$ 103.8	5
CysLT	139.0 $\pm$ 27.3	9
LTB ₄	6.4 $\pm$ 2.3	9
LXA ₄	15.0 $\pm$ 3.3	5

*Mice were sensitized to OVA and challenged daily with an aerosol of 6% OVA for 4 consecutive days (see *Methods*). BAL was performed 24 h after the final OVA aerosol. After centrifugation, eicosanoid levels were determined using ELISA's with the cell-free BAL supernatants. Values are the mean (pg/ml) of duplicate determinations.

[0145] References

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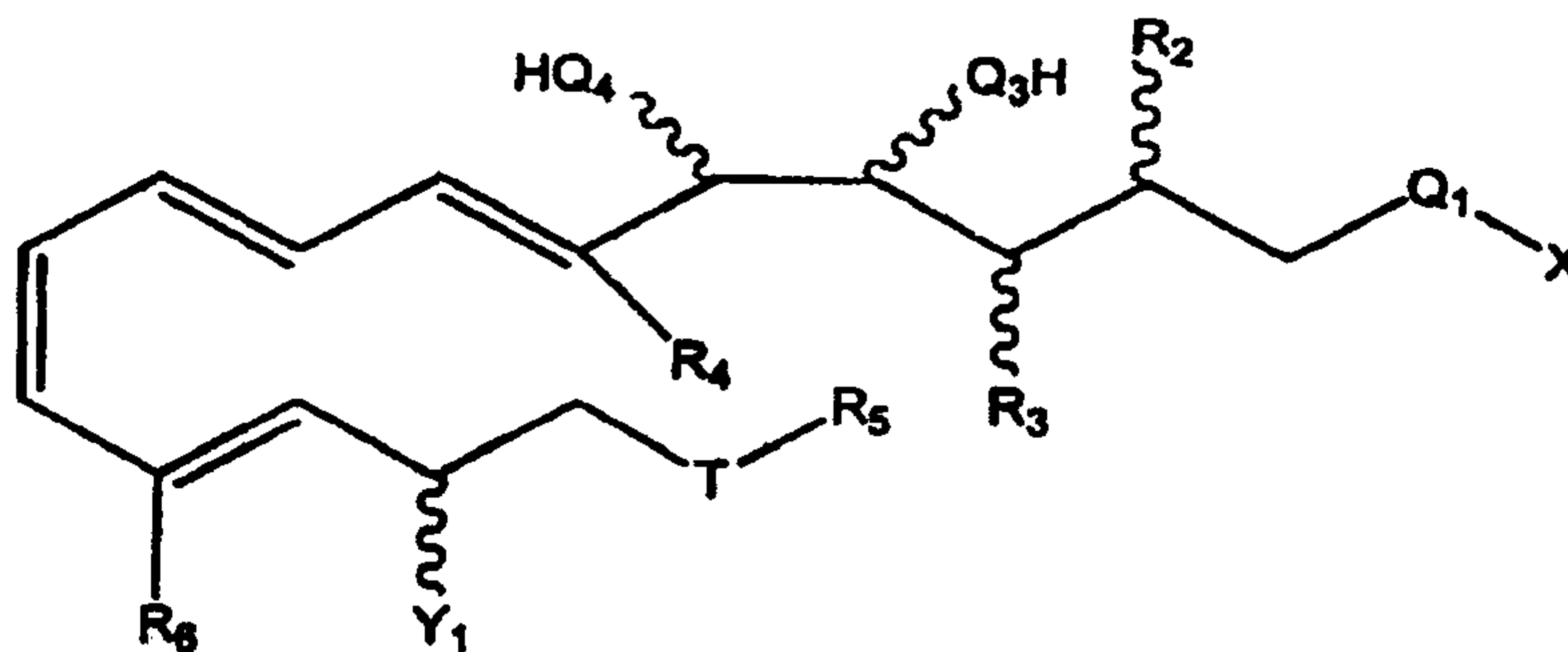
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[0146] One having ordinary skill in the art will appreciate further features and
10 advantages of the invention based on the above-described embodiments. Accordingly, the invention is not to be limited by what has been particularly shown and described, except as indicated by the appended claims.

WE CLAIM:

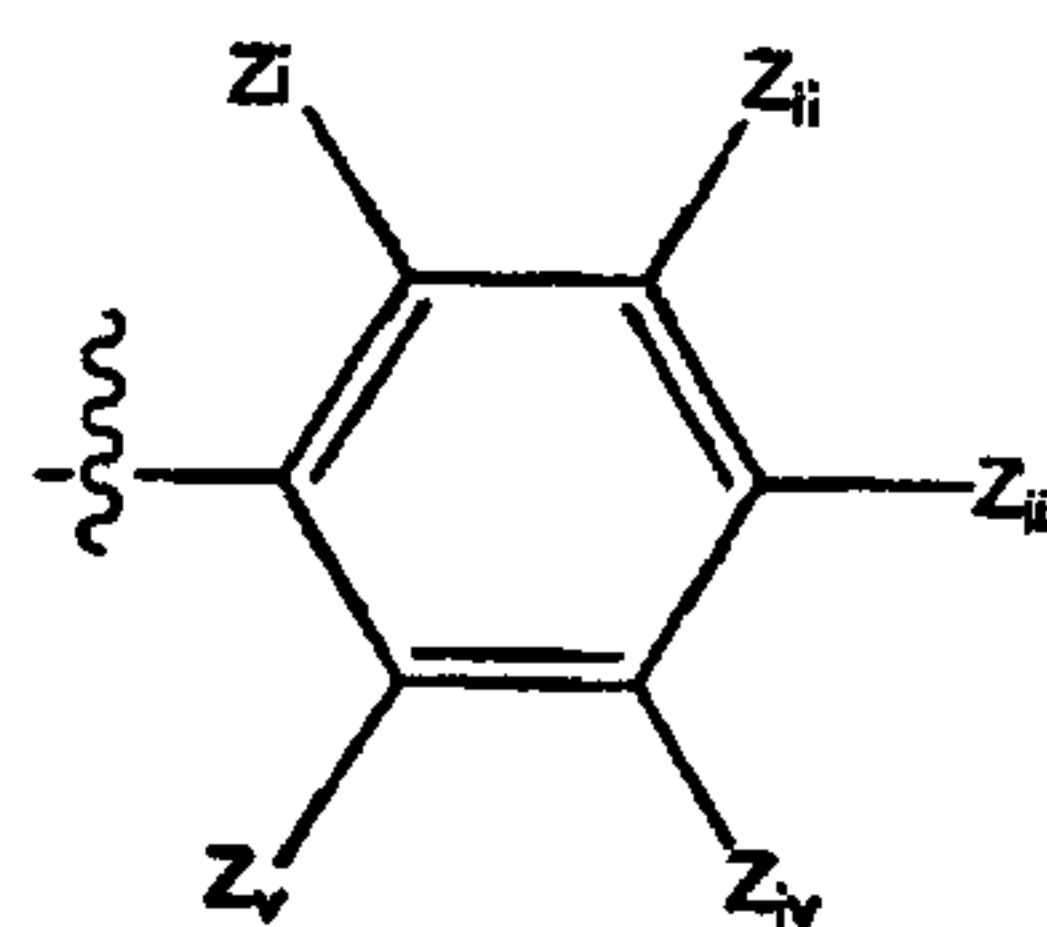
1. Use of a lipoxin analog for treating or preventing disorders of the respiratory tract or lung selected from the group consisting of chronic bronchitis, bronchiectasis, eosinophilic lung diseases, allergic bronchopulmonary aspergillosis, bronchiolitis, bronchiolitis obliterans, bronchiolitis obliterans with organizing pneumonia, eosinophilic granuloma, Wegener's granulomatosis, sarcoidosis, hypersensitivity pneumonitis, lung injury associated with parasite infection and chronic obstructive pulmonary disease (COPD), said lipoxin analog having the formula



wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl;

(vii) a detectable label molecule; or

(viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) , provided when Q_1 is CN , then X is absent;

wherein Q_3 and Q_4 are each independently O , S or NH ;

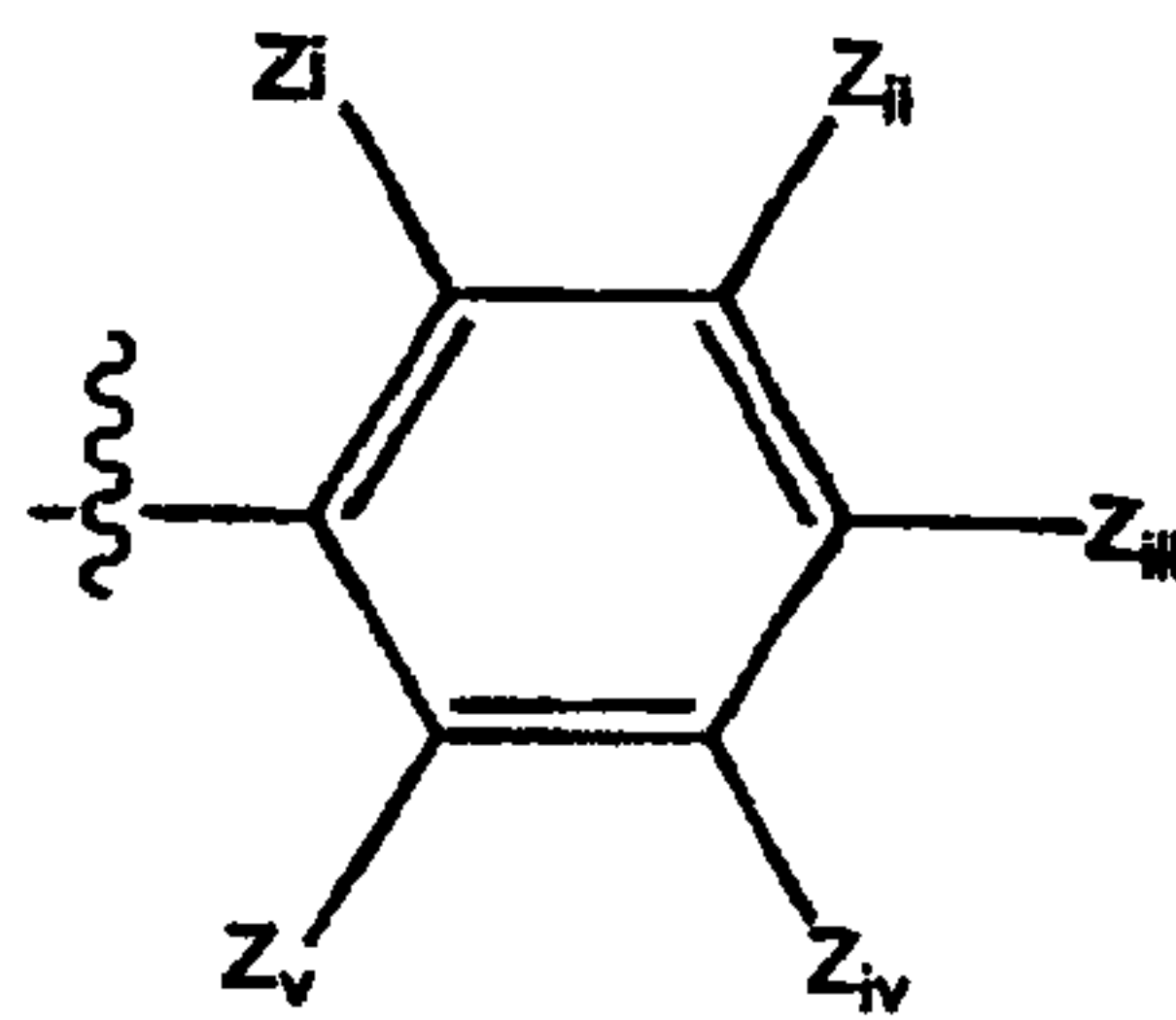
wherein one of R_2 and R_3 is a hydrogen atom and the other is

- (a) H ;
- (b) an alkyl of 1 to 8 carbon atoms, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms;
- (d) an alkenyl of 2 to 8 carbon atoms, which may be straight chain or branched; or
- (e) $R_a Q_2 R_b$ wherein Q_2 is $-\text{O}-$ or $-\text{S}-$; wherein R_a is alkylene of 0 to 6 carbon atoms, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

- (a) H ;
- (b) an alkyl of 1 to 6 carbon atoms, which may be a straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

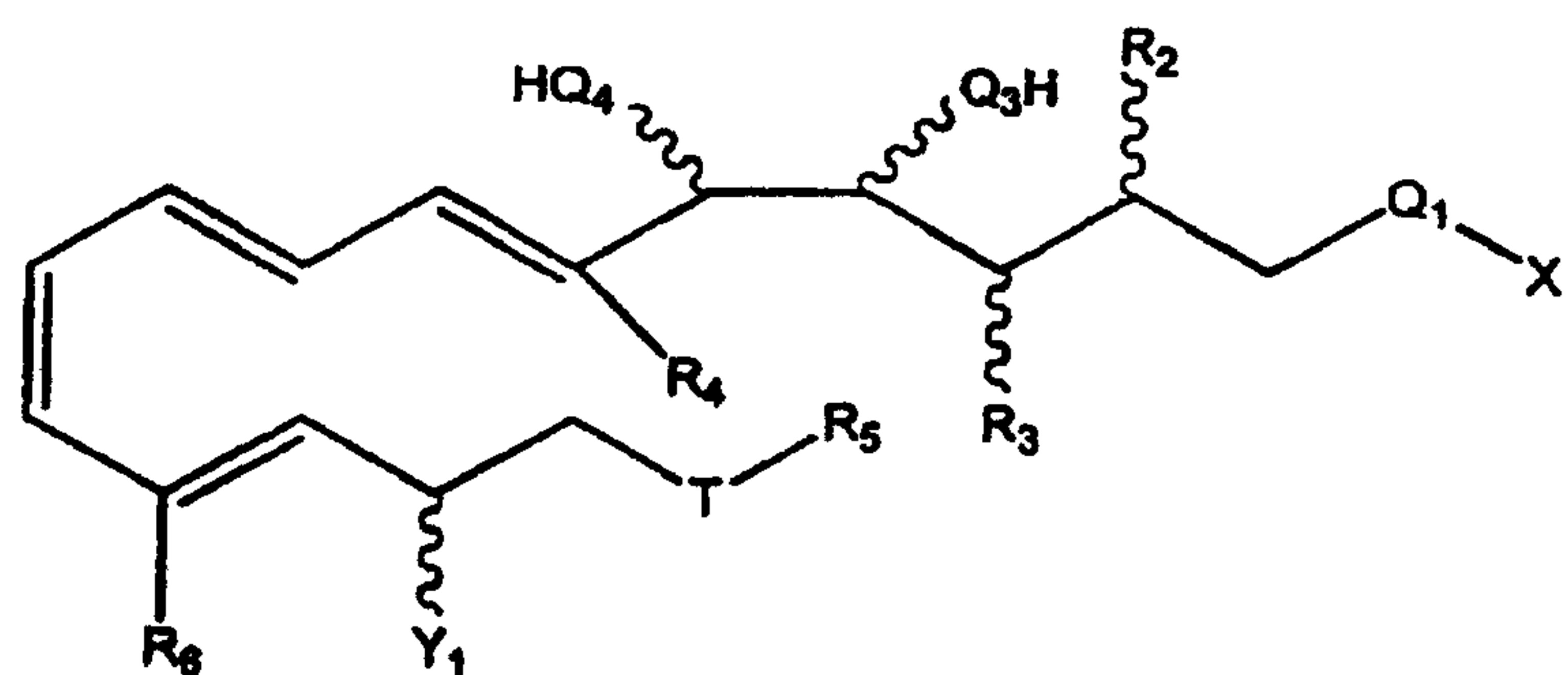
wherein Y_1 is $-\text{OH}$, methyl, $-\text{SH}$, an alkyl of 2 to 4 carbon atoms, straight chain or branched, an alkoxy of 1 to 4 carbon atoms, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano, nitro or a halogen;

wherein R_6 is

- (a) H ;
- (b) an alkyl from 1 to 4 carbon atoms, straight chain or branched;

wherein T is O or S , and pharmaceutically acceptable salts thereof; with the proviso that the lipoxin analog is not 16-phenoxy lipoxin A_4 .

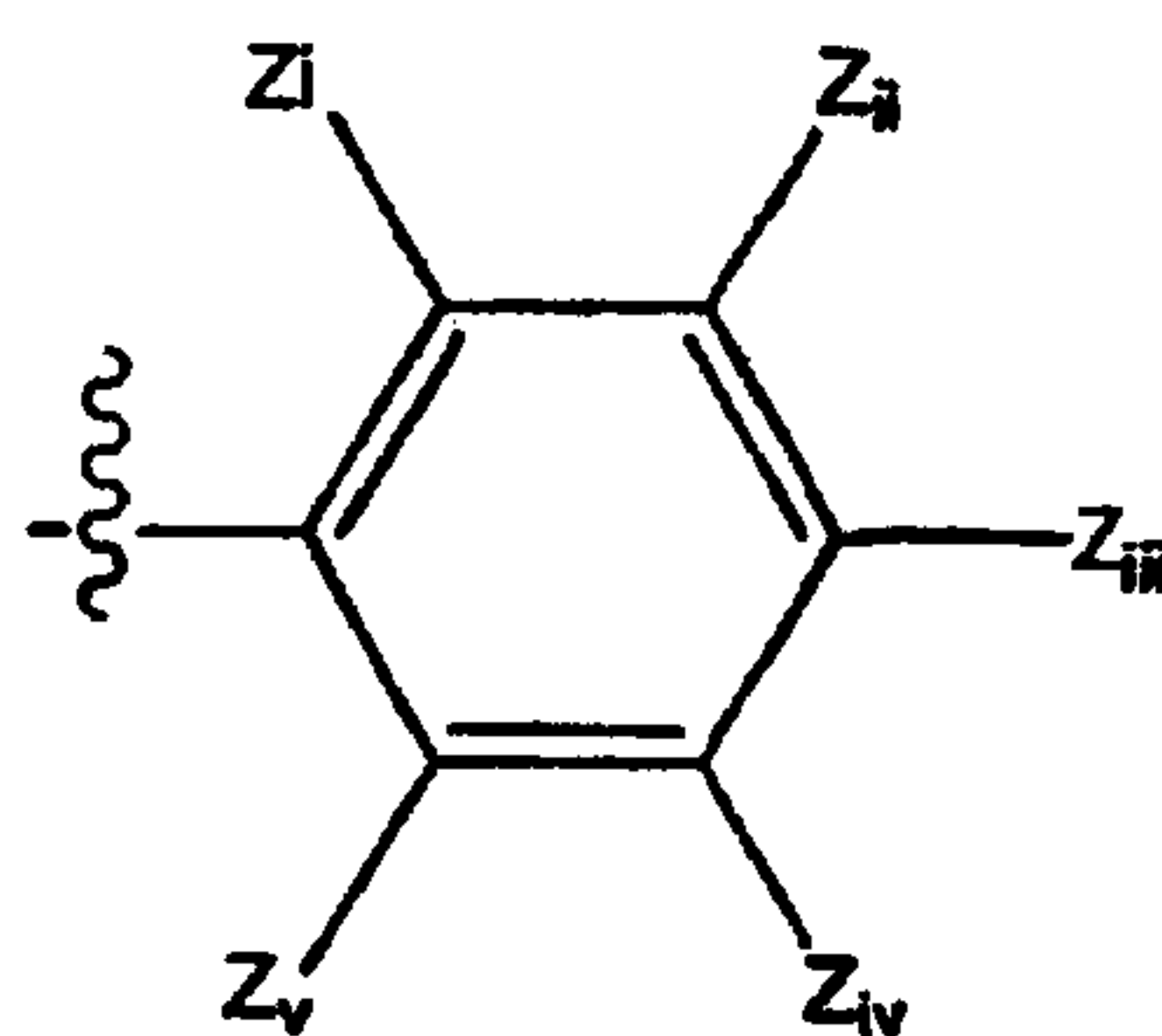
2. A pharmaceutical composition for use in treating or preventing disorders of the respiratory tract or lung in a subject selected from the group consisting of chronic bronchitis, bronchiectasis, eosinophilic lung diseases, allergic bronchopulmonary aspergillosis, bronchiolitis, bronchiolitis obliterans, bronchiolitis obliterans with organizing pneumonia, eosinophilic granuloma, Wegener's granulomatosis, sarcoidosis, hypersensitivity pneumonitis, lung injury associated with parasite infection and chronic obstructive pulmonary disease (COPD), comprising a therapeutically effective amount of at least one lipoxin analog having the formula



wherein X is R_1 , OR_1 , or SR_1 ;

wherein R_1 is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, which may be straight chain or branched;
- (iii) a cycloalkyl of 3 to 10 carbon atoms;
- (iv) an aralkyl of 7 to 12 carbon atoms;
- (v) phenyl;
- (vi) substituted phenyl



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-NO_2$, $-CN$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl;

- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms;

wherein Q_1 is $(C=O)$, SO_2 or (CN) , provided when Q_1 is CN , then X is absent;

wherein Q_3 and Q_4 are each independently O, S or NH ;

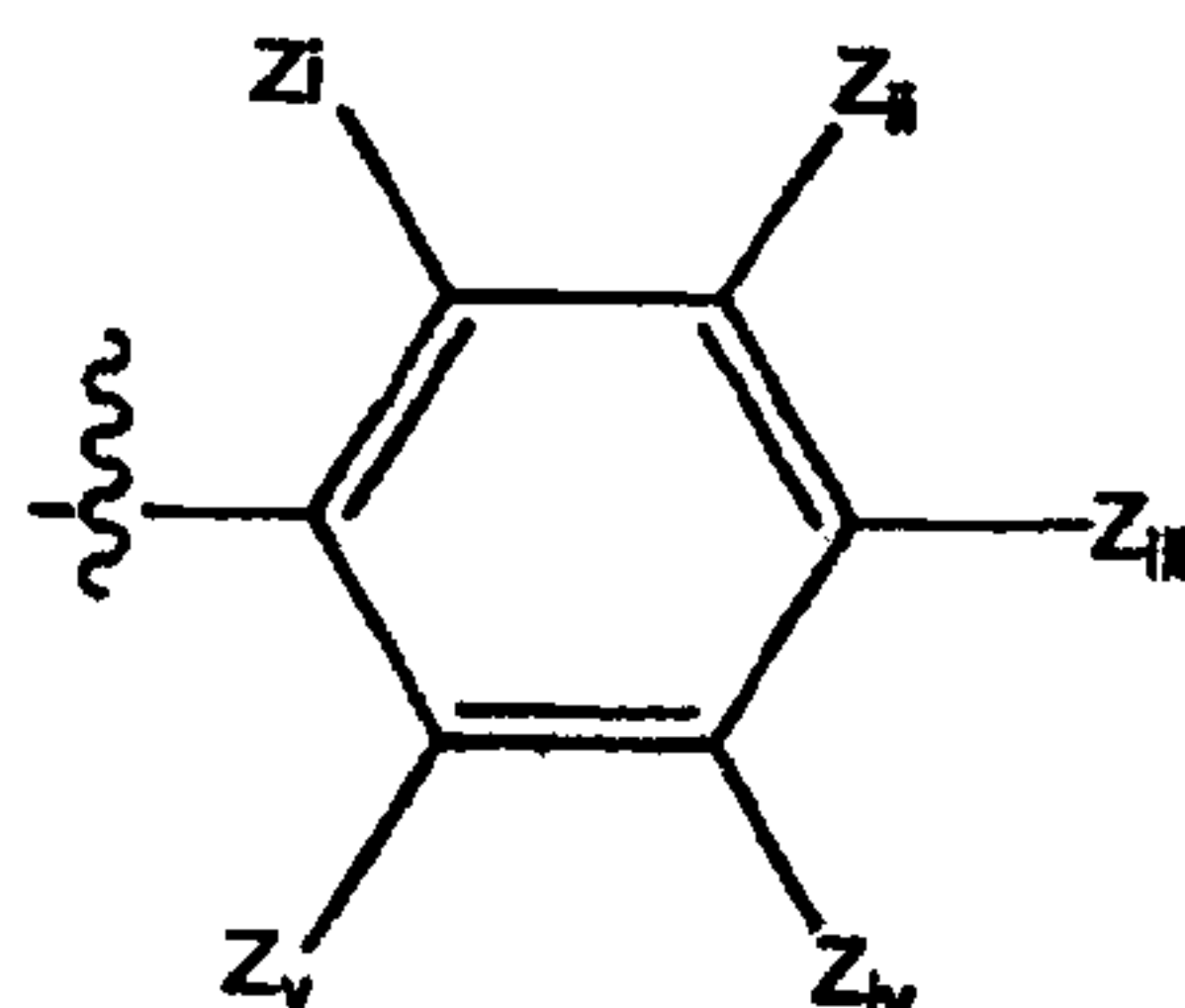
wherein one of R_2 and R_3 is a hydrogen atom and the other is

- (a) H;
- (b) an alkyl of 1 to 8 carbon atoms, which may be a straight chain or branched;
- (c) a cycloalkyl of 3 to 6 carbon atoms;
- (d) an alkenyl of 2 to 8 carbon atoms, which may be straight chain or branched; or
- (e) $R_aQ_2R_b$ wherein Q_2 is -O- or -S-; wherein R_a is alkylene of 0 to 6 carbon atoms, which may be straight chain or branched and wherein R_b is alkyl of 0 to 8 carbon atoms, which may be straight chain or branched, provided when R_b is 0, then R_b is a hydrogen atom;

wherein R_4 is

- (a) H;
- (b) an alkyl of 1 to 6 carbon atoms, which may be a straight chain or branched;

wherein R_5 is



wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} , and Z_v are each independently selected from $-NO_2$, $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is an alkyl group of 1 to 8 carbon atoms, which may be a straight chain or branched, and hydroxyl or a substituted or unsubstituted, branched or unbranched alkyl group;

wherein Y_1 is $-OH$, methyl, $-SH$, an alkyl of 2 to 4 carbon atoms, straight chain or branched, an alkoxy of 1 to 4 carbon atoms, or CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3 and Z is cyano, nitro or a halogen;

wherein R_6 is

- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, straight chain or branched;

wherein T is O or S, and pharmaceutically acceptable salts thereof with the proviso that the lipoxin analog is not 16-phenoxy lipoxin A₄; and a pharmaceutically acceptable carrier.

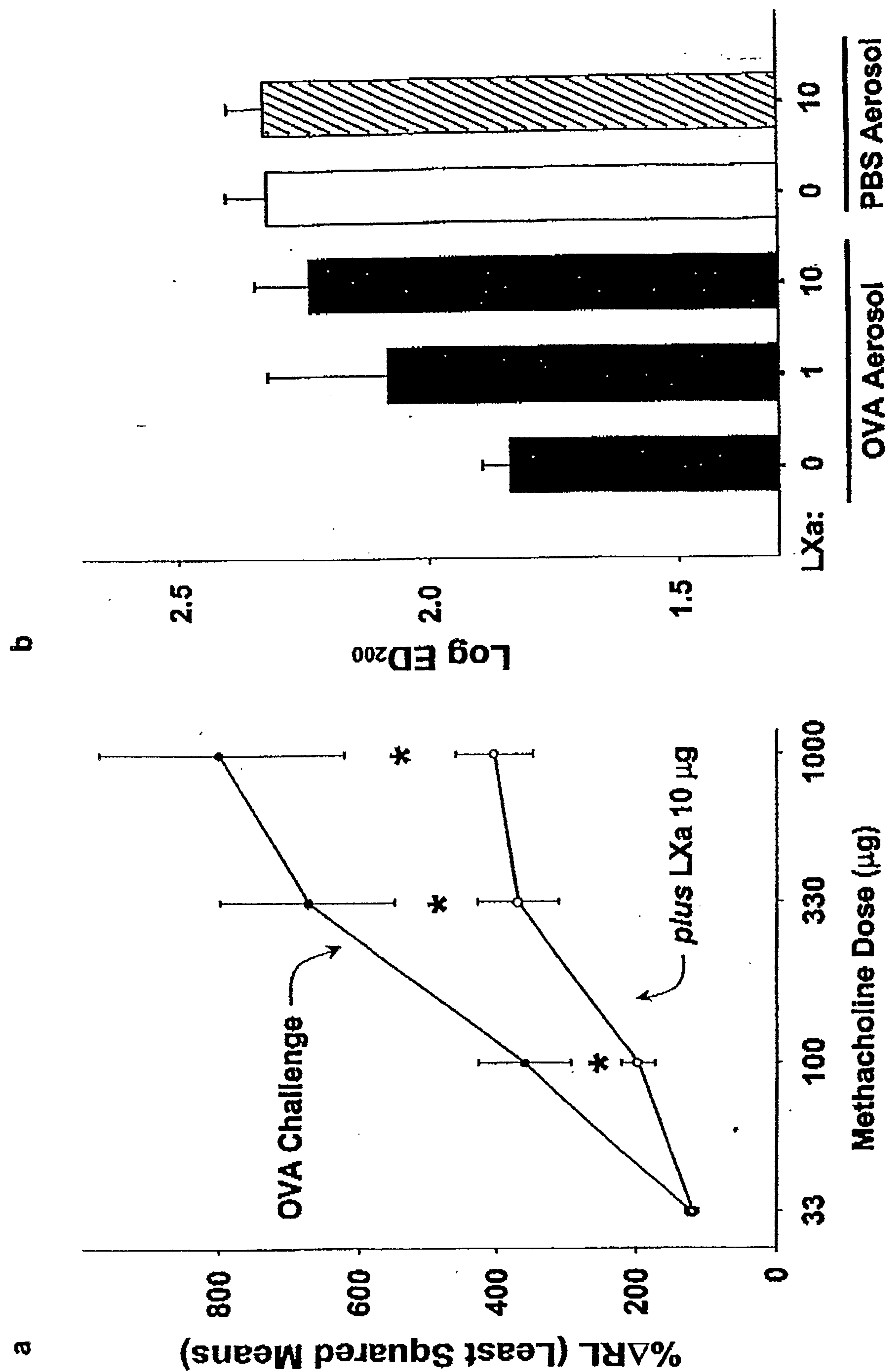
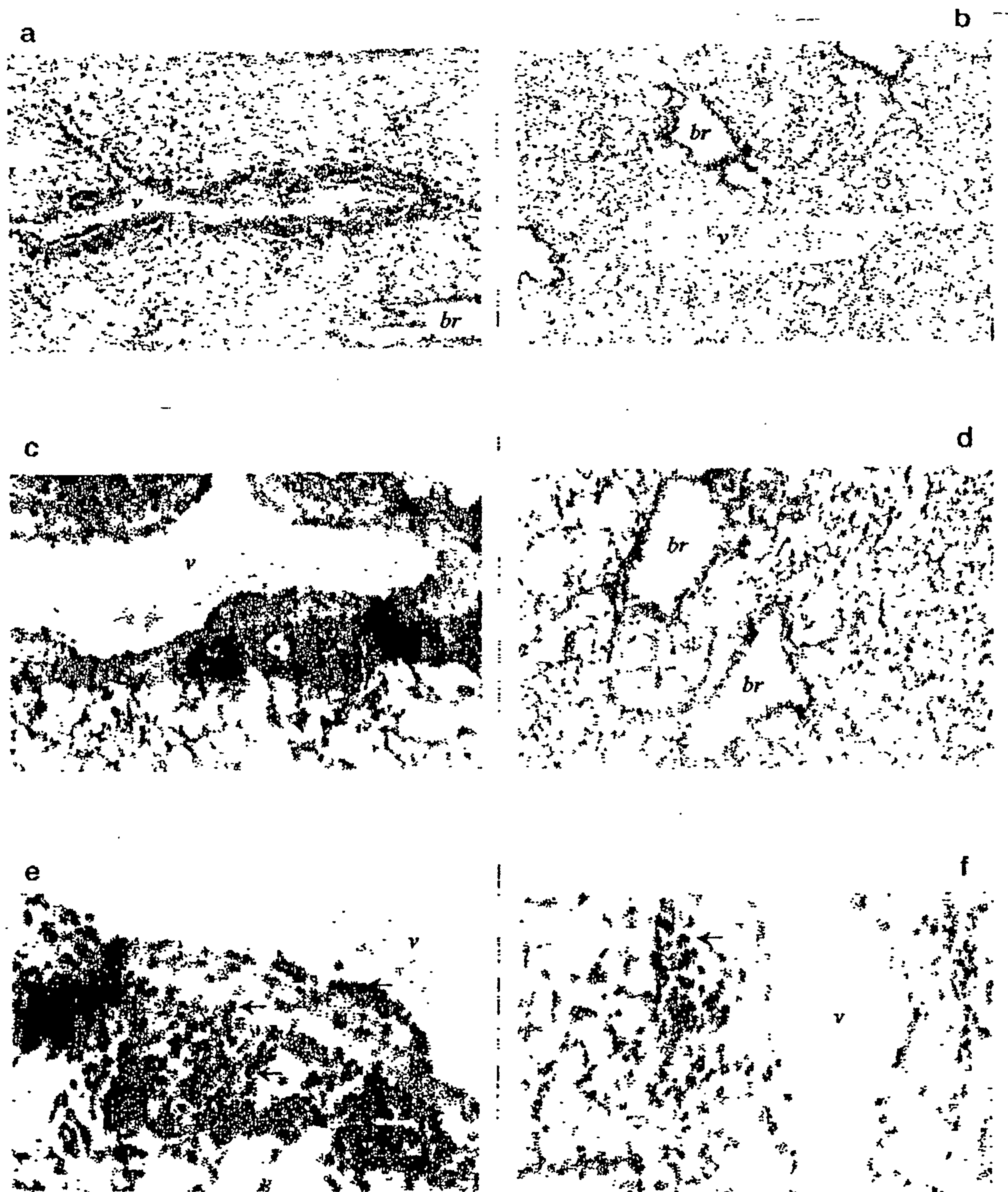


Fig. 1

Fig. 2

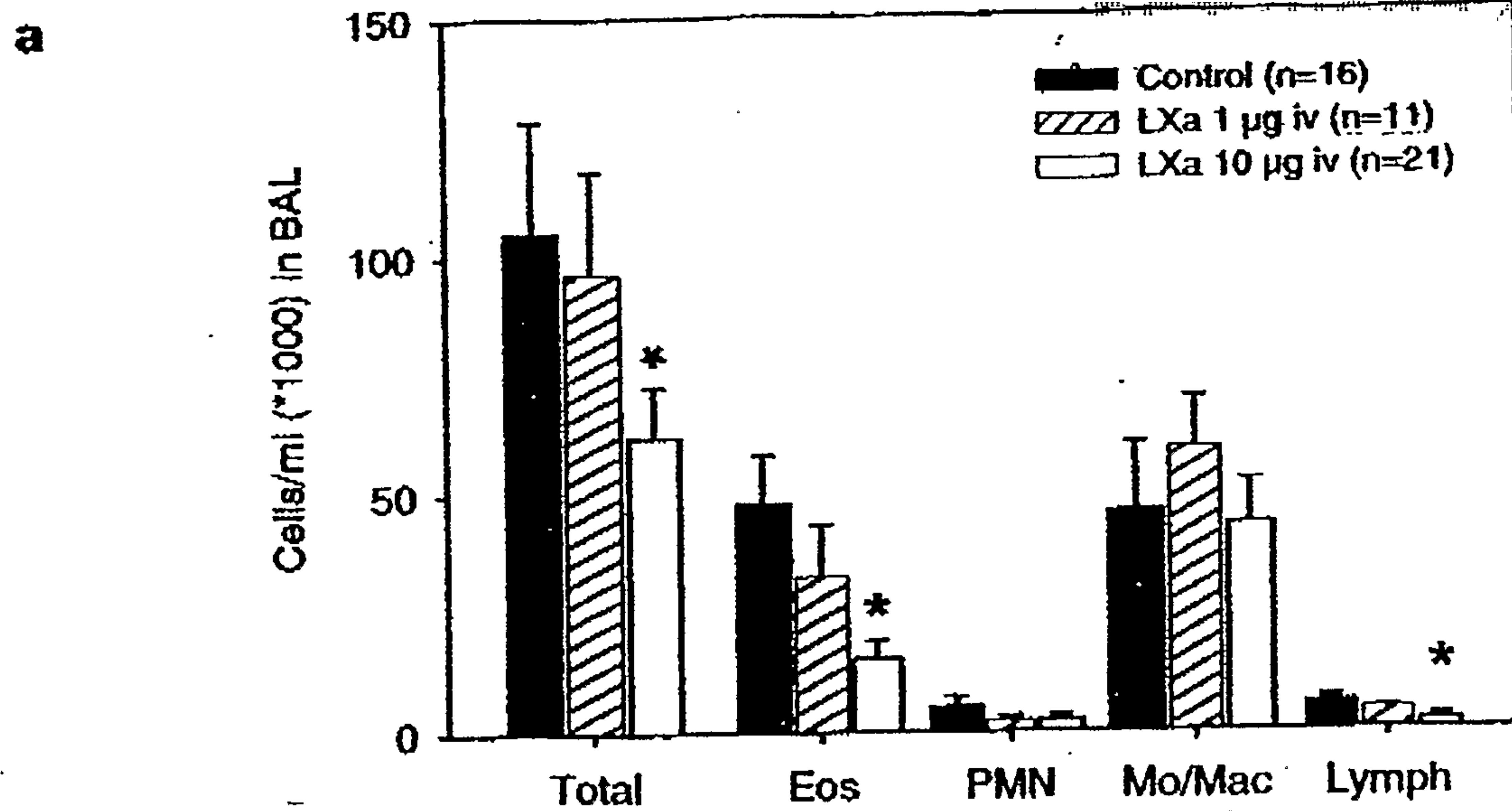
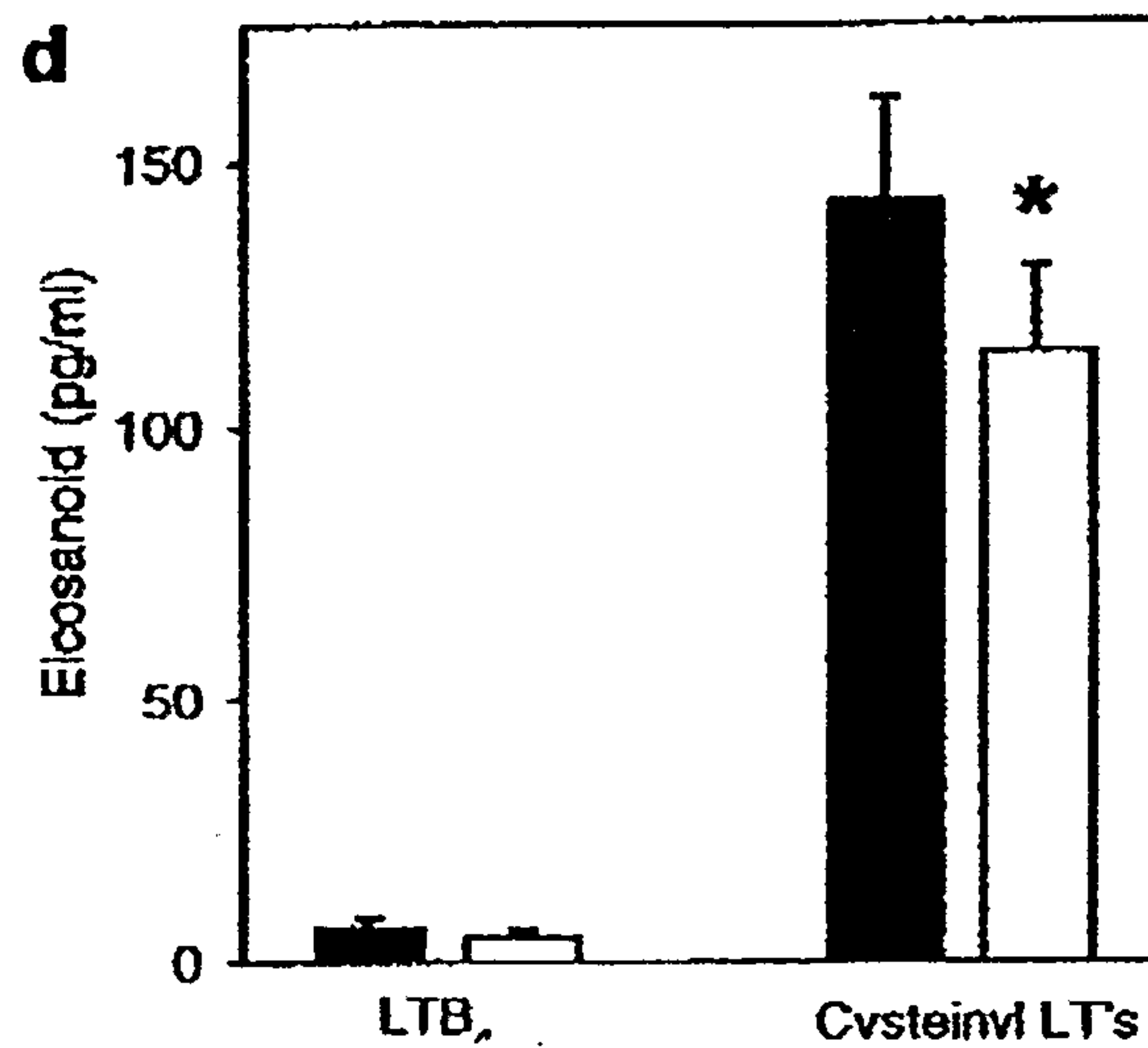
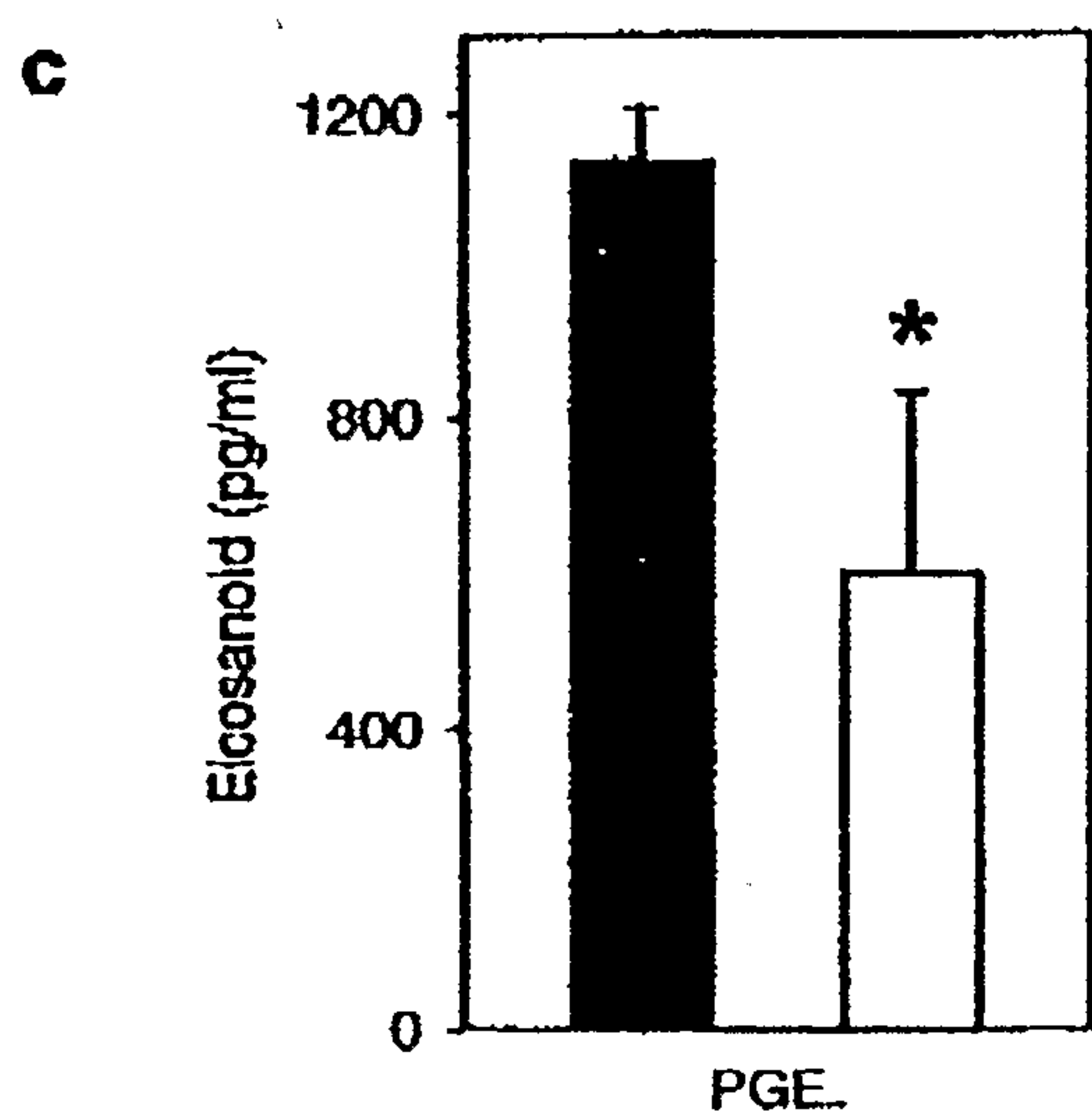
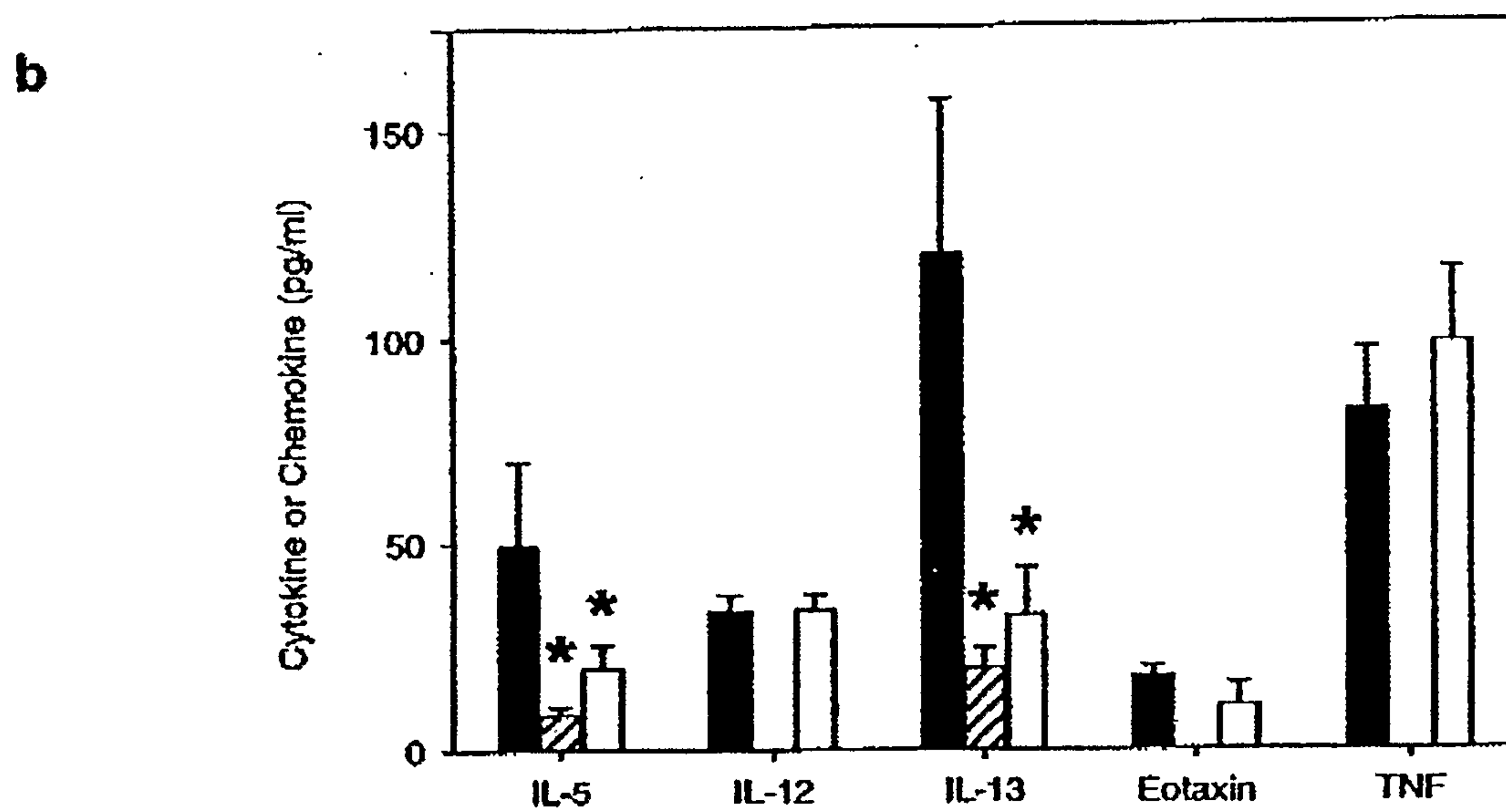
**Fig. 3**

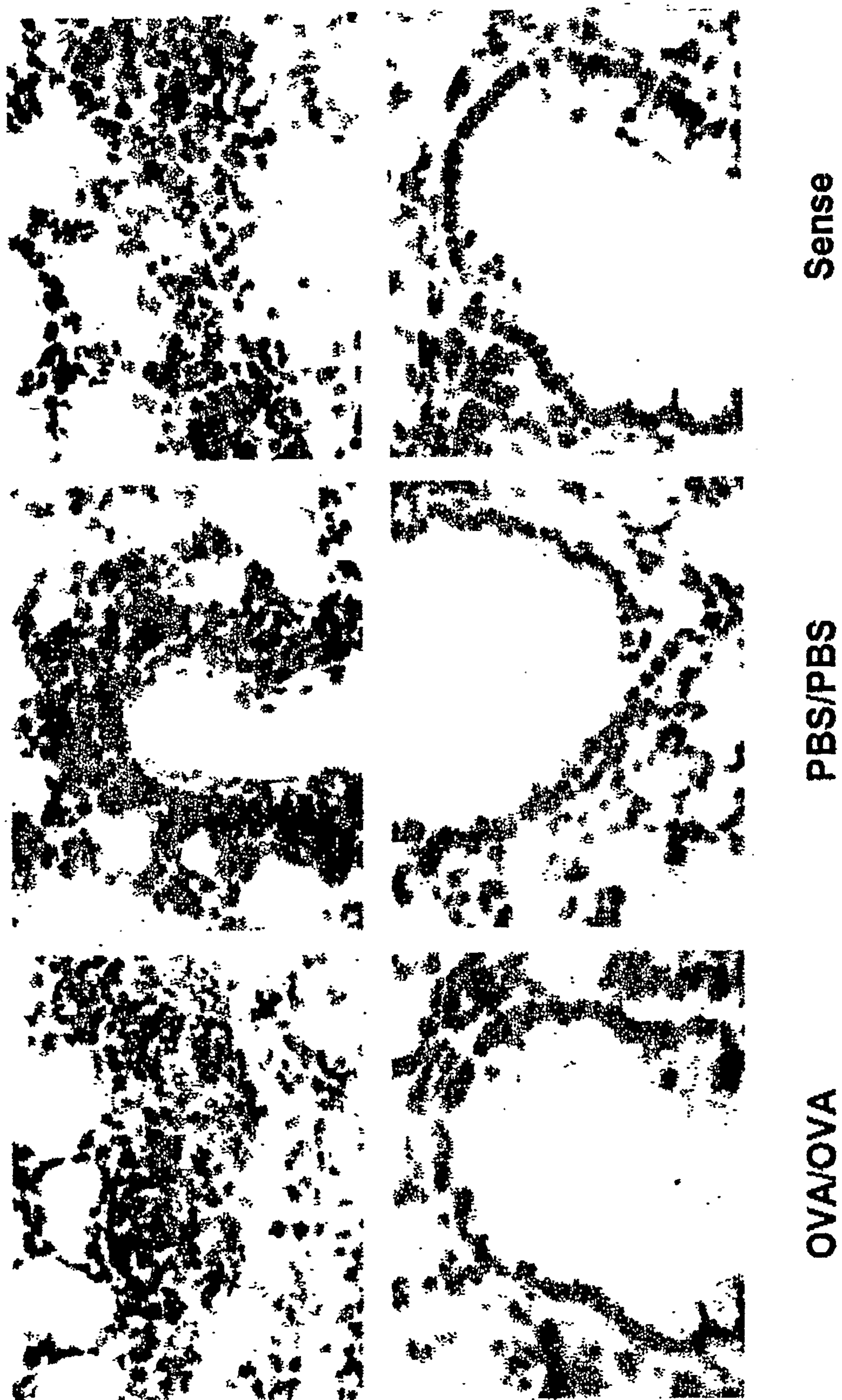
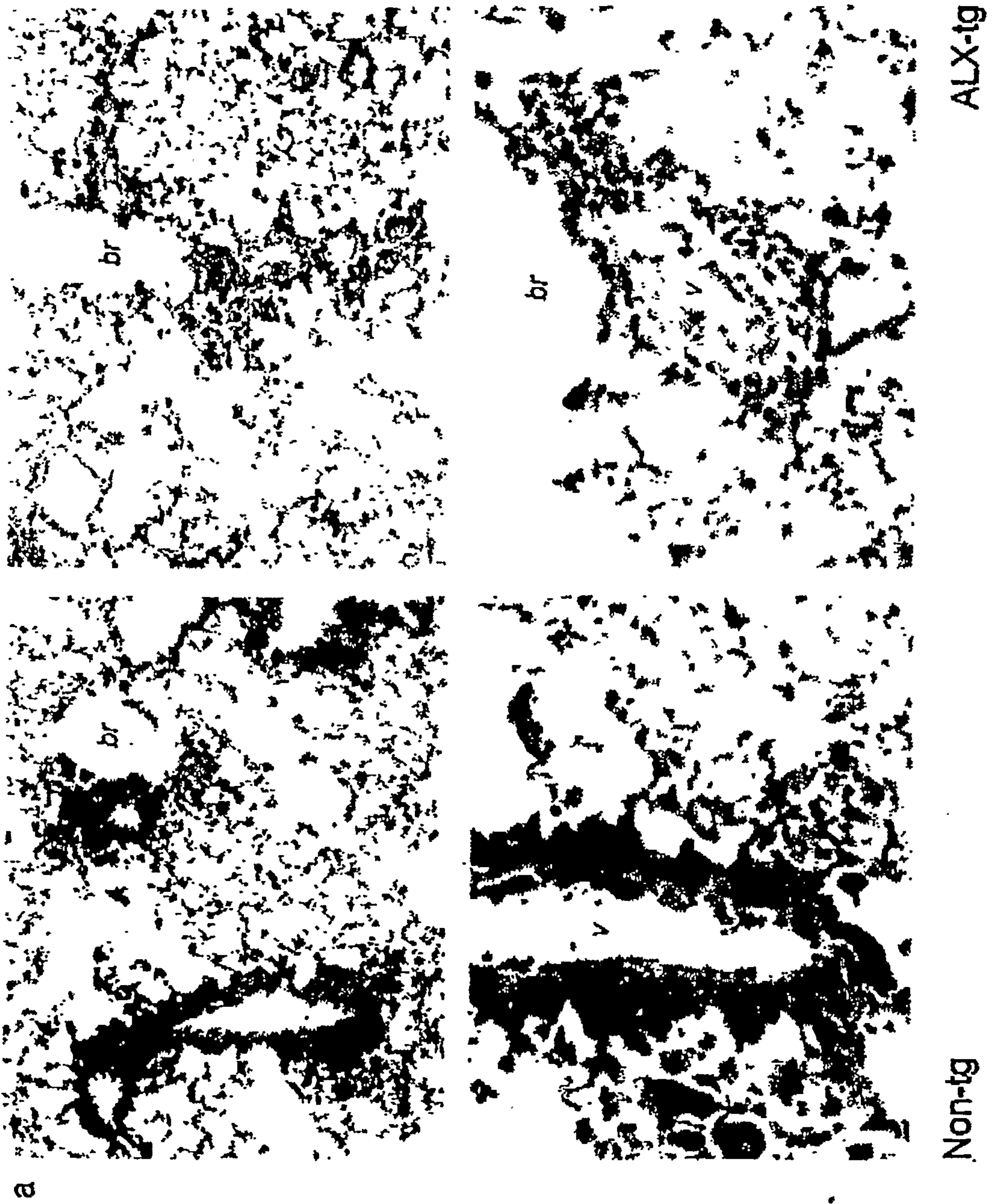
Fig. 4

Fig. 5a

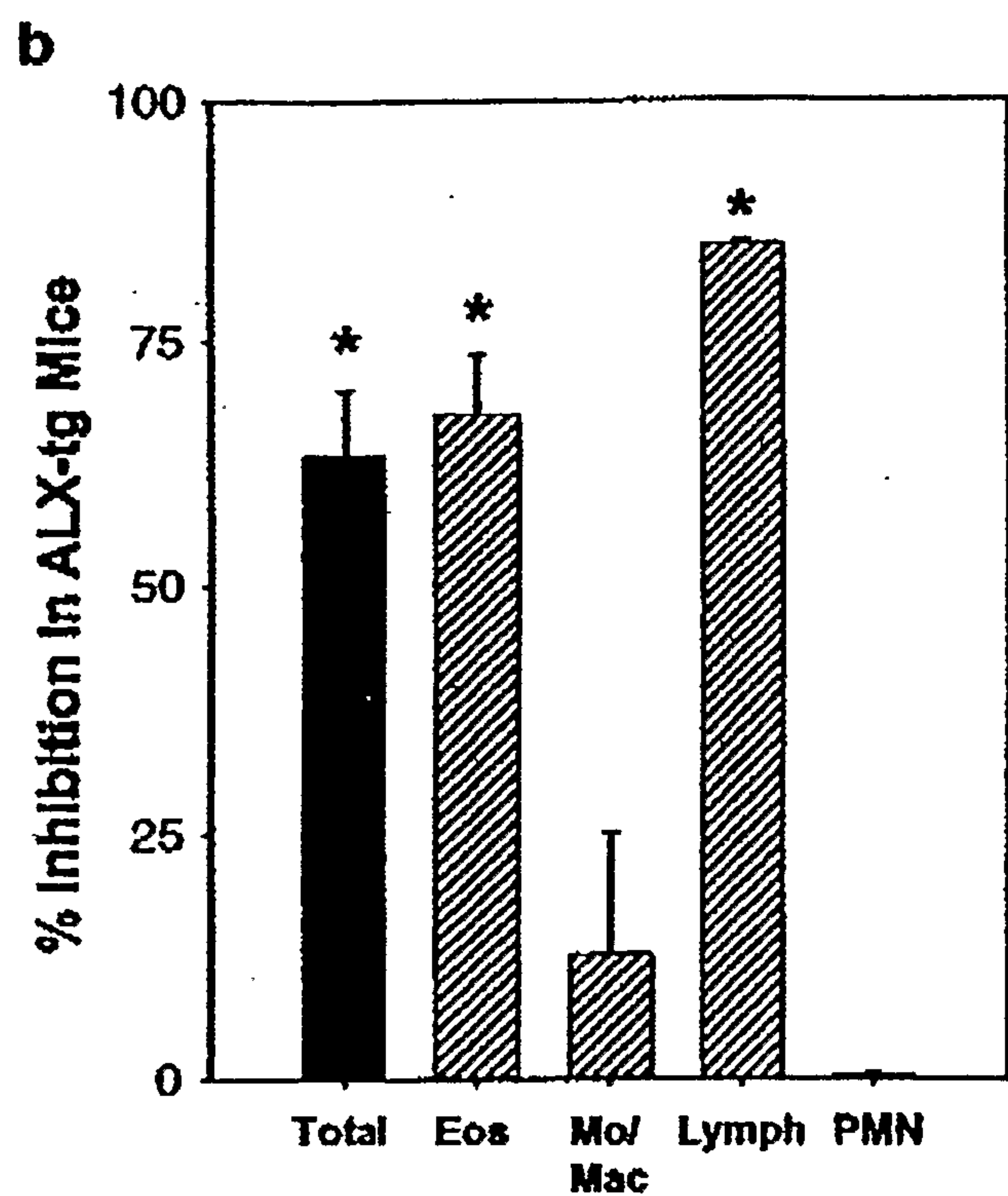
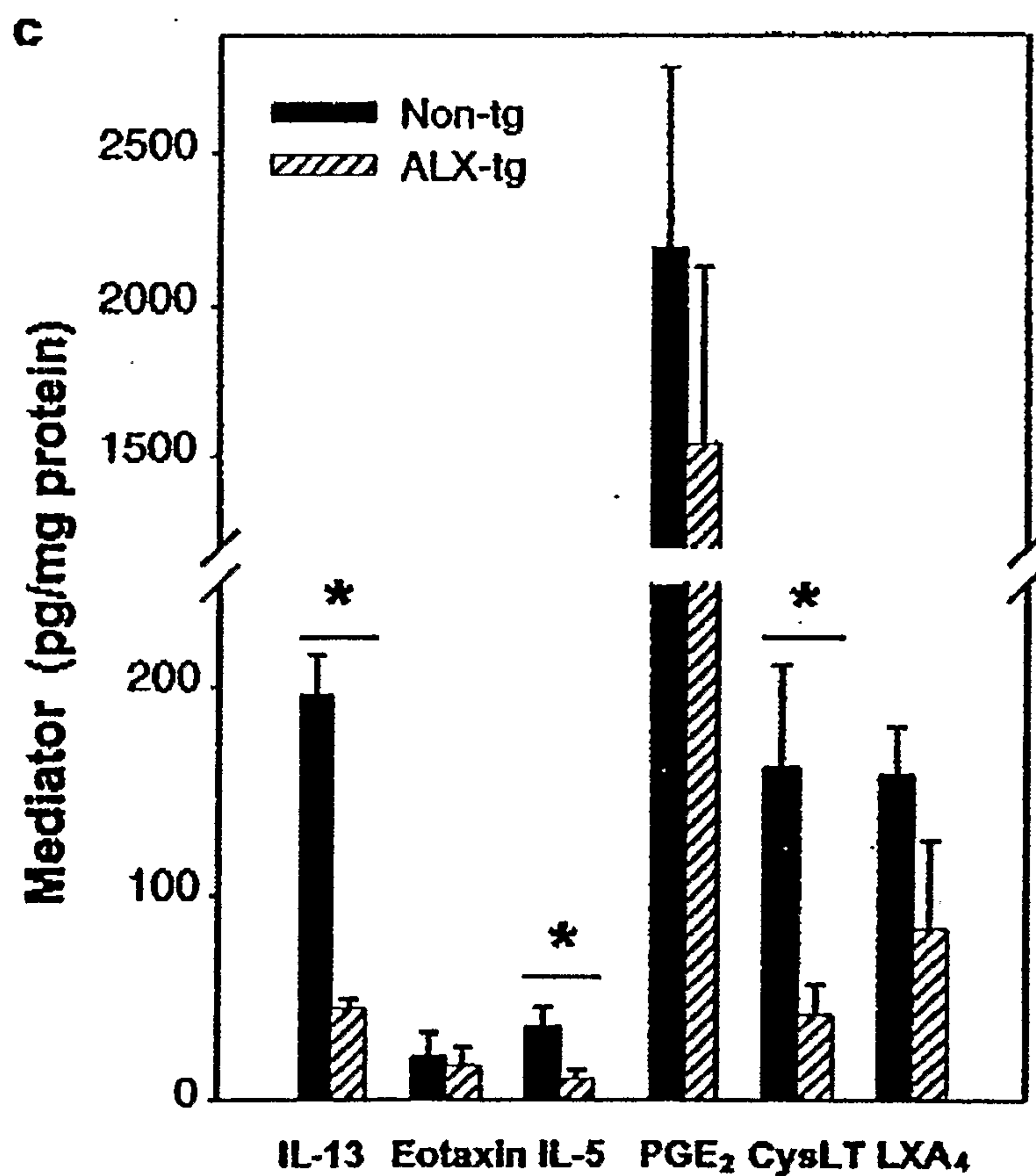
**Fig. 5b****Fig. 5c**

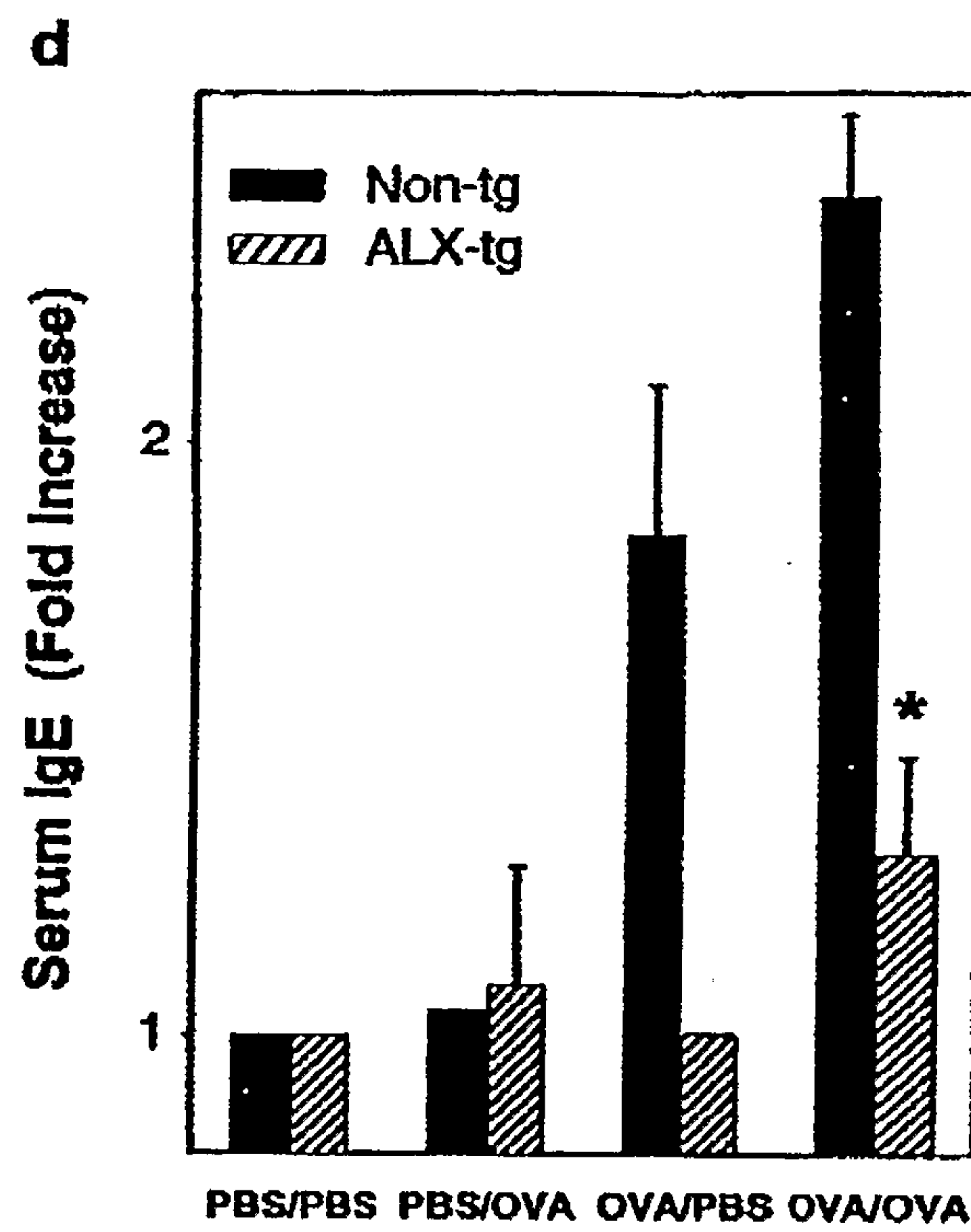
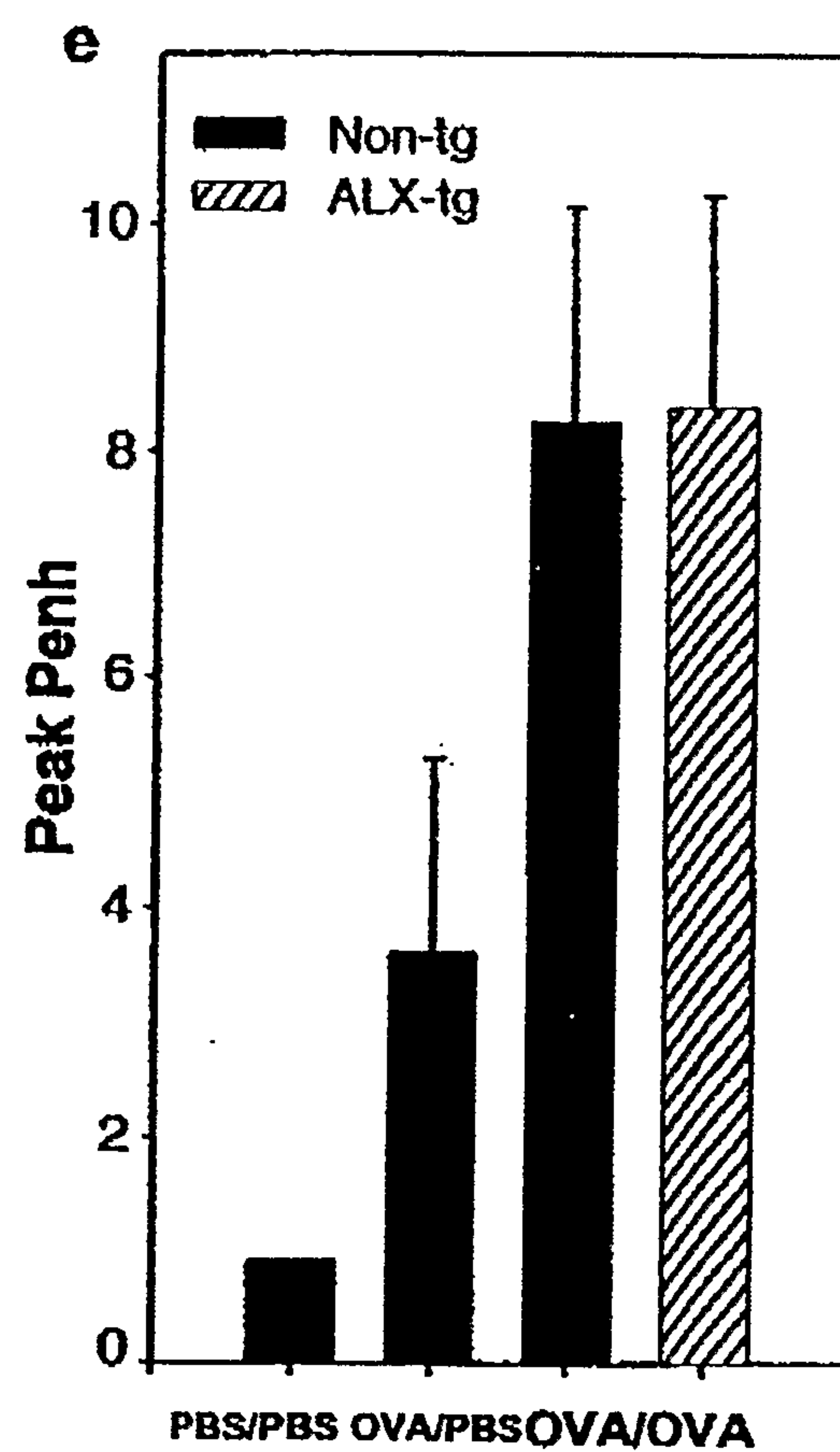
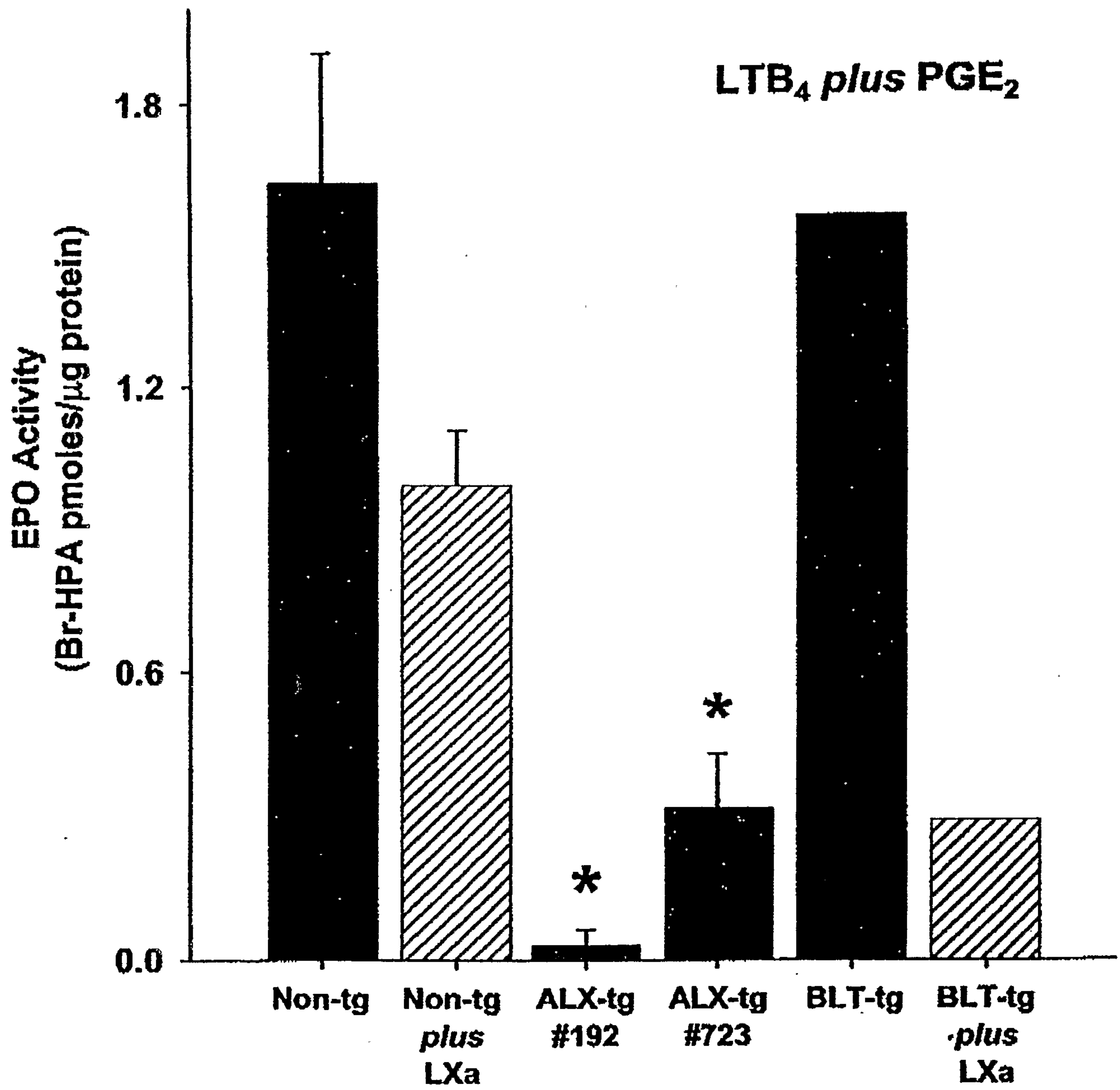
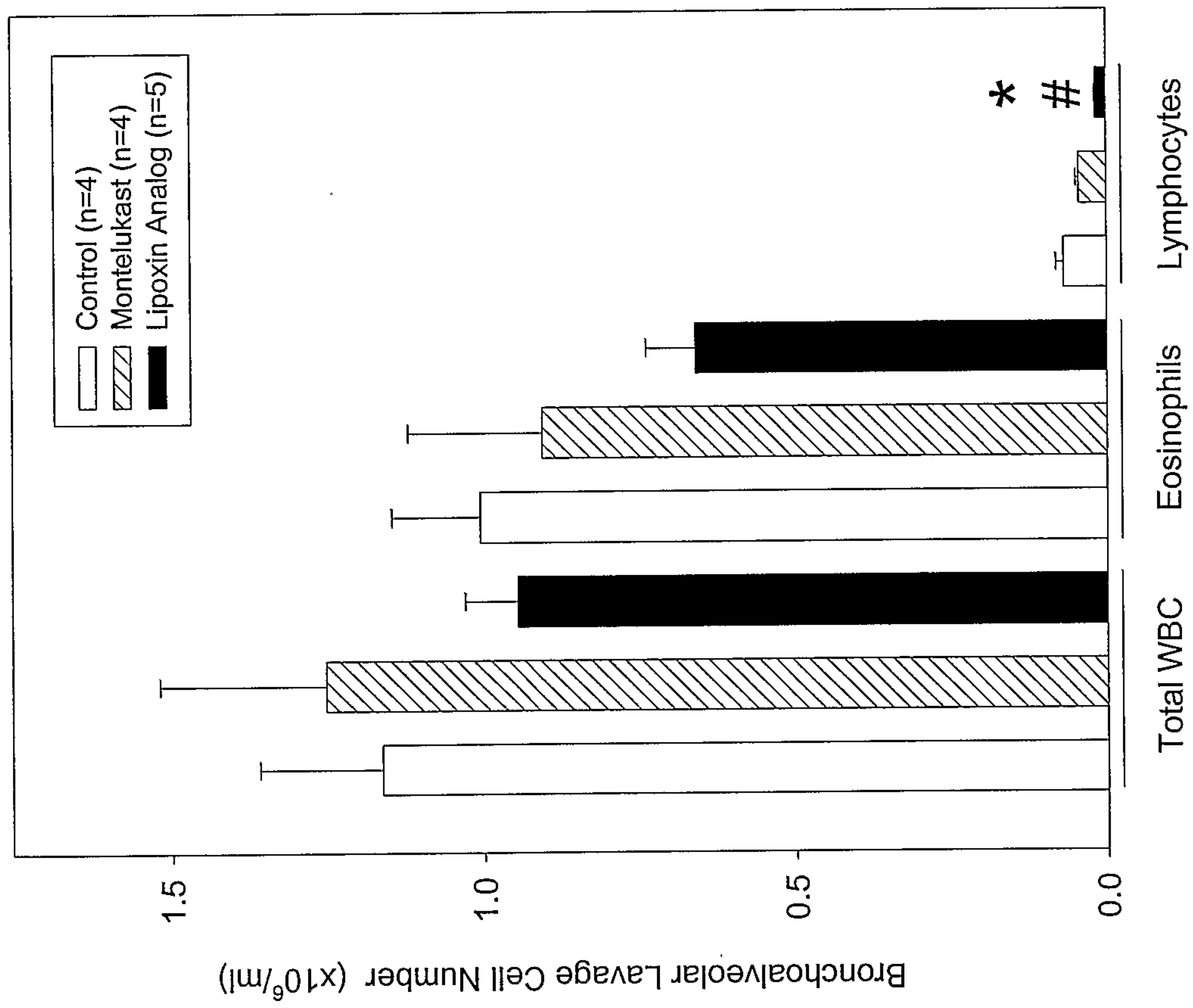
Fig. 5d**Fig. 5e**

Fig. 6

Regulation of Allergic Airway Inflammation By Lipoxin Analog Comparison with Montelukast

FIG. 7



* P < 0.01 LXa versus Control

P < 0.01 LXa versus ML

