

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

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
12 September 2002 (12.09.2002)

PCT

(10) International Publication Number
WO 02/070068 A2

(51) International Patent Classification⁷: A61P 9/00, (81) Designated States (national): AE, AG, AL, AM, AT, AU, 9/10, 17/02, 35/00 AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW.

(21) International Application Number: PCT/US02/06404

(22) International Filing Date: 1 March 2002 (01.03.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/272,931 2 March 2001 (02.03.2001) US
10/086,609 1 March 2002 (01.03.2002) US

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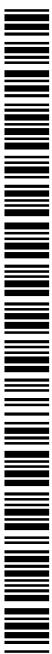
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(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



WO 02/070068 A2

(54) Title: LIPOXIN ANALOGS AS NOVEL INHIBITORS OF ANGIOGENESIS

(57) Abstract: The present invention is directed to methods for the prevention or inhibition of angiogenesis. The method is accomplished by the administration of an effective amount of 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄, LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ and pharmaceutically acceptable salts, esters, amides, carboxylic acids, or prodrugs thereof, to a subject in need thereof. As a consequence of the action of the therapeutic agent, angiogenesis is prevented or inhibited in the subject.

TITLE: Lipoxin Analogs as Novel Inhibitors of Angiogenesis**STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH**

The U.S. government may have rights in this invention pursuant to National Institute of Health Grant Nos. GM38765 and P01-DE13499.

5 RELATED APPLICATION

This application claims priority to U.S. Provisional Application No. 60/272,931, filed March 2, 2001, entitled "A Novel Inhibitor of Angiogenesis: Aspirin-Triggered-15R Lipoxin A₄", the contents of which are incorporated herein by reference.

10 BACKGROUND OF THE INVENTION

Angiogenesis is a fundamental process by which new capillaries are formed from existing blood vessels. This process plays important roles in physiological events such as formation of the corpus luteum, development of the embryo and wound healing, including recovery from both myocardial ischemia 15 and peptic ulcer (1). Unregulated growth of blood vessels can contribute to tissue injury in a large number of diseases such as arthritis, diabetes, and tumor progression (2). Endothelial cells are normally quiescent and are activated during the angiogenic response. Upon stimulation, endothelial cells can degrade their basement membrane and proximal extracellular matrix, migrate directionally, 20 then divide and organize into functional capillaries invested by a new basal lamina (3).

There is a growing body of evidence demonstrating that the angiogenic switch is regulated by the net balance between positive and negative regulators of new capillary growth (2). Persistence of neovascularization requires a pro- 25 angiogenic environment, with the expression of angiogenic factors outweighing that of angiostatic factors. A range of peptides can influence this balance, including mitogenic factors such vascular endothelial growth factor (VEGF) (3), nonmitogenic factors (selected cytokines, CXC chemokines), and internal peptide fragments of angiotatin and endostatin (3). Certain eicosanoids also have potent 30 biologic actions on vascular endothelial cells. In rabbits, PGE₂, PGR_{2a}, and prostacyclin (PGI₂) stimulate angiogenesis where prostaglandin E series, in particular PGE₁, is most potent. PGE₂ is a potent inducer of VEGF expression in

synovial fibroblasts. In addition to its known vasodilator and antiplatelet properties, PGI₂ can also induce VEGF gene expression and protein synthesis (4).

It was recently reported that 12-lipoxygenase activity and one of its products, 12(S)-HETE, is required for angiogenic responses (5), and that P450-derived 12R-HETE stimulates angiogenesis via NF-kB (6). The cyclooxygenase-2 (COX-2) gene in endothelial cells is rapidly upregulated by several growth factors as well as inducers of angiogenesis (7). Along these lines, results using three different endothelial cell models show that COX-2 is an essential component of angiogenesis, at least *in vitro* (8). Nonsteroidal anti-inflammatory drugs such as aspirin (ASA) have been implicated in the prevention of certain cancers such as lung and colon cancer (9, 10) that might be related to ASA's ability to reduce angiogenesis (7).

A need therefore exists, for compositions and methods to prevent angiogenesis that are directed toward the disease process, such that angiogenesis is prevented or inhibited physiologically. A need also exists for compositions and methods that induce angiogenesis in tissue that is lacking the requisite or essential physiological requirements for sustainability.

SUMMARY

Proliferative states such as chronic inflammation, ischemic diseases and cancer are often accompanied by intense angiogenesis, a highly orchestrated process involving vessel sprouting, endothelial cell migration, proliferation and maturation. Aspirin-triggered lipoxins (ATL), the 15R enantiomeric counterparts of lipoxins (LXs), are endogenous mediators generated during multicellular responses that display potent immunomodulatory actions. Surprisingly, it has been discovered that LXs, ATLs, and more specifically, the ATL stable analogs, 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (denoted ATL-1), LX_{A4}, 15-epi-LX_{A4} and 15-R/S-methyl, LX_{A4} are potent inhibitors of angiogenesis. For example, ATL-1, LX_{A4}, 15-epi-LX_{A4} and 15-R/S-methyl, LX_{A4} each inhibited endothelial cell proliferation in the 1-10 nM range by ~50% in cells stimulated with either vascular endothelial growth factor (VEGF) at 3 ng/ml or leukotriene D₄ (10 nM). In addition, ATL-1 (in a 10-100 nM range) inhibited VEGF (3

ng/ml) -induced endothelial cell chemotaxis. In a granuloma *in vivo* model of inflammatory angiogenesis, ATL-1 treatment (10 µg/mouse) reduced by ~50% the angiogenic phenotype, as assessed by both vascular casting and fluorescence. Together, these results identified a novel and potent previously unappreciated 5 action of aspirin-triggered 15-epi-LX.

In one aspect, the present invention pertains to methods for the prevention, reduction or inhibition of angiogenesis. The method is accomplished by the administration of an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or 10 prodrugs thereof, to a subject in need thereof. As a consequence of the action of the therapeutic agent, angiogenesis is prevented or inhibited in the subject.

In another aspect, the present invention also pertains to methods for the prevention or inhibition of angiogenesis. The method is accomplished by the administration of an effective amount of an aspirin triggered lipoxin (ATL) (15-15 epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof. As a consequence of the action of the therapeutic agent, angiogenesis is prevented or inhibited in the subject.

In still another aspect, the present invention pertains to methods for the 20 prevention or inhibition of solid tumor tissue growth undergoing neovascularization in a subject. The method is accomplished by the administration of an effective amount of an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject 25 in need thereof, to a subject in need thereof.

In another aspect, the present invention pertains to methods for the 30 prevention or inhibition of solid tumor tissue growth undergoing neovascularization in a subject. The method is accomplished by the administration of an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

5 In yet another aspect, the present invention is directed to methods to inhibit or prevent neovascularization from occurring in a subject. The method is accomplished by the administration of an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

10 In yet another aspect, the present invention is directed to methods to inhibit or prevent neovascularization from occurring in a subject. The method is accomplished by the administration of an effective amount of an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

15 The invention is also directed to methods for treating a subject in which neovascularization is occurring in retinal tissue. The neovascularization in the retinal tissue can be prevented or inhibited by administering an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

20 The present invention is further directed to methods for treating a subject in which neovascularization is occurring in retinal tissue. The neovascularization in the retinal tissue can be prevented or inhibited by administering an effective amount of an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

25 The invention is further directed to methods for treating a subject for restenosis in tissue wherein smooth muscle cell migration occurs following angioplasty. The restenosis can be prevented or inhibited by administering an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

30 The invention is directed to methods for treating a subject for restenosis in tissue wherein smooth muscle cell migration occurs following angioplasty. The restenosis can be prevented or inhibited by administering an effective amount of

an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

In still another embodiment, the invention pertains to methods of reducing
5 blood supply to tissue required to support new growth of the tissue in a subject. This reduction or elimination of new undesired growth of tissue can be accomplished by the administration of a composition comprising an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject
10 in need thereof.

The invention further pertains to methods of reducing blood supply to tissue required to support new growth of the tissue in a subject. This reduction or elimination of new undesired growth of tissue can be accomplished by the administration of a composition comprising an effective amount of an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof.

Further, the present invention pertains to methods for the prevention, diminishment or inhibition the production of new vessels in a subject associated with or stimulated by the production or release of VEGF. The method is
20 accomplished by the administration of an effective amount of a therapeutic agent, including LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof. Alternatively, an effective amount of an aspirin triggered lipoxin
25 (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof can be utilized. As a consequence of the action of the therapeutic agent, growth of new vessels associated with the production of VEGF is prevented or inhibited and thus the growth of endothelial cells in the subject. For example, VEGF is
30 associated with tumor genesis, lymphangiogenesis and proliferative disorders. The invention therefore, can be used to inhibit, reduce or prevent metasasis of tumors.

Surprisingly, configurational isomers of LXA₄, LXA₄ analogs and ATL analogs, LXB₄ and LXB₄ analogs and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, provide the opposite effects with regard to revascularization of tissue by the above-identified compounds of the invention.

5 That is, it has been surprisingly discovered that LXB₄ and LXB₄ analogs have the ability to stimulate regeneration and ingrowth of vascular or epithelial tissue in tissues that are in need of such stimulation. This is especially important in tissue grafting, tissue engineering and prosthetic group sites of attachment.

BRIEF DESCRIPTION OF THE DRAWINGS

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

Figure 1 demonstrates that ATL-1 inhibits VEGF-stimulated HUVEC proliferation. HUVEC (5×10^3) were plated in 96-well culture plates and cell proliferation was stimulated with 3 ng/ml VEGF. Three days after treatment, cell numbers were measured using MTT assay. Results are expressed as percent inhibition of proliferation relative to vehicle and represent mean \pm SE for four independent experiments performed in triplicate. *Inset:* Representative experiment showing the time course of cell proliferation induced by 3 ng/ml (filled triangle) or 10 ng/ml (filled square) VEGF. Vehicle (open circles) and 15 ATL-1 (100 nM) (open triangle).

Figure 2 demonstrates that ATL-1, as well as LXA₄, 15-epi-LXA₄ and 15-R/S-methyl, LXA₄ each inhibit endothelial cell chemotaxis. (A) Chemotaxis was initiated by addition of VEGF (3 ng/ml) or ATL-1 (100 nM) to the lower compartment of a 48-well chemotaxis chamber. Results are expressed as percent of cell migration compared to vehicle alone and represent mean \pm SE for three independent experiments performed in triplicate ($P < 0.05$). (B) HUVEC were incubated with vehicle or indicated concentrations of ATL-1 (15 min, 37° C) and added to the upper compartment of the microchamber (1×10^6 /well). Chemotaxis was initiated by addition of VEGF (3 ng/ml) to the lower compartment. Results are expressed as percent inhibition of migration relative to VEGF for a representative experiment performed in triplicate.

Figure 3 is a graphical representation demonstrating inhibition of HUVEC proliferation. (A) ATL-1 inhibits LTD₄-stimulated HUVEC proliferation. HUVECs (5×10^3) were plated in 96-well culture plates, cell proliferation was stimulated with 10 nM LTD₄, and cell numbers were determined after 3 days using MTT. (B), concentration dependent cell proliferation induced by LTD₄ and LTB₄. Results are expressed as mean \pm S.E. for four independent experiments performed in triplicate.

Figure 4 demonstrates that ATL-1 inhibits angiogenic phenotype *in vivo*. (A) Vascular index (VI = mg carmine dye/mg weight of tissue) in day-6 murine air pouch. Animals received a local injection of ATL-1 (10 μ g /pouch) or vehicle immediately before VEGF (1 μ g/pouch), 24 h after raising the pouch. Results are expressed as the mean \pm SE for n=4 animals per group. *Denotes statistically significant difference ($P<0.05$) from VEGF alone; (B) Cedar wood oil histology of air pouch. Carmine dye vascular casts were made in day-6 air pouch from mice treated locally with vehicle, ATL-1 (10 μ g), VEGF (1 μ g) or VEGF plus ATL-1. Tissue was fixed in ethanol and cleared in cedarwood oil.

Figure 5 depicts Anti-angiogenic action of ATL-1: fluorescent microscopy. Representative fluorescence photomicrographs showing the anti-angiogenic action of ATL-1 (10 μ g/pouch) in the murine air pouch (see methods).

Figure 6 are photomicrographs of a murine air pouch. Immunohistochemistry for murine air pouch CD31. Paraffin-embedded air pouch sections were stained for CD31 from mice as in Figures 4 and 5 and treated with vehicle alone (A), analog alone (B), VEGF-treated mice (C), and VEGF plus ATL-treated mice (D). Results are representative of eight separate mice each in duplicate. Magnification is 200 X power and horseradish peroxidase with hematoxylin counterstain.

DETAILED DESCRIPTION

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 invention.

Abbreviations used throughout the present application include the following and are included here for convenience. ASA, aspirin; ATL, aspirin-triggered 15-epi-lipoxins; ATL-1, 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄; COX, cyclooxygenase; HETE, hydroxyeicosatetraenoic acid; HUVEC, human umbilical vein endothelial cells; IL, interleukin; LO, lipoxygenase; LT, leukotriene; LX, lipoxin; LXA₄, 5*S*, 6*R*, 15*S*-trihydroxy-7, 9, 13-*trans*-11-*cis*-eicosatetraenoic acid; 15-epi-LXA₄, 5*S*, 6*R*, 15*R*-trihydroxy-7, 9, 13-*trans*-11-*cis*-eicosatetraenoic acid; 15-*R/S*-methyl, LXA₄, 5*S*, 6*R*, 15*R/S*-trihydroxy-15-methyl-7, 9, 13-*trans*-11-*cis*-eicosatetraenoic acid, methyl ester; LXB₄, 5*S*, 14*R*, 15*S*-trihydroxy-6, 8, 12-*trans*-10-*cis*-eicosatetraenoic acid; MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium; PBS, phosphate buffered saline; PG, prostaglandin; PMN, neutrophils; VEGF, vascular endothelial growth factor.

It is to be understood, that throughout the present specification, reference is often made to the therapeutic compounds of the inventions as esters, for example, ATL-1 as a carboxylic ester, i.e., methyl ester. However, all pharmaceutically acceptable salts, esters, amides, and prodrugs, including the carboxylic acid, are considered within the scope of the invention for the LXA₄, ATL and LXB₄ compounds. For convenience, this terminology has been minimized throughout the description but should be considered as part of the invention. Additionally, it should be understood that the terms LXA₄, 15-epi-LXA₄ and 15-*R/S*-methyl, LXA₄ also include all pharmaceutically acceptable salts, esters, amides, prodrugs and carboxylic acids.

Additionally, the hydroxyl(s) of ATLs, LXA₄s, and LXB₄s can be protected by various protecting groups, such as those known in the art. An artisan skilled in the art can readily determine which protecting group(s) can be useful for the protection of the hydroxyl group(s). Standard methods are known in the art and are more fully described in literature. For example, suitable protecting groups can be selected by the skilled artisan and are described in Green and Wuts, "Protecting Groups in Organic Synthesis", John Wiley and Sons, 1991, the teachings of which are incorporated herein by reference.

Preferred protecting groups include TMS or TIPPS groups, and preferably acetate or propionate groups.

For example, one or more hydroxyl groups can be treated with a mild base, such as triethylamine in the presence of an acid chloride or silyl chloride to 5 facilitate a reaction between the hydroxyl ion and the halide. Alternatively, an alkyl halide can be reacted with the hydroxyl ion (generated by a base such as lithium diisopropyl amide) to facilitate ether formation.

It should also be understood that not all hydroxyl groups need be 10 protected. One, two or all three hydroxyl groups can be protected. This can be accomplished by the stoichiometric choice of reagents used to protect the hydroxyl groups. Methods known in the art can be used to separate the mono, di- or tri-protected hydroxy compounds, e.g., HPLC, LC, flash chromatography, gel permeation chromatography, crystallization, distillation, etc.

One advantage of protecting one or more hydroxyl groups of ATL, LXA₄, 15 or LXB₄ compounds, e.g., via acetates, is the ability to delay the complete metabolic uptake of the compound(s). This is one means by which the compound(s) can remain active over a prolonged period of time as the subject's body slowly removes the protecting group from the hydroxyl under normal physiological conditions. Additionally, by protecting one or more of the 20 hydroxyl groups of these compounds, hydrolysis of the protecting group allows the medication to enter the biochemical pathway of the subject prior to degradation of the parent, unprotected, compound.

Methods to prepare lipoxin analogs (ATLs, LXA₄s, or LXB₄s) are known 25 in the art. For example, U.S. Patent Nos. 4,576,758, 4,560,514, 5,079,261, 5,049,681, 5,441,951, 5,648,512, 5,650,435, 6,048,897, 6,100,296, 6,177,468 and 6,316,648 and Japanese Patent Nos. 3,227,922, 63,088,153, 62,198,677 and 1,228,994 describe approaches to prepare lipoxin analogs. Publications by K.C. Nicolaou et al. include approaches to various lipoxin compounds. (For example see, Nicolaou, K.C. et al. *Biochim. Biophys. Acta* 1003:44-53; Nicolaou, K.C. et 30 al. *J. Org. Chem.* 54:5527-5535; and Nicolaou, K.C. *Angew. Chem. Int. Ed. Engl.* 30:1100-1116. Additional literature references for the preparation of lipoxin analogs include Takano, T., S. Fiore, J.F. Maddox, H.R. Brady, N.A.

Petasis, and C.N. Serhan. 1997. Aspirin-triggered 15-epi-lipoxin A₄ and LXA₄ stable analogs are potent inhibitors of acute inflammation: Evidence for anti-inflammatory receptors. *J. Exp. Med.* 185:1693-1704 and Serhan, Charles N., Maddox, Jane F., Petasis, Nicos A., Akritopoulou-Zanze, Irini, Papayianni, 5 Aikaterina, Brady, Hugh R., Colgan, Sean P., and Madara, James L. (1995), Biochemistry, 34, pp.14609-14614.

Aspirin's therapeutic mechanism of action includes inhibition of COX-derived prostanoids (10). It was discovered that COX-2, when acetylated by ASA, blocks the ability of COX-2 to generate prostanoids, yet this enzyme 10 remains active in endothelial cells, epithelial cells and mononuclear cells and initiates the biosynthesis of new products of cell-cell interactions or transcellular biosynthesis termed aspirin-triggered-15-epi-lipoxins (ATLs) (11). These novel endogenous lipid mediators are the carbon 15 epimers of LX that carry their 15 alcohol in the *R* configuration compared to their native lipoxin (LX) counterparts 15 and appear to mimic most if not all endogenous LX bioactivities.

To date the actions of ATLs appear to be most relevant in regulating inflammatory responses, since they are generated during cell-cell interactions that can involve, for example, endothelial cells-neutrophils *in vivo* (12), and display potent inhibitory actions in several key and strategic events in inflammation (12-20). Both LX and ATL actions include inhibiting adhesion and transmigration of neutrophils, and hence can serve as counterregulatory signals to limit and/or regulate leukocyte accumulation that are potentially operative in the dampening and resolution of inflammatory sites (14). Since LX are rapidly generated and inactivated in the local microenvironment, to investigate these actions *in vivo*, 25 stable analogs of both lipoxins, i.e., LXA₄ and ATL were designed that enhance bioavailabilities and these natural compounds bioactivities compared to their native products (14) and also proved to be ~100 times the potency of ASA (13). The present invention establishes that LXA₄s and ATLs can regulate angiogenesis, a previously unknown and surprising application of these 30 compounds. For example, using a metabolically more stable ATL synthetic analog [15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ (denoted ATL-1)], ATLs and LXA₄ compounds, including, 15-epi-LXA₄ or 15-*R/S*-methyl, LXA₄ and LXA₄ proved to be potent angiostatic eicosanoid *in vivo*, identifying a new

activity for these endogenous mediators that is in sharp contrast to the actions of other eicosanoids and is relevant in several human diseases.

In one aspect, the present invention pertains to methods for the prevention, diminishment or inhibition of angiogenesis. The method is 5 accomplished by the administration of an effective amount of LXA₄ and analogs thereof, such as 15-R/S methyl, LXA₄, and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof. As a consequence of the action of the therapeutic agent, angiogenesis is prevented, reduced or inhibited in the subject. More specifically, the therapeutic agents can be used in 10 the treatment of the disease states and conditions of the angiogenic disease processes as described below. More specifically, the LXA₄ and ATL therapeutic compounds described throughout the specification can be used for the treatment of restenosis, solid tumor tissue growth, neovascularization, e.g., retinal tissue, and reducing blood supply to tissue required to support new growth of tissue in a 15 subject.

In another aspect, the present invention also pertains to methods for the prevention, reduction or inhibition of angiogenesis in tissue of a subject. The method is accomplished by the administration of an effective amount of an aspirin triggered lipoxin (ATL) (15-epi-LXA₄, such as 15-epi-16-(*para*-fluoro)- 20 phenoxy-lipoxin A₄ (ATL-1)), and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, to a subject in need thereof. As a consequence of the action of the therapeutic agent, angiogenesis is prevented or inhibited in the subject.

The invention thus provides for a method for the general inhibition of 25 angiogenesis in tissue, and thereby inhibits or prevent events in the tissue which depend upon angiogenesis. Generally, the method comprises administering to the tissue a composition comprising an angiogenesis-inhibiting amount of, for example, ATL-1, LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄.

As used herein, the term "angiogenesis" means the formation of new 30 blood vessels into a tissue or organ. Under normal physiological conditions, humans or animals only undergo angiogenesis in very specific restricted situations. For example, angiogenesis is associated with wound healing, fetal and

embryonal development and formation of the corpus luteum, endometrium and placenta. The biochemical aspects of angiogenesis are associated with a highly regulated system of angiogenic stimulators and inhibitors. Controlled angiogenesis has been found to be altered in certain disease states and, in many 5 cases, the pathological damage associated with the disease is related to the uncontrolled angiogenesis.

The bioprocesses of controlled and uncontrolled angiogenesis are thought to occur in a similar manner. Endothelial cells and pericytes, surrounded by a basement membrane, form capillary blood vessels. Erosion of the basement 10 membrane promotes angiogenesis by enzymes released by endothelial cells and leukocytes. The endothelial cells, which line the lumen of blood vessels, then break through the basement membrane. Angiogenic stimulants induce the endothelial cells to migrate through the eroded basement membrane. The migrating cells form an offshoot from the parent blood vessel, where the 15 endothelial cells undergo mitosis and proliferate. The endothelial offshoots can merge with each other to form capillary loops, creating a new blood vessel. In the disease state, prevention of angiogenesis could avert the damage caused by the invasion of the new microvascular system.

Persistent, unregulated angiogenesis can occur in a multiplicity of disease 20 states, tumor metastasis and abnormal growth by endothelial cells and supports the pathological damage seen in these conditions. The diverse pathological states created due to unregulated angiogenesis have been grouped together as angiogenic dependent or angiogenic associated diseases. The present invention provides therapies that are directed to control the angiogenic processes thus 25 leading to the abrogation or mitigation of these diseases. With the exception of traumatic wound healing, corpus leuteum formation and embryogenesis, it is believed that angiogenesis processes are associated with undesired, and often life threatening, disease processes and therefore, the use of the present therapeutic methods are selective for the disease, i.e., angiogenesis, and do not have 30 deleterious side effects.

The following angiogenic diseases can be treated according to the present invention by use of the afore-mentioned ATLs, such as ATL-1 or LXA₄s such

as 15-R/S-methyl, LXA₄. These angiogenic diseases include, but are not limited to, the following:

One example of a disease mediated by angiogenesis is ocular neovascular disease. This disease is characterized by invasion of new blood vessels into the 5 structures of the eye such as the retina or cornea. It is perhaps, one of the most common causes of blindness and is involved in over twenty eye diseases. For example, in age-related macular degeneration, the associated visual problems are caused by an ingrowth of chorioidal capillaries through defects in Bruch's membrane with proliferation of fibrovascular tissue beneath the retinal pigment 10 epithelium. Angiogenic damage is also associated with diabetic retinopathy, retinopathy of prematurity, corneal graft rejection, neovascular glaucoma and retrobulbar fibroplasia. Other diseases associated with corneal neovascularization include, but are not limited to, Vitamin A deficiency, contact lens overwear, atopic keratitis, superior limbic keratitis, epidemic keratoconjunctivitis, 15 pterygium keratitis sicca, sjogrens, acne rosacea, lipid degeneration, chemical burns, bacterial ulcers, fungal ulcers, phlyctenulosis, syphilis, Mycobacteria infections, Herpes simplex infections, Herpes zoster infections, Wegener's sarcoidosis, Scleritis, Steven's Johnson syndrome, periphigoid radial keratotomy, protozoan infections, Kaposi sarcoma, Mooren ulcer, Terrien's marginal 20 degeneration, marginal keratolysis, rheumatoid arthritis, systemic lupus, polyarteritis, trauma, and corneal graft rejection.

Diseases associated with retinal/choroidal neovascularization include, but are not limited to, diabetic retinopathy, vein occlusion, artery occlusion, carotid 25 obstructive disease, chronic uveitis/vitritis, mycobacterial infections, Lyme's disease, systemic lupus erythematosus, macular degeneration, sickle cell anemia, sarcoid, syphilis, pseudoxanthoma elasticum, Pagets disease, retinopathy of prematurity, Eales disease, Bests disease, myopia, optic pits, Stargarts disease, pars planitis, chronic retinal detachment, hyperviscosity syndromes, toxoplasmosis, Bechets disease, infections causing a retinitis or choroiditis, 30 presumed ocular histoplasmosis, trauma and post-laser complications. Other diseases include, but are not limited to, diseases associated with rubeosis and diseases caused by the abnormal proliferation of fibrovascular or fibrous tissue including all forms of proliferative vitreoretinopathy.

An even more prevalent disease in which angiogenesis is believed to be involved is rheumatoid arthritis. For example, the blood vessels in the synovial lining of the joints undergo angiogenesis. In addition to forming new vascular networks, the endothelial cells release factors and reactive oxygen species that 5 lead to pannus growth and cartilage destruction. It is believed that the factors involved in angiogenesis can actively contribute to, and help maintain, the chronically inflamed state of rheumatoid arthritis.

It is believed that factors associated with angiogenesis can also have a role in osteoarthritis. The activation of the chondrocytes by angiogenic-related 10 factors contributes to the destruction of the joint. At a later stage, the angiogenic factors can promote new bone formation. The present invention provides therapeutic intervention that prevents the bone destruction and can halt the progress of the disease and provide relief for persons suffering with arthritis.

Both ulcerative colitis and Crohn's disease are known to have histological 15 changes with the ingrowth of new blood vessels into the inflamed tissues. Bartonellosis, can result in a chronic stage that is characterized by proliferation of vascular endothelial cells. An even more insidious pathological role associated with angiogenesis is found in arteriosclerosis. The plaquing of the lumen of blood vessels has been shown to have angiogenic stimulatory activity.

20 A frequent angiogenic disease of childhood is hemangioma. Generally, the tumors associated with the disease are benign and regress without intervention. In more severe cases, the tumors grow and create clinical complications. Systemic forms of hemangiomas, the hemangiomatoses, have a high mortality rate. Therapy-resistant hemangiomas exist that cannot be treated 25 with therapeutics currently in use.

Angiogenesis is also responsible for damage found in hereditary diseases such as Osler-Weber-Rendu disease, or hereditary hemorrhagic telangiectasia. These diseases are characterized by multiple small angiomas, tumors of blood or lymph vessels. The angiomas are found in the skin and mucous membranes, 30 often accompanied by epistaxis (nosebleeds) or gastrointestinal bleeding and sometimes with pulmonary or hepatic arteriovenous fistula.

Of great concern is the disease state(s) associated with cancer(s). Often times, the cancer is associated with angiogenesis and is identified by solid tumor formation and metastasis. Angiogenic factors are associated with several solid tumors such as neuroblastoma, rhabdomyosarcomas, retinoblastoma, Ewing 5 sarcoma, and osteosarcoma. It is known that a tumor cannot expand without a blood supply to provide nutrients and remove cellular wastes. Tumors in which angiogenesis is important include solid tumors, and benign tumors such as acoustic neuroma, neurofibroma, trachoma and granulomas. Prevention or inhibition of angiogenesis could prevent or halt the growth of these tumors and 10 the subsequent degenerative condition due to the presence of the tumor.

Angiogenesis has also been associated with blood-born tumors including leukemias, any of the various acute or chronic neoplastic diseases of bone marrow in which unrestrained proliferation of white blood cells occurs, usually accompanied by anemia, impaired blood clotting, and enlargement of the lymph 15 nodes, liver, and spleen. It is believed that angiogenesis is significant as a caustive factor in the abnormalities in the bone marrow that give rise to leukemia-like tumors.

Angiogenesis is important in two stages of tumor metastasis. The first stage where angiogenesis stimulation is important is in the vascularization of the 20 tumor which allows tumor cells to enter the blood stream and to circulate throughout the body. Once the tumor cells leave the primary site, and find a secondary metastasis site, angiogenesis must occur before the new tumor can grow and expand. Therefore, prevention of angiogenesis could prevent metastasis of tumors and contain the neoplastic growth at the primary site.

25 In a related embodiment, the present invention can be used in combination with other therapies such as conventional chemotherapy directed against solid tumors and metastases. ATLs, such as ATL-1 or LXA₄s, such as 15-R/S-methyl, LXA₄, can be administered during or after chemotherapy. In a preferred embodiment, the drug should be administered when the tumor tissue is 30 responding to the toxic assault when vascular tissue is being reorganized to supply blood and nutrients to the tumor tissue. Additionally, the use of ATLs, such as ATL-1 or LXA₄s can be used as a prophylactic treatment after surgical removal of a tumor to prevent angiogenesis from occurring at the treatment site.

Knowledge of the role of angiogenesis in the maintenance and metastasis of tumors has led to a prognostic indicator for breast cancer. The amount of neovascularization found in the primary tumor was determined by counting the microvessel density in the area of the most intense neovascularization in invasive 5 breast carcinoma. A high level of microvessel density was found to correlate with tumor recurrence. Control of angiogenesis by therapeutic means could possibly lead to cessation of the recurrence of the tumors.

Angiogenesis is also involved in normal physiological processes such as reproduction and wound healing. Angiogenesis is an important step in ovulation 10 and also in implantation of the blastula after fertilization. Prevention of angiogenesis could be used to induce amenorrhea, to block ovulation or to prevent implantation by the blastula.

In wound healing, excessive repair or fibroplasia can be a detrimental side effect of surgical procedures and can be caused or exacerbated by angiogenesis. 15 Adhesions are a frequent complication of surgery and lead to problems such as small bowel obstruction.

Restenosis is a process of smooth muscle cell (SMC) migration and proliferation at the site of percutaneous transluminal coronary angioplasty which hampers the success of angioplasty. The migration and proliferation of SMC's 20 during restenosis can be considered a process of angiogenesis which is inhibited by the present methods. Therefore, the invention also contemplates inhibition, reduction or prevention of restenosis by inhibiting, reducing or preventing angiogenesis according to the present methods in a subject following angioplasty procedures. For inhibition or prevention of restenosis, an ATL, such as ATL-1 or 25 an LXA₄, such as 15-R/S-methyl, LXA₄, can be administered, preferably via intravenous injection, several days before the operation or after the angioplasty procedure for from about 2 to about 28 days, and more typically for about the first 14 days following the procedure.

The term "subject" as used herein refers to any living organism in which 30 an angiogenic response is elicited. The term subject includes, but is not limited to, humans, nonhuman primates such as chimpanzees and other apes and monkey species; farm animals such as cattle, sheep, pigs, goats and horses; domestic

mammals such as dogs and cats; laboratory animals including rodents such as mice, rats and guinea pigs, and the like. The term does not denote a particular age or sex. Thus, adult and newborn subjects, as well as fetuses, whether male or female, are intended to be covered.

5 The term “mammal” as used herein refers to a living organism capable of eliciting an immune response to an antigen. The term subject includes, but is not limited to, nonhuman primates such as chimpanzees and other apes and monkey species, sheep, pigs, goats, horses, dogs, cats, mice, rats and guinea pigs, and the like.

10 The 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 can 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, 15 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 which is incorporated herein by reference).

20 The term “prodrug” refers to compounds that are rapidly transformed in vivo to yield the parent compound of the above formulae, for example, by hydrolysis in blood. A thorough discussion is provided in T. Higuchi and V. Stella, “Pro-drugs as Novel Delivery Systems,” Vol. 14 of the A.C.S. Symposium Series, and in Bioreversible Carriers in Drug Design, ed. Edward B. Roche,

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American Pharmaceutical Association and Pergamon Press, 1987, both of which are hereby incorporated by reference. As used herein, a prodrug is a compound that, upon *in vivo* administration, is metabolized or otherwise converted to the biologically, pharmaceutically or therapeutically active form of the compound.

5 To produce a prodrug, the pharmaceutically active compound is modified such that the active compound will be regenerated by metabolic processes. The prodrug can be designed to alter the metabolic stability or the transport characteristics of a drug, to mask side effects or toxicity, to improve the flavor of a drug or to alter other characteristics or properties of a drug. By virtue of

10 knowledge of pharmacodynamic processes and drug metabolism *in vivo*, once a pharmaceutically active compound is identified, those of skill in the pharmaceutical art generally can design prodrugs of the compound [see, e.g., Nogradi (1985) *Medicinal Chemistry A Biochemical Approach*, Oxford University Press, New York, pages 388-392]. Conventional procedures for the

15 selection and preparation of suitable prodrug derivatives are described, for example, in "Design of Prodrugs," ed. H. Bundgaard, Elsevier, 1985. Suitable examples of prodrugs include methyl, ethyl and glycerol esters of the corresponding acid.

The compounds of the invention can be formulated into pharmaceutical compositions as described, *vide infra*. In a preferred embodiment, the compound can be administered over an extended period of time in a sustained release composition. Sustained release compositions are known in the art and one skilled in the art can formulate an acceptable composition based on generally recognized parameters in the art. In a most preferred embodiment, the glycerol ester can be used in the treatment of inflammatory conditions, described herein, in sustained release compositions, i.e., a transdermal patch, as known in the art. Suitable methods to prepare a transdermal patch can be found in U.S. Patent No., 5,814,599, 5,846,974 or 4,201,211, the contents of which are incorporated herein by reference. More particularly, the compounds can be delivered transdermally using the types of patch technologies available from Ciba-Geigy Corporation and Alza Corporation. The administration of the pharmaceutical compositions of the present invention can be intermittent, or at a gradual, continuous, constant or controlled rate to a warm-blooded animal, such as a human being. In addition, the time of day and the number of times per day that the pharmaceutical

formulation is administered can vary. Administration preferably is such that the active ingredients of the pharmaceutical formulation interact with the inflammatory condition.

A “therapeutically effective amount” is an amount of an ATL, such as
5 ATL-1 or an LXA₄, such as 15-R/S-methyl, LXA₄, sufficient to produce a measurable inhibition, reduction or prevents angiogenesis in the tissue being treated, i.e., an angiogenesis-inhibiting amount. Inhibition of angiogenesis can be measured in situ by immunohistochemistry or by other methods known to one skilled in the art such as measurement by FAC analysis that monitors P-selectin
10 or VEGF receptors. See also Figures 4 and 5.

More specifically, the pharmaceutical compositions of the invention can include a “therapeutically effective amount” or a “prophylactically effective amount” of an antiangiogenic of the invention. A “therapeutically effective amount” refers to an amount effective, at dosages and for periods of time
15 necessary, to achieve the desired therapeutic result, e.g., a diminishment, reduction or prevention of angiogenic factors associated with various disease states or conditions. A therapeutically effective amount of the antiangiogenic can vary according to factors such as the disease state, age, sex, and weight of the individual, and the ability of the antiangiogenic to elicit a desired response in the
20 individual. A therapeutically effective amount is also one in which any toxic or detrimental effects of the antibody or antibody portion are outweighed by the therapeutically beneficial effects.

A “prophylactically effective amount” refers to an amount effective, at dosages and for periods of time necessary, to achieve the desired prophylactic result, i.e. prevent. Typically, since a prophylactic dose is used in subjects prior to or at an earlier stage of disease, the prophylactically effective amount will be less than the therapeutically effective amount.
25

Dosage regimens can be adjusted to provide the optimum desired response (e.g., a therapeutic or prophylactic response). For example, a single
30 bolus can be administered, several divided doses can be administered over time or the dose can be proportionally reduced or increased as indicated by the exigencies of the therapeutic situation. It is especially advantageous to formulate

parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the mammalian subjects to be treated; each unit containing a predetermined quantity of active compound calculated to 5 produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the invention are dictated by and directly dependent on (a) the unique characteristics of the active compound and the particular therapeutic or prophylactic effect to be achieved, and (b) the limitations inherent in the art of compounding such an 10 active compound for the treatment of sensitivity in individuals.

An exemplary, non-limiting range for a therapeutically or prophylactically effective amount of an antiangiogenic of the invention is 0.1-20 mg/kg, more preferably 1-10 mg/kg. It is to be noted that dosage values can vary with the type and severity of the condition to be alleviated. It is to be further understood that 15 for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions, and that dosage ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed composition.

20 The antiangiogenic compounds of the invention, e.g., an ATL, such as ATL-1 or an LXA₄, such as 15-*R/S*-methyl, LXA₄, can be incorporated into pharmaceutical compositions suitable for administration to a subject. Typically, the pharmaceutical composition comprises an antiangiogenic of the invention and a pharmaceutically acceptable carrier. As used herein, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like that are physiologically compatible. Examples of pharmaceutically acceptable carriers include one or more of water, saline, phosphate buffered saline, dextrose, glycerol, ethanol and the like, as well as combinations thereof. 25 In some instances, it can be beneficial to include isotonic agents, for example, sugars, polyalcohols such as mannitol, sorbitol, or sodium chloride in the composition. Pharmaceutically acceptable carriers can further comprise minor amounts of auxiliary substances such as wetting or emulsifying agents, 30

preservatives or buffers, which enhance the shelf life or effectiveness of the antiangiogenic.

Additionally, the mono-, di- and/or tri- protected alcohols of the compounds of the invention provide for sustained release of the 5 inhibitory/preventative compound. For example, the tri-acyl analogs of the invention provide for such a sustained release of the hydrolyzed tri-hydroxy alcohol(s). The triacylated analogs are thus de-esterified in the blood of the subject.

The antiangiogenics of the invention can be incorporated into a 10 pharmaceutical composition suitable for parenteral administration. Other suitable buffers include but are not limited to, sodium succinate, sodium citrate, sodium phosphate or potassium phosphate. Sodium chloride can be used to modify the toxicity of the solution at a concentration of 0-300 mM (optimally 150 mM for a liquid dosage form). Cryoprotectants can be included for a lyophilized dosage 15 form, principally 0-10% sucrose (optimally 0.5-1.0%). Other suitable cryoprotectants include trehalose and lactose. Bulking agents can be included for a lyophilized dosage form, principally 1-10% mannitol (optimally 2-4%). Stabilizers can be used in both liquid and lyophilized dosage forms, principally 1- 20 50 mM L-Methionine (optimally 5-10 mM). Other suitable bulking agents include glycine, arginine, can be included as 0-0.05% polysorbate-80 (optimally 0.005-0.01%). Additional surfactants include but are not limited to polysorbate 20 and BRIJ surfactants.

The compositions of this invention can be in a variety of forms. These 25 include, for example, liquid, semi-solid and solid dosage forms, such as liquid solutions (*e.g.*, injectable and infusible solutions), dispersions or suspensions, tablets, pills, powders, liposomes and suppositories. The preferred form depends on the intended mode of administration and therapeutic application. Typical preferred compositions are in the form of injectable or infusible solutions, such as compositions similar to those used for passive immunization of humans. The 30 preferred mode of administration is parenteral (*e.g.*, intravenous, subcutaneous, intraperitoneal, intramuscular). In a preferred embodiment, the antiangiogenic is administered by intravenous infusion or injection. In another preferred embodiment, the antiangiogenic is administered by intramuscular or

subcutaneous injection. In the most preferred embodiment, the antiangiogenic is administered orally.

Alternatively, a preferred embodiment includes the use of the compounds of the invention in eye-drop solutions. This provides for application to the ease 5 to inhibit or prevent ocular diseases such as glaucoma. Generally, the active ingredient, i.e., the compounds of the invention, would be dissolved in an aqueous solution that can be applied directly to the eye.

Therapeutic compositions typically must be sterile and stable under the 10 conditions of manufacture and storage. The composition can be formulated as a solution, microemulsion, dispersion, liposome, or other ordered structure suitable to high drug concentration. Sterile injectable solutions can be prepared by incorporating the active compound (*i.e.*, antigen, antibody or antibody portion) in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization.

15 Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile, lyophilized powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and spray-drying that yields a powder of the 20 active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof. The proper fluidity of a solution can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prolonged absorption of injectable compositions can be brought about by 25 including in the composition an agent that delays absorption, for example, monostearate salts and gelatin.

The antiangiogenics of the present invention can be administered by a variety of methods known in the art. As will be appreciated by the skilled artisan, the route and/or mode of administration will vary depending upon the desired 30 results. In certain embodiments, the active compound can be prepared with a carrier that will protect the compound against rapid release, such as a controlled release formulation, including implants, transdermal patches, and

microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Many methods for the preparation of such formulations are patented or generally known to those skilled in the art.

5 *See, e.g., Sustained and Controlled Release Drug Delivery Systems*, J.R. Robinson, ed., Marcel Dekker, Inc., New York, 1978.

In certain embodiments, an antiangiogenic of the invention can be orally administered, for example, with an inert diluent or an assimilable edible carrier. The compound (and other ingredients, if desired) can also be enclosed in a hard 10 or soft shell gelatin capsule, compressed into tablets, or incorporated directly into the subject's diet. For oral therapeutic administration, the compounds can be incorporated with excipients and used in the form of ingestible tablets, buccal tablets, troches, capsules, elixirs, suspensions, syrups, wafers, and the like. To administer a compound of the invention by other than parenteral administration, 15 it can be necessary to coat the compound with, or co-administer the compound with, a material to prevent its inactivation.

The present invention also provides for packaged pharmaceutical compositions useful in the prevention or inhibition of angiogenic activity in a subject. The packaged pharmaceutical composition includes a container holding 20 a therapeutically effective amount of at least one ATL, such as ATL-1 or an LXA₄, such as 15-R/S-methyl, LXA₄, or a pharmaceutically acceptable salt, esters, amide, or prodrug thereof and instructions for using the therapeutic compound for preventing, reducing or inhibiting angiogenic activity in the subject. Additionally, the present invention provides therapeutically effective 25 amounts of packaged pharmaceutical compositions, e.g., ATL-1, LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ or pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, and instructions useful to treat, i.e., inhibit or prevent, solid tumor tissue growth from undergoing neovascularization, neovascularization from occurring, neovascularization from occurring in retinal 30 tissue, restenosis from occurring following angioplasty in a tissue wherein smooth muscle cell migration occurs, or reducing blood supply to a tissue required to support new growth of new angiogenic tissue.

The present invention also provides angiogenic compounds that facilitate angiogenesis. Surprisingly, configurational isomers of LXA₄, LXA₄ analogs and ATL analogs, LXB₄ and LXB₄ analogs and pharmaceutically acceptable salts, esters, amides or prodrugs thereof, provide the opposite effects with regard to 5 revascularization of tissue by the above-identified compounds of the invention. That is, it has been surprisingly discovered that LXB₄ and LXB₄ analogs have the ability to stimulate regeneration and ingrowth of vascular or epithelial tissue in tissues that are in need of such stimulation. This is especially important in tissue 10 grafting, tissue engineering and prosthetic group sites of attachment. Therefore, the present invention provides methods of tissue regeneration, compounds for 15 such application and packaged pharmaceuticals to accomplish such results.

For example, cardiovascular disease occurs as a consequence of the 20 partial or complete blockage of vessels carrying blood in the coronary vascular system and in peripheral vasculature. Occlusion of the vessel can result in death 15 of tissue previously nourished by the occluded vessels or inability of the vessels to transport sufficient blood supply to regions requiring high blood consumption and accompanying nutrients. Blood vessel occlusion can be partially compensated by the natural process of angiogenesis, in which new conduits are 25 formed to replace the function of the impaired vessels. These new conduits are referred to as "collateral" vessels and can help in the restoration of blood flow to the deprived tissue, thereby constituting natural bypasses around the occluded vessels. However, some individuals for various reasons are unable to generate 30 sufficient collateral vessels to manage the consequences of diminished blood flow from cardiovascular disease.

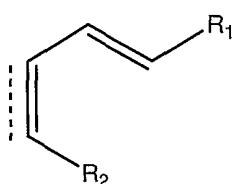
25 The LXB₄ compounds of the invention can be used to enhance the body's natural ability to repair itself by undergoing natural angiogenesis. As can be seen from the contents of the specification and Figures, vessel growth is stimulated by these unique compounds. This process and use of the present compounds can be utilized for the treatment of wounds. The LXB₄ compounds help stimulate the 30 healing process, causing re-epithelialization and vascularization to occur.

**Suitable lipoxin analogs, including ATLs, LXA₄s and LXB₄s
encompassed by the present invention include those having the following
characteristics.**

The instant lipoxins comprising an "active region" and a "metabolic transformation region" as both terms are defined herein are generally of the following structure:

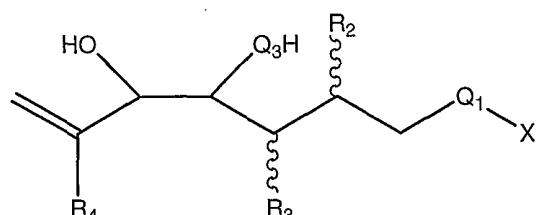
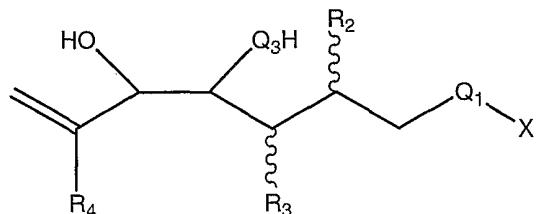
wherein R₁ can be

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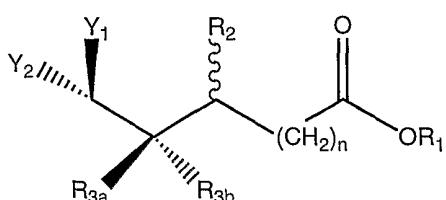


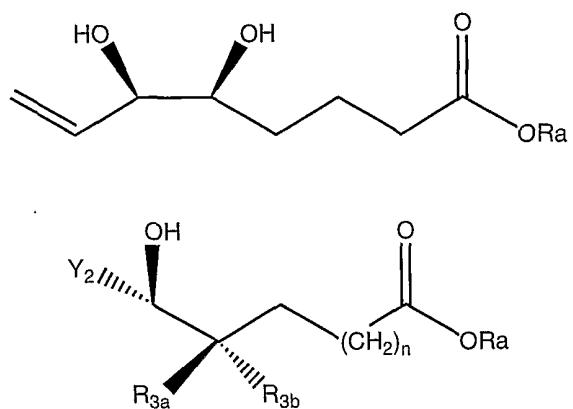
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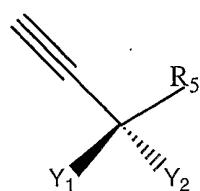
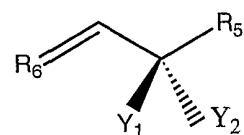


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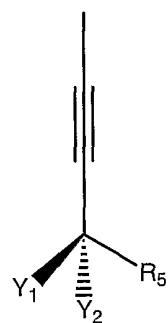


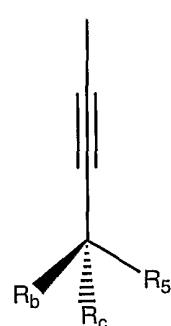
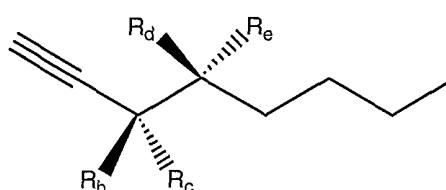
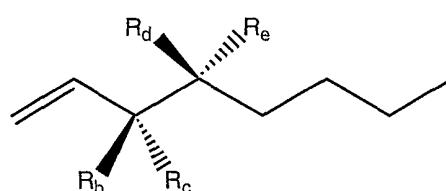
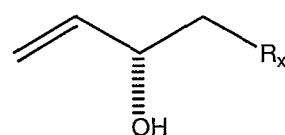
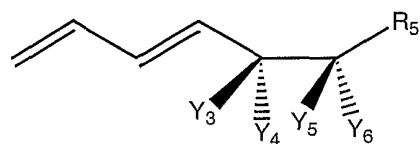
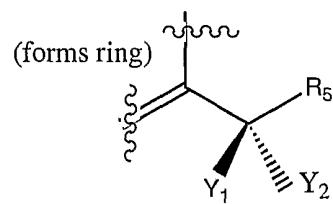


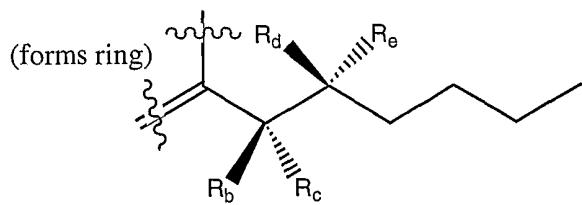
5 and R₂ can be



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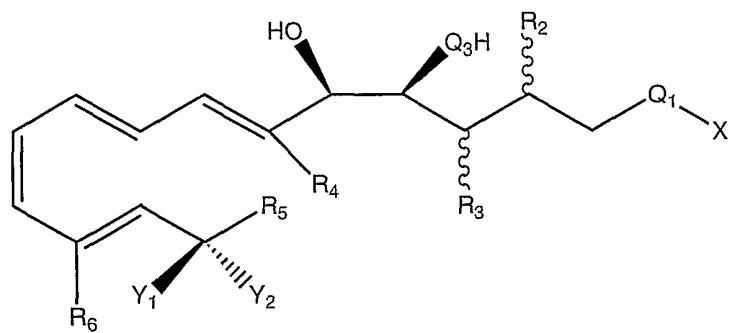






In one embodiment, the lipoxin analogs of this invention have the following structural formula I:

5



wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

(i) a hydrogen atom;

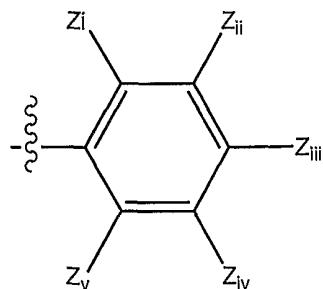
10 (ii) an alkyl of 1 to 8 carbon atoms, inclusive, which can be straight chain or branched;

(iii) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(iv) an aralkyl of 7 to 12 carbon atoms;

(v) phenyl;

15 (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 can 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;

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) ;

10 wherein Q_3 is O, S or NH;

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

(a) a hydrogen atom;

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

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

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

(e) $\text{R}_a \text{Q}_2 \text{R}_b$

wherein Q_2 is $-\text{O}-$ or $-\text{S}-$;

20 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be
straight chain or branched; and wherein R_b is alkyl of 0 to 8 carbon atoms,
inclusive, which can be straight chain or branched;

wherein R_4 is

(a) a hydrogen atom;

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

wherein Y_1 or Y_2 is $-\text{OH}$, methyl, or $-\text{SH}$ and wherein the other is

(a) a hydrogen atom

(b) CH_aZ_b

30 where $a+b=3$, $a=0$ to 3, $b=0$ to 3; and
each Z , independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

or

(d) an alkoxy of 1 to 4 carbon atoms, inclusive;

or Y_1 and Y_2 taken together are

5 (a) =NH; or

(b) =O;

wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or

branched;

10 (b) $-(CH_2)_n-R_i$

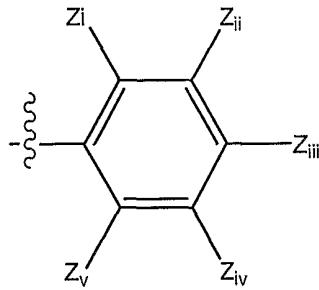
wherein $n=0$ to 4 and R_i is

(i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) a phenyl; or

(iii) substituted phenyl

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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, -

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

(c) $R_aQ_aR_b$

wherein Q_a is O or S;

25 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

(d) $—C(R_{iii})(R_{iv})—R_i$

wherein R_{iii} and R_{iv} are each, independently:

5 (i) a hydrogen atom;

(ii) CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0+3$, and wherein each Z , independently, is a cyano, a nitro, or a halogen atom;

(e) a haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms, inclusive, straight chain or branched; and

10 wherein R_6 is

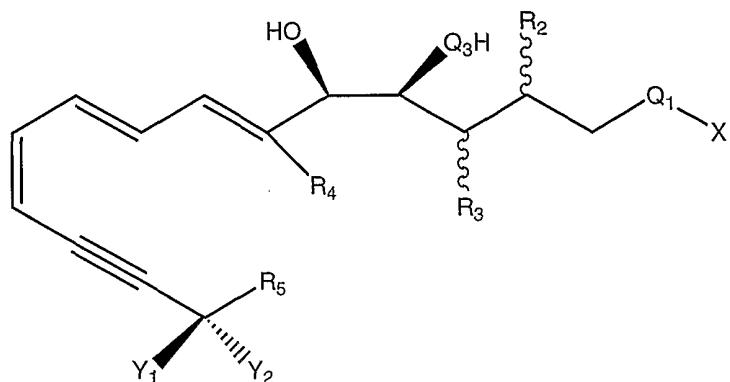
(a) a hydrogen atom;

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

(c) a halogen.

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In one embodiment of this invention, the lipoxin analogs have the following structure II:

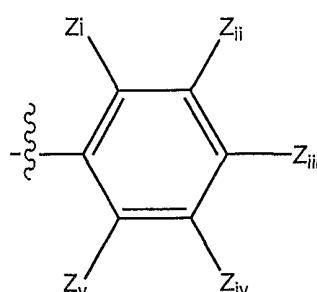


5

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

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

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wherein Z_i, Z_{ii}, Z_{iii}, Z_{iv} and Z_v 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 can be a straight chain or branched, and hydroxyl;

(vii) a detectable label molecule, such as but not limited to fluorescent labels; or

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

wherein Q_1 is (C=O), SO₂ or (C=N);

wherein Q_3 is O, S or NH;

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

10 (a) a hydrogen atom;

(b) an alkyl of 1 to 8 carbon atoms, inclusive, which can be 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 can be straight

15 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 carbons atoms, inclusive, which can be straight chain or branched;

20 wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

wherein R_4 is

(a) a hydrogen atom;

(b) alkyl of 1 to 6 carbon atoms, inclusive, which can be straight chain or

25 branched;

wherein Y_1 or Y_2 is —OH, methyl, -H or -SH and wherein the other is

(a) a hydrogen atom;

(b) CH_aZ_b

where $a+b=3$, $a=0$ to 3, $b=0$ to 3

30 wherein each Z , independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

(d) an alkoxy of 1 to 4 carbon atoms, inclusive; or Y_1 and Y_2 taken together are

(a) $=NH$; or

(b) $=O$;

5 wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or branched;

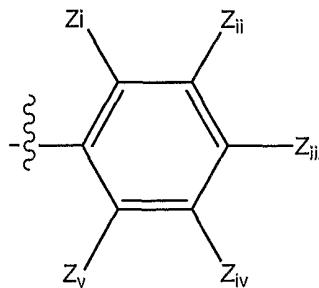
(b) $-(CH_2)_n-R_i$

wherein $n=0$ to 4 and R_i is

10 (i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) phenyl; or

(iii) 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 can be a straight chain or branched, and hydroxyl;

(c) $-R_a Q_a R_b$

20 wherein Q_a is $-O-$ or $-S-$; and

wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

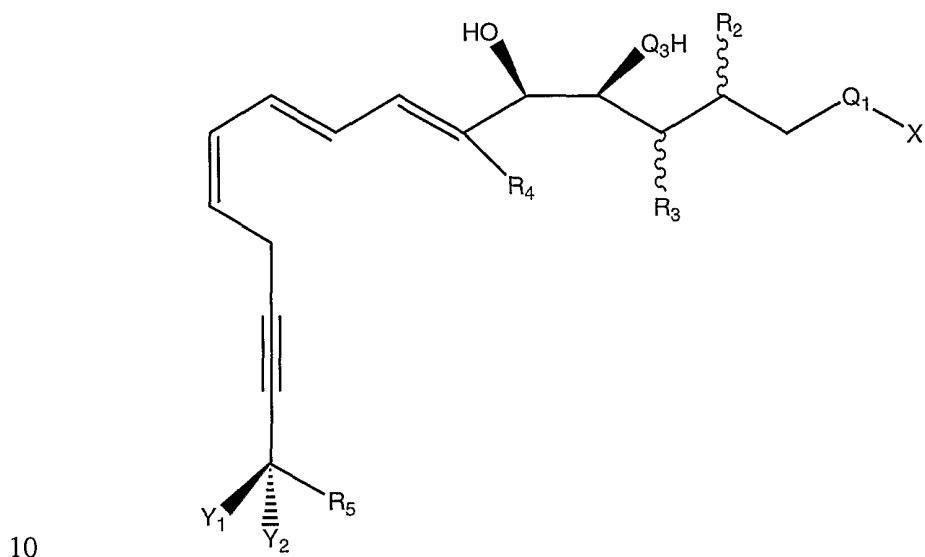
wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

25 (d) $-C(R_{iii})(R_{iv})-R_i$

wherein R_{iii} and R_{iv} are each independently:

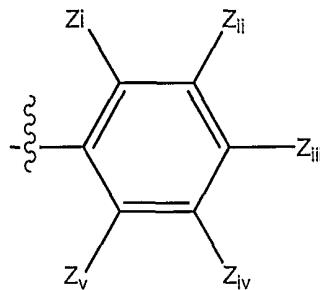
- (i) a hydrogen atom; or
- (ii) CH_aZ_b where $a+b=3$, $a=0$ to 3 , $b=0+3$
wherein each Z , independently, is a cyano, a nitro, or a halogen atom,
- (e) a haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms, inclusive, straight chain or branched.

In one embodiment of this invention, the lipoxin analogs have the following structure III:



- 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 carbons atoms, inclusive, which can be straight chain or branched;
 - (iii) a cycloalkyl of 3 to 10 carbon atoms, inclusive;
 - (iv) an aralkyl of 7 to 12 carbon atoms;
 - (v) phenyl;
 - (vi) substituted phenyl

15
20



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, $-\text{OR}_x$,

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

(vii) a detectable label molecule; or

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

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or $(\text{C}=\text{N})$;

wherein Q_3 is O, S or NH;

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

(a) a hydrogen atom;

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

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

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

(e) $\text{R}_a\text{Q}_2\text{R}_b$

20 wherein Q_2 is $-\text{O}-$ or $-\text{S}-$;

wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be
straight chain or branched;

wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be
straight chain or branched;

25 wherein R_4 is

(a) a hydrogen atom; or

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

wherein Y_1 or Y_2 is hydroxyl, methyl, hydrogen or thiol and wherein the other is

5 (a) a hydrogen atom;

(b) CH_aZ_b

where $a+b=3$, $a=0$ to 3, $b=0$ to 3

wherein each Z , independently, is a cyano, a nitro, or a halogen atom ;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

10 (d) an alkoxy of 1 to 4 carbon atoms, inclusive; or Y_1 and Y_2 taken together are

(a) =NH; or

(b) =O; and

wherein R_5 is

15 (a) an alkyl of 1 to 9 carbon atoms which can be straight chain or branched;

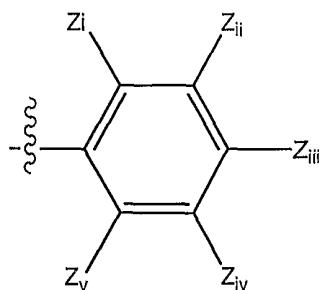
(b) $-(CH_2)_n-R_i$

wherein $n=0$ to 4 and R_i is

(i) cycloalkyl of 3 to 10 carbon atoms, inclusive;

20 (ii) phenyl;

(iii) substituted phenyl



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

25 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 can be a straight chain or branched, and hydroxyl;

(c) $R_a Q_a R_b$

wherein Q_a is $-O-$ or $-S-$;

5 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched; or

(d) $—C(R_{iii})(R_{iv})—R_i$

10 wherein R_{iii} and R_{iv} are each independently:

(i) a hydrogen atom; or

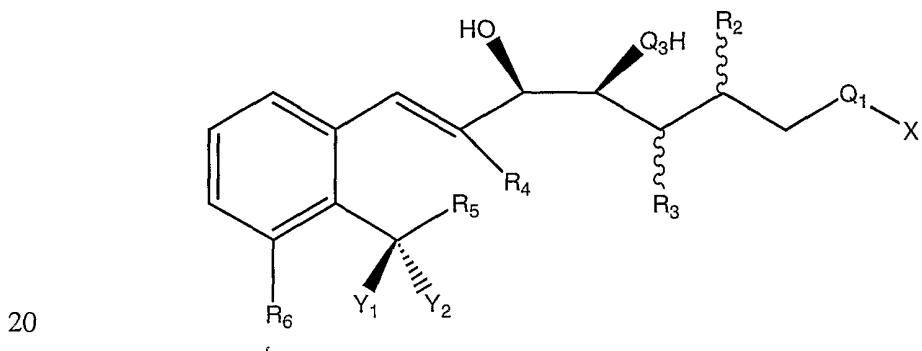
(ii) $CH_a Z_b$ where $a+b=3$, $a=0$ to 3, $b=0+3$

wherein each Z , independently, is a cyano, a nitro, or a halogen atom,

(e) a haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen

15 atoms, inclusive, straight chain or branched.

In another embodiment of this invention, lipoxin analogs have the following structural formula IV:



wherein X is R_1 , OR_1 , or SR_1 ; wherein R_1 is

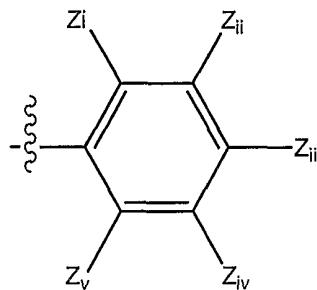
(i) a hydrogen atom;

(ii) an alkyl of 1 to 8 carbons atoms, inclusive, which can be straight

25 chain or branched;

- (iii) a cycloalkyl of 3 to 10 carbon atoms, inclusive;
- (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, $-\text{OR}_x$,
wherein R_x is 1 to 8 carbon atoms, inclusive, which can be a straight chain or
10 branched, and hydroxyl;

(vii) a detectable label molecule; or

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

wherein Q_1 is $(\text{C}=\text{O})$, SO_2 or (CN) ;

15 wherein Q_3 is O, S or NH;

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

(a) a hydrogen atom;

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

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

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

(e) $\text{R}_a\text{Q}_2\text{R}_b$

wherein Q_2 is $-\text{O}-$ or $\text{S}-$;

25 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be
straight chain or branched;

wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

wherein R_4 is

(a) a hydrogen atom; or

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

wherein Y_1 or Y_2 is $-\text{OH}$, methyl, or $-\text{SH}$ and wherein the other is

(a) a hydrogen atom;

(b) CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0$ to 3, wherein each Z ,

10 independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

or

(d) an alkoxy of 1 to 4 carbon atoms, inclusive;

or Y_1 and Y_2 taken together are

15 (a) $=\text{NH}$; or

(b) $=\text{O}$;

wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or branched;

20 (b) $-(\text{CH}_2)_n-\text{R}_i$

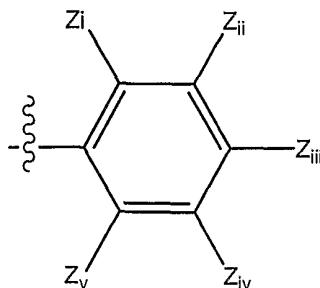
wherein $n=0$ to 4 and R_i is

(i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) phenyl; or

(iii) substituted phenyl

25

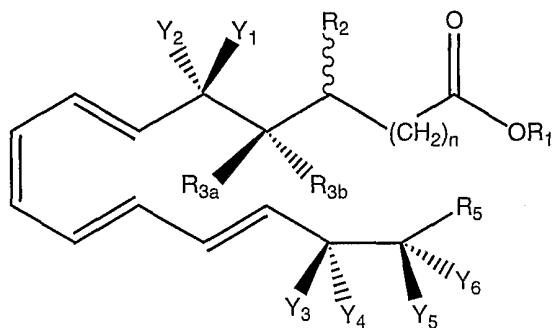


wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from – NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which can be a straight chain or branched,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

5 (c) $\text{R}_a\text{Q}_a\text{R}_b$
wherein Q_a is $-\text{O}-$ or $-\text{S}-$;
wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;
wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

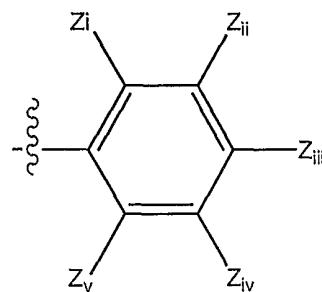
10 (d) $-\text{C}(\text{R}_{iii})(\text{R}_{iv})-\text{R}_i$
wherein R_{iii} and R_{iv} are each independently:
(i) a hydrogen atom; or
(ii) CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0+3$ and
15 wherein each Z , independently, is a cyano, a nitro, or a halogen atom; or
(e) haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms, inclusive, straight chain or branched; and
wherein R_6 is
(a) a hydrogen atom;
20 (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched; or
(c) a halogen atom.

In another embodiment of this invention, lipoxin analogs have the
25 following structural formula V:



wherein R_1 is

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



wherein Z_1 , 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, $-OR_x$,

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

(vii) a detectable label molecule; or

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

20 wherein $n = 1$ to 10, inclusive;

wherein R_2 , R_{3a} , and R_{3b} are each independently:

(a) a hydrogen atom;

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

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

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

(e) $R_aQ_2R_b$

wherein Q_2 is $-O-$ or $-S-$;

10 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched; and wherein R_b is alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched;

wherein Y_1 or Y_2 is $-OH$, methyl, hydrogen, or $-SH$ and

wherein the other is

15 (a) a hydrogen atom;

(b) CH_aZ_b

where $a+b=3$, $a=0$ to 3, $b=0$ to 3, and

wherein each Z , independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

20 (d) an alkoxy of 1 to 4 carbon atoms, inclusive, straight chain or branched;

or Y_1 and Y_2 taken together are

(a) $=NH$; or

(b) $=O$;

25 wherein Y_3 or Y_4 is $-OH$, methyl, hydrogen, or $-SH$ and

wherein the other is

(a) a hydrogen atom;

(b) CH_aZ_b

wherein $a+b=3$, $a=0$ to 3, $b=0$ to 3,

30 and wherein each Z , independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

(d) an alkoxy of 1 to 4 carbon atoms, inclusive, straight chain or branched;

or Y_3 and Y_4 taken together are

(a) $=\text{NH}$; or

5 (b) $=\text{O}$;

wherein Y_5 or Y_6 is $-\text{OH}$, methyl, hydrogen, or $-\text{SH}$ and wherein the other is

(a) a hydrogen atom;

(b) CH_aZ_b

10 where $a+b=3$, $a=0$ to 3, $b=0$ to 3
wherein each Z , independently, is a cyano, a nitro, or a halogen atom;

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

(d) an alkoxy of 1 to 4 carbon atoms, inclusive, straight chain or branched;

15 or Y_5 and Y_6 taken together are

(a) $=\text{NH}$; or

(b) $=\text{O}$;

wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or

20 branched;

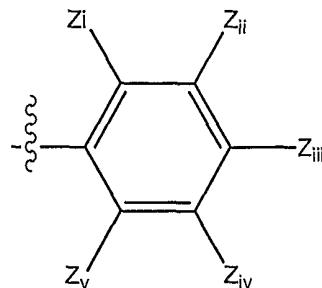
(b) $-(\text{CH}_2)_n-\text{R}_i$

wherein $n=0$ to 4 and R_i is

(i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) phenyl; or

25 (iii) substituted phenyl



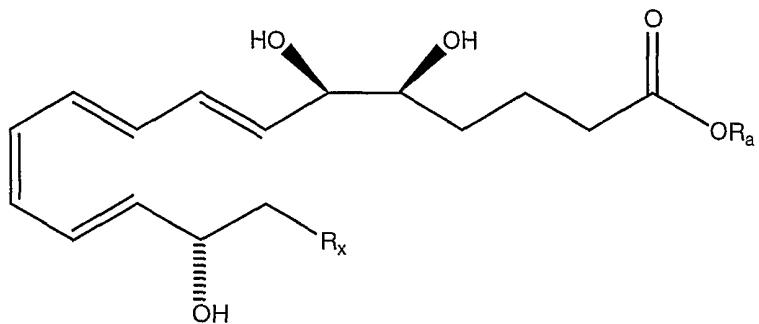
wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –
 NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$,
wherein R_x is 1 to 8 carbon atoms, inclusive, which can be a straight chain or
5 branched,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

(c) $-\text{R}_a\text{Q}_a\text{R}_b$
wherein Q_a is $-\text{O}-$ or $-\text{S}-$; and
wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

10 wherein R_b is either alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched or substituted phenyl;

(d) $-\text{C}(\text{R}_{iii})(\text{R}_{iv})-\text{R}_i$
wherein R_{iii} and R_{iv} are each independently:
(i) a hydrogen atom; or
15 (ii) CH_aZ_b where $a+b=3$, $a=0$ to 3, $b=0+3$, and
wherein each Z , independently, is a cyano, a nitro, or a halogen atom; or
(e) haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms, inclusive, straight chain or branched.

20 In another embodiment of this invention, lipoxin analogs have the structural formula VI:



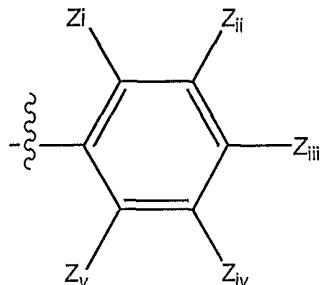
wherein R_a is

(a) a hydrogen atom; or

5 (b) alkyl of 1 to 8 carbon atoms;

wherein R_x is

(a) substituted phenyl

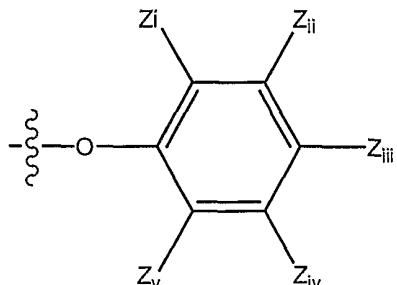


10 wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from –

NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$,

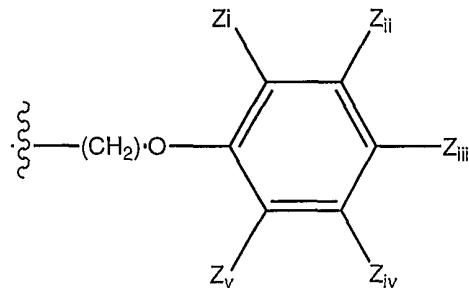
wherein R_x is 1 to 8 carbon atoms, inclusive, which can be a straight chain or branched, and hydroxyl;

(b) a substituted phenoxy



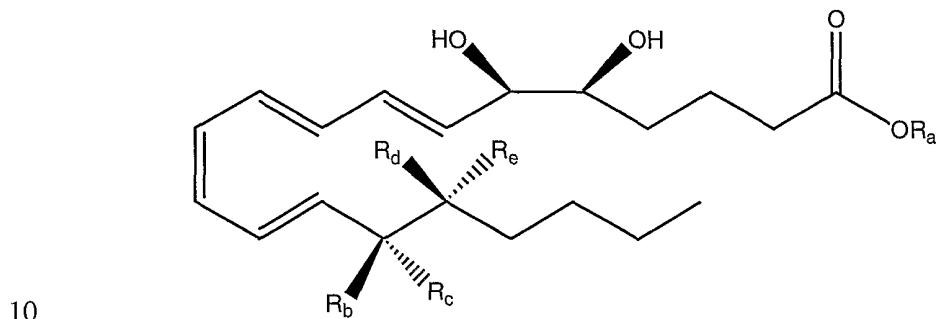
15

wherein Z_i through Z_v are as defined above; or



5 wherein Z_i through Z_v are as defined above.

In another preferred embodiment of this invention, lipoxin analogs have the following structural formula VII:



10

wherein R_a is

- (a) a hydrogen atom; or
- (b) an alkyl of 1 to 8 carbon atoms;

15 wherein R_b and R_c are each independently:

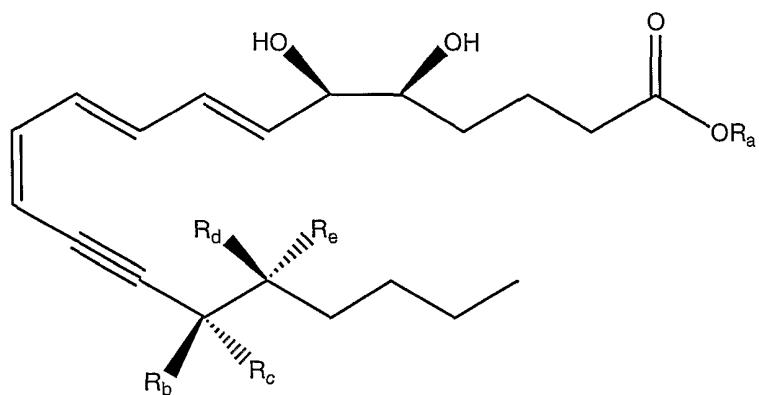
- (a) a hydrogen atom;
- (b) a hydroxyl, or a thiol;
- (c) a methyl or a halomethyl;
- (d) a halogen;
- (e) an alkoxy of 1 to 3 carbon atoms;

20

wherein R_d and R_e are each independently:

- (a) a hydrogen atom;
- (b) a hydroxyl, or thiol;
- (c) a methyl or halomethyl;
- 5 (d) a halogen;
- (e) an alkoxy of 1 to 3 carbon atoms; or
- (f) an alkyls or haloalkyl of 2 to 4 carbon atoms, inclusive, which can be straight chain or branched.

10 In another preferred embodiment of this invention, the lipoxin analogs have the structural formula VIII:



15 wherein R_a is

- (a) a hydrogen atom; or
- (b) an alkyl of 1 to 8 carbon atoms;

wherein R_b and R_c are each independently:

- (a) a hydrogen atom;
- 20 (b) a hydroxyl or a thiol;
- (c) a halomethyl;
- (d) a halogen;
- (e) an alkyl of 1 to 3 carbon atoms, inclusive, straight chain or branched;

or

(f) an alkoxy of 1 to 3 carbon atoms, inclusive;

wherein R_d and R_e are each independently:

(a) a hydrogen atom;

(b) a hydroxyl, or a thiol;

5 (c) a methyl or a halomethyl;

(d) a halogen;

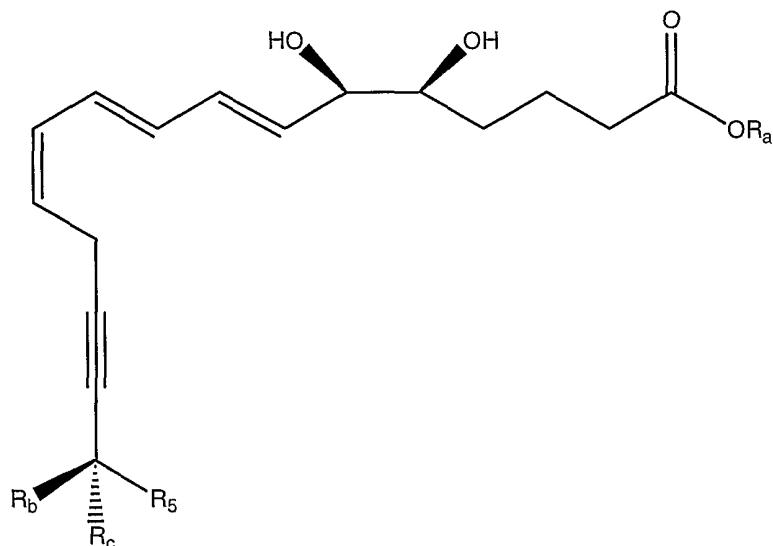
(e) an alkoxy of 1 to 3 carbon atoms, inclusive; or

(f) an alkyl or haloalkyl of 2 to 4 carbon atoms, inclusive, which can be

straight chain or branched.

10

In another embodiment of this invention, the lipoxin analogs have the structural formula IX:



15

wherein R_a is

(a) a hydrogen atom; or

(b) an alkyl of 1 to 8 carbon atoms;

wherein R_b and R_c are each independently:

20 (a) a hydrogen atom;

(b) a hydroxyl or thiol;

(c) a halomethyl;

(d) a halogen;

(e) an alkyl of 1 to 3 carbon atoms, inclusive, straight chain or branched;

(f) an alkoxy of 1 to 3 carbon atoms, inclusive; and

5 wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or branched;

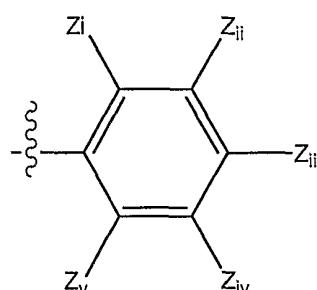
(b) $-(CH_2)_n-$, $-R_i$

wherein $n = 0$ to 4 and R_i is

10 (i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) phenyl; or

(iii) 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 can be a straight chain or branched, and hydroxyl;

(c) $R_aQ_aR_b$

20 wherein Q_a is $-O-$ or $-S-$;

wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

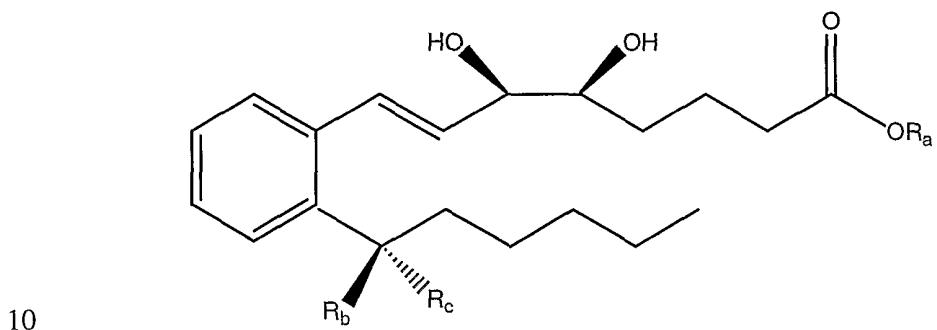
wherein R_b is either alkyl of 0 to 8 carbon atoms, inclusive, which can be straight chain or branched or substituted phenyl;

25 (d) $-C(R_{iii})(R_{iv})-R_i$

wherein R_{iii} and R_{iv} are each, independently:

- (i) a hydrogen atom; or
- (ii) CH_aZ_b where $a+b=3$, $a=0$ to 3 , $b=0+3$
wherein each Z , independently, is a cyano, a nitro, or a halogen atom; or
- (e) haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms,
5 inclusive, straight chain or branched.

In another preferred embodiment, the compounds have the structural formula X:



wherein R_a is

- (a) a hydrogen atom; or
- (b) alkyl of 1 to 8 carbon atoms, inclusive, straight chain or branched; and

15 wherein R_b and R_c are each, independently:

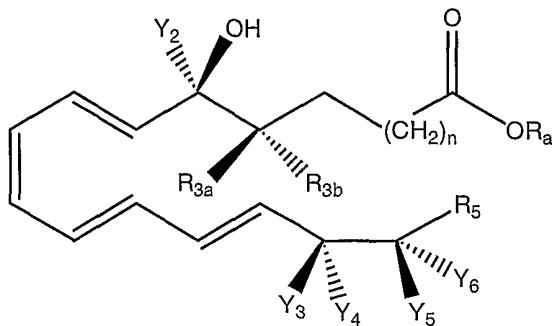
- (a) a hydrogen atom;
- (b) a hydroxyl or a thiol;
- (c) a halomethyl;
- (d) a halogen;

20 (e) an alkyl of 1 to 3 carbon atoms, inclusive, straight chain or branched;

- (f) an alkoxy of 1 to 3 carbon atoms, inclusive.

In another preferred embodiment, the compounds have the structural formula XI:

25



wherein R_a is

(i) a hydrogen atom;

5 (ii) an alkyl of 1 to 8 carbon atoms, inclusive, which can be straight chain or branched; or

(iii) a detectable label molecule;

wherein $n=1$ to 10, inclusive;

wherein Y_2 , R_{3a} , and R_{3b} are each, independently:

10 (a) a hydrogen atom;

(b) an alkyl of 1 to 8 carbon atoms, inclusive, which can be 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 can be straight

15 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, inclusive, which can be

straight chain or branched; and wherein R_b is alkyl of 0 to 8 carbon atoms,

20 inclusive, which can be straight chain or branched;

wherein Y_1 is $—OH$, methyl, or $—SH$;

wherein Y_2 is

(a) a hydrogen atom;

(b) CH_aZ_b

25 where $a+b=3$, $a=0$ to 3, $b=0$ to 3

wherein each Z , independently, is a cyano, a nitro, or a halogen atom; or

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched; wherein Y_3 and Y_5 are each independently:

(a) a hydrogen atom;

5 (b) CH_aZ_b

wherein $a+b=3$, $a=0$ to 3, $b=0$ to 3 and wherein each Z , independently, is a cyano, a nitro, or a halogen atom; or

(c) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched; wherein Y_4 and Y_6 are each, independently

10 (a) a hydrogen atom;

(b) an alkyl of 2 to 4 carbon atoms, inclusive, straight chain or branched;

(c) an alkoxy of 1 to 4 carbon atoms, inclusive, straight chain or branched;

or

(d) a hydroxyl or thiol; and

15 wherein R_5 is

(a) an alkyl of 1 to 9 carbon atoms which can be straight chain or branched;

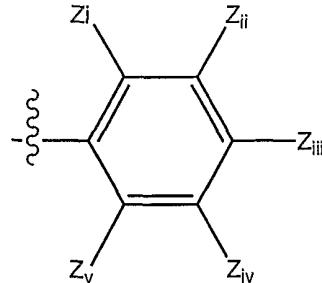
(b) $-(\text{CH}_2)_n-\text{R}_i$

wherein $n=0$ to 3 and R_i is

20 (i) a cycloalkyl of 3 to 10 carbon atoms, inclusive;

(ii) phenyl;

(iii) substituted phenyl



25 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 can be a straight chain or branched, and hydroxyl;

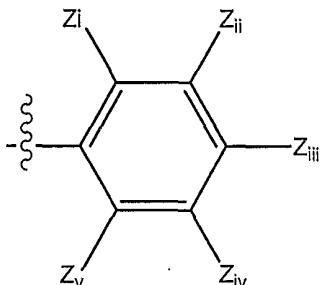
(c) $-R_a Q_a R_b$

wherein Q_a is $-O-$ or $-S-$;

5 wherein R_a is alkylene of 0 to 6 carbons atoms, inclusive, which can be straight chain or branched;

wherein R_b is

(a) a substituted phenyl

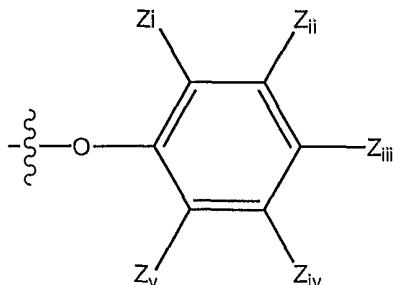


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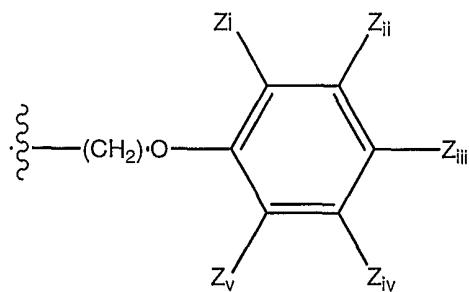
wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-NO_2$, $-CN$, $-C(=O)-R_1$, $-SO_3H$, a hydrogen atom, halogen, methyl, $-OR_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which can be a straight chain or branched, and hydroxyl;

15

(b) a substituted phenoxy



wherein Z_i through Z_v are as defined above; or

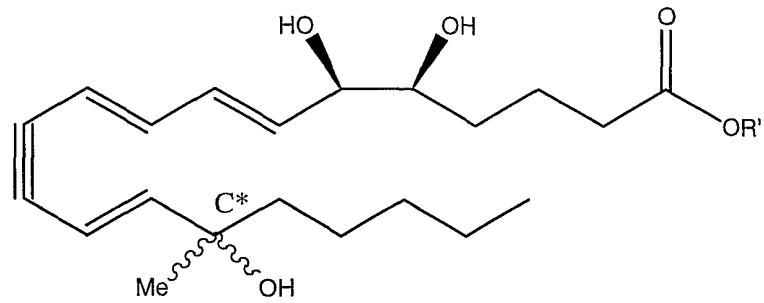


wherein Z_i through Z_v are as defined above;

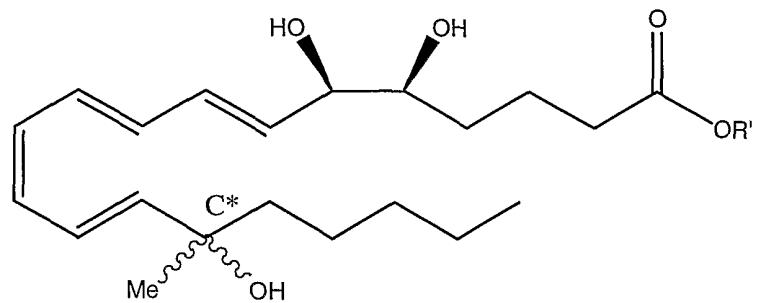
(d) a haloalkyl of 1 to 8 carbon atoms, inclusive, and 1 to 6 halogen atoms, inclusive, straight chain or branched.

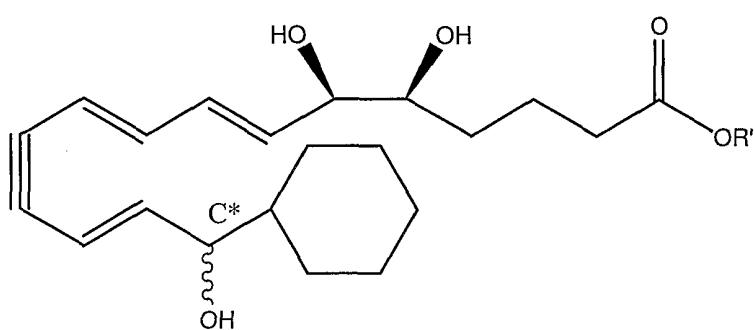
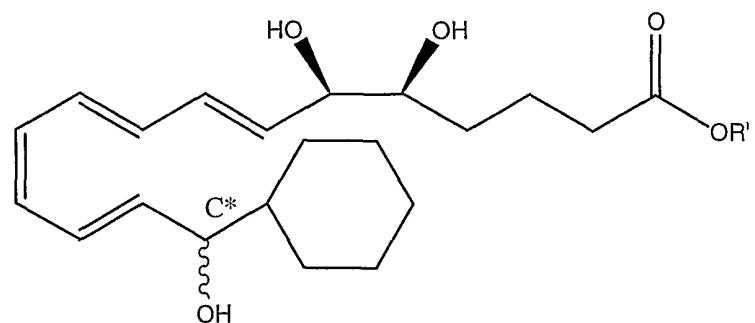
In certain embodiments of this invention, the compounds of this invention have the following structural formulas:

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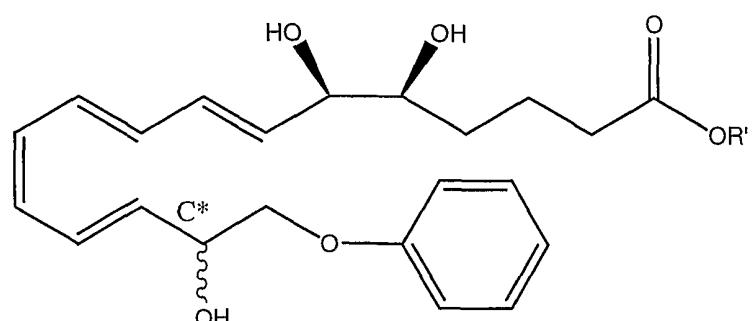
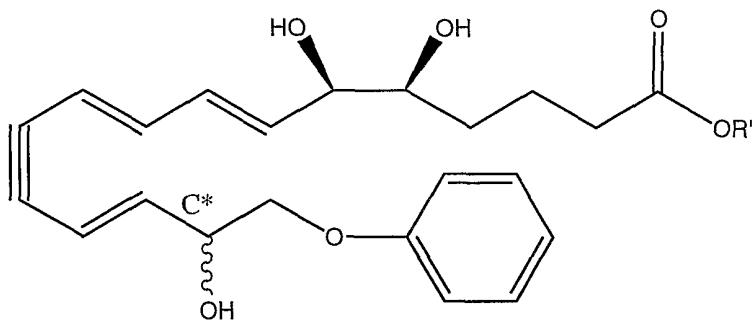


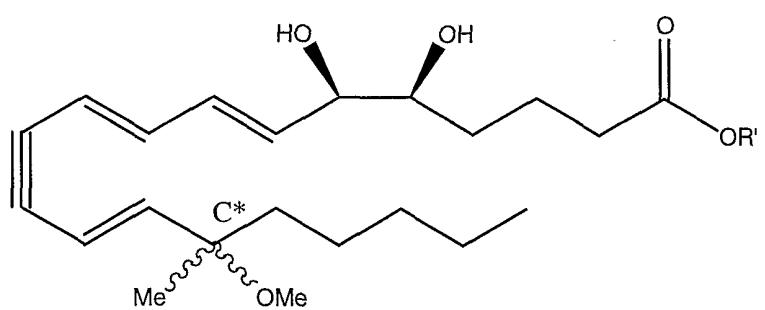
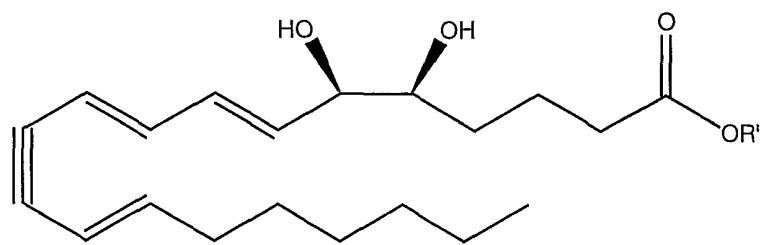
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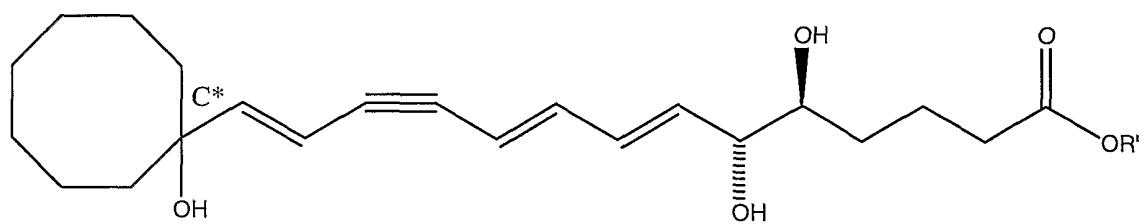


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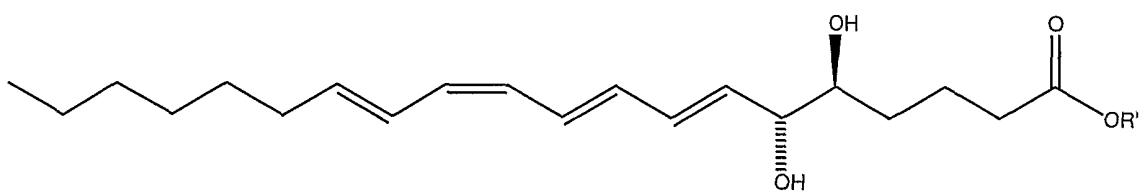
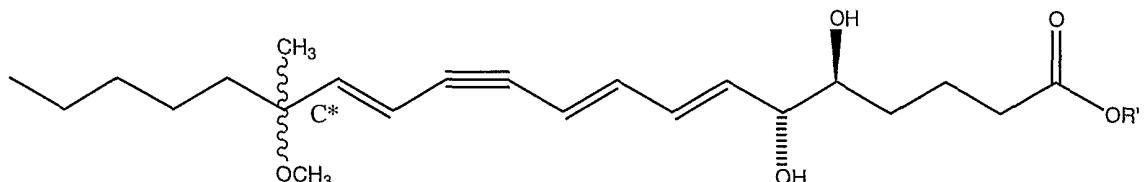




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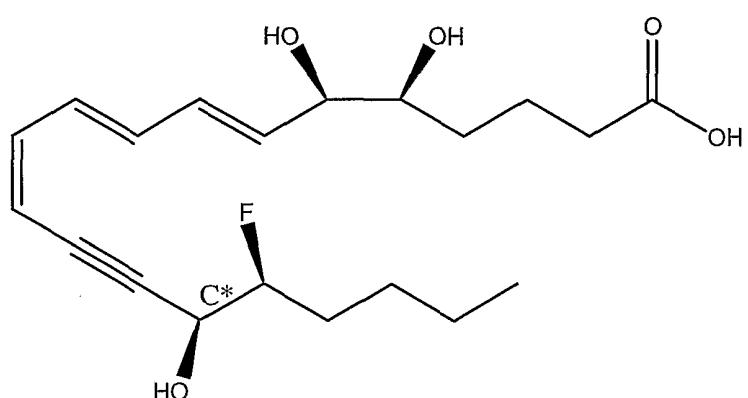
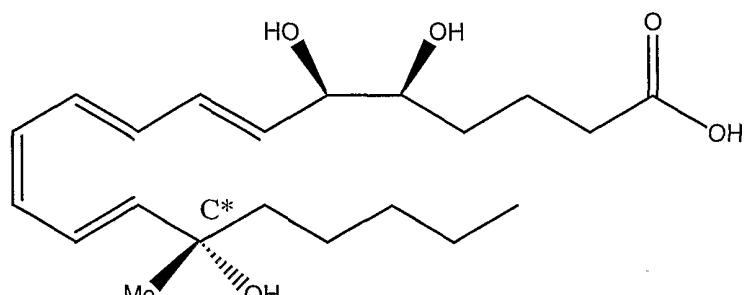


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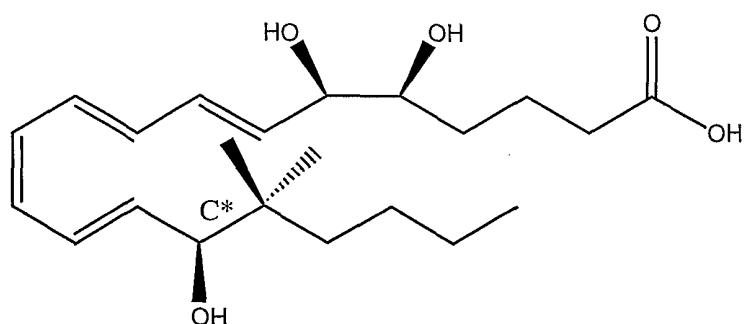


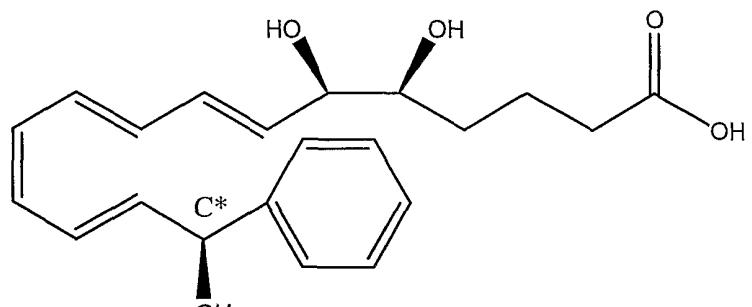
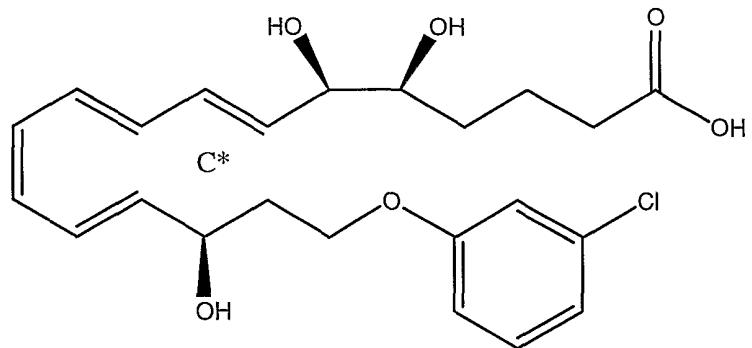
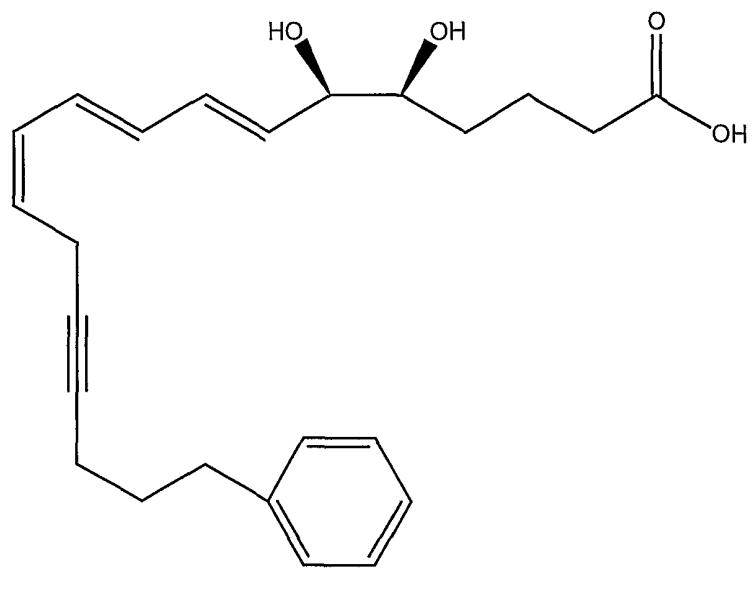
where R' is H or CH₃; and where the substituents at C* are in the R configuration.

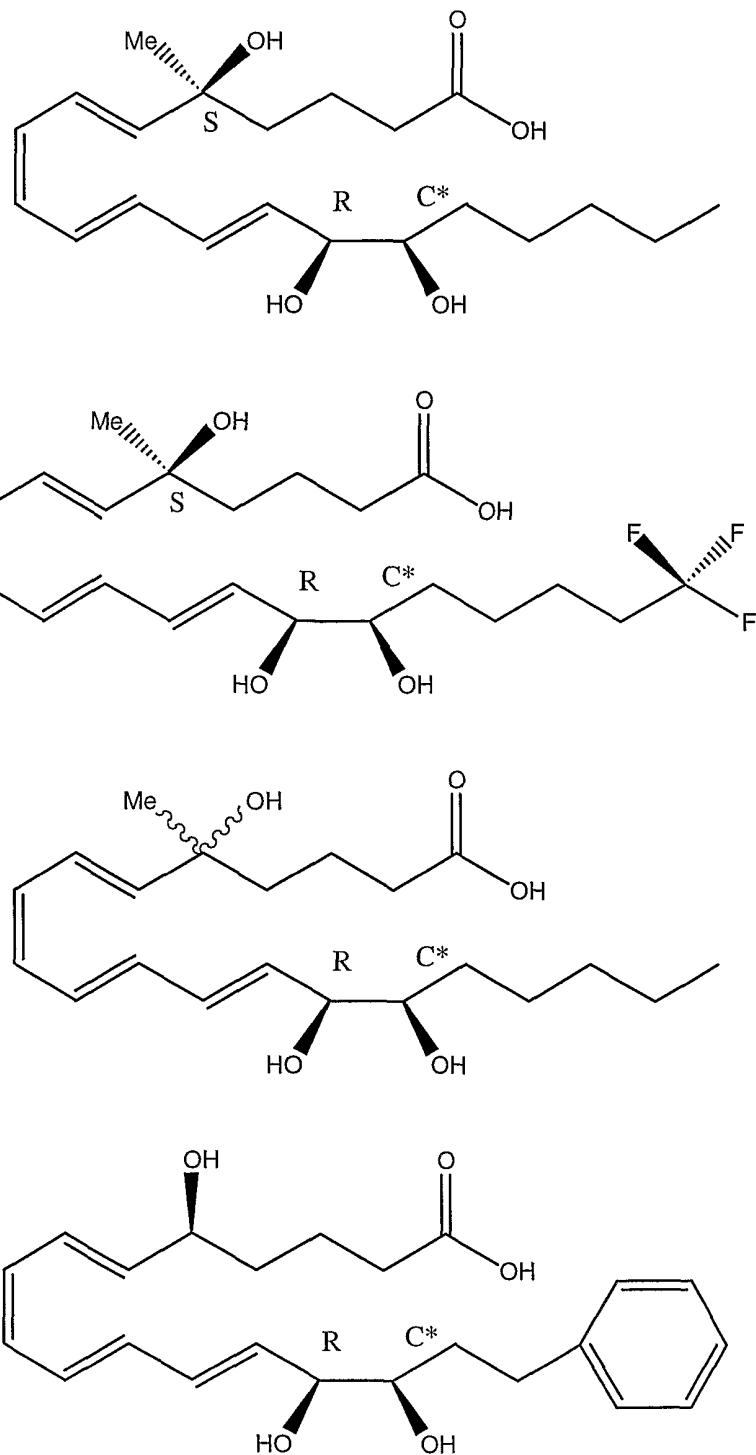
In other preferred embodiments of this invention, the compounds of this
5 invention have the following structural formulas:



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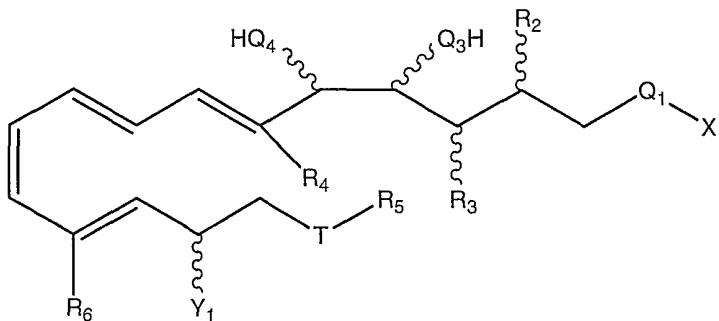
where the substituents at the C^* are in the R configuration.

It is to be understood that the carboxylic acids and esters of the invention can be converted, if necessary, into pharmaceutically acceptable salts.

5 In certain embodiments of the invention, LXB₄ or the C5 and C14 and C15 alkanoates (acetates) of LXB₄ may be excluded.

LIPOXINS HAVING PHENOXY OR THIOPHENOXY SUBSTITUENTS

In another aspect, lipoxins and lipoxin analogs useful as therapeutic agents in the treatment of the maladies, disease states or conditions described 10 throughout the specification has the formula:

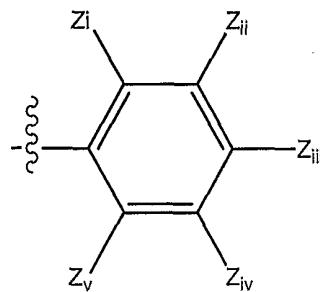


15 wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
20 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

25



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;

- (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 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

15

- (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) $\text{R}_a\text{Q}_2\text{R}_b$ 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

20

25

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

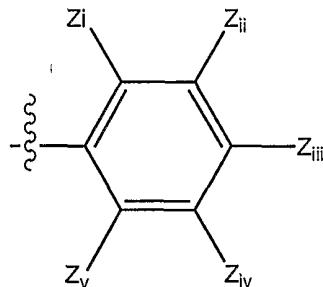
wherein R_4 is

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 , –CN, –C(=O)– R_1 , – SO_3H , a hydrogen atom, halogen, methyl, –
15 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,
20 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

25

- (a) H;

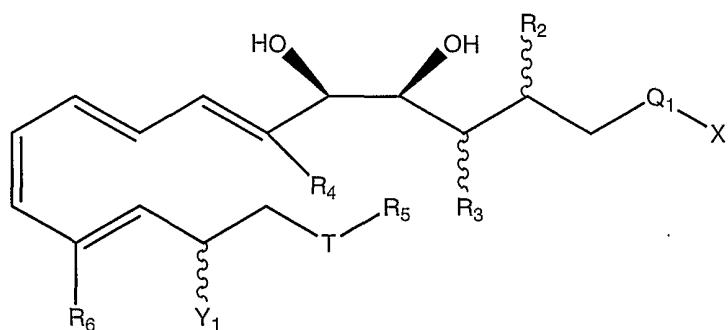
(b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

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

5

In yet another aspect, lipoxins and lipoxin analogs useful as therapeutic agents in the treatment of the maladies, disease states or conditions described throughout the specification has the formula:

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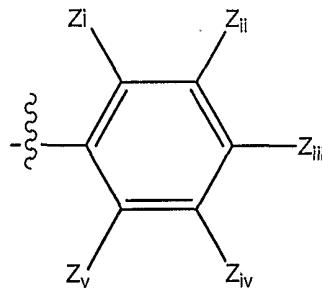
wherein X is R₁, OR₁, or SR₁;

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

20



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;

10 (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;

15 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;

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

(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

25 carbon atoms, inclusive, which may be straight chain or

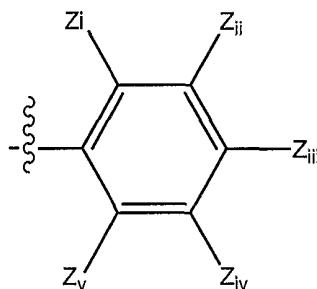
branched, provided when R_b is 0, then R_b is a hydrogen atom;

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 , -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 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;

20

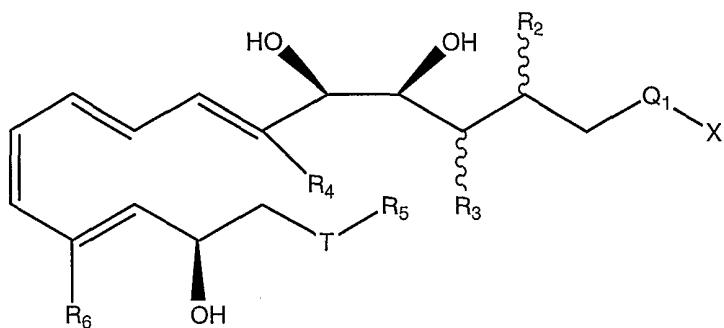
wherein R_6 is

25

- (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.

In still another aspect, lipoxins and lipoxin analogs useful as therapeutic agents in the treatment of the maladies, disease states or conditions described throughout the specification has the formula:

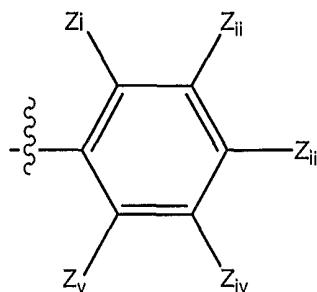


wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, 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

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, – 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;

10

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

15

- (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 a straight chain or branched; or
- (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;

20

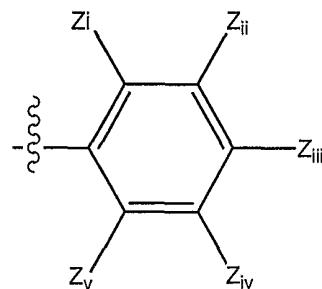
25

wherein R_4 is

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

5

wherein R_5 is



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

15

wherein R_6 is

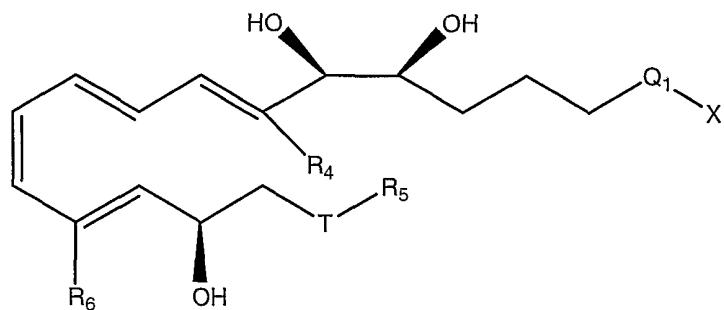
- (a) H;
- (b) an alkyl from 1 to 4 carbon atoms, inclusive, straight chain or branched;

20

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

In yet another aspect, lipoxins and lipoxin analogs useful as therapeutic agents in the treatment of the maladies, disease states or conditions described throughout the specification has the formula:

5



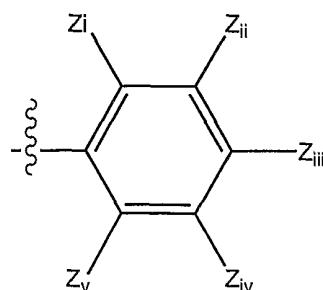
wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

10

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, $-\text{OR}_x$, 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;

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

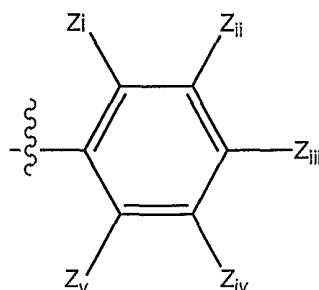
wherein R_4 is

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

20



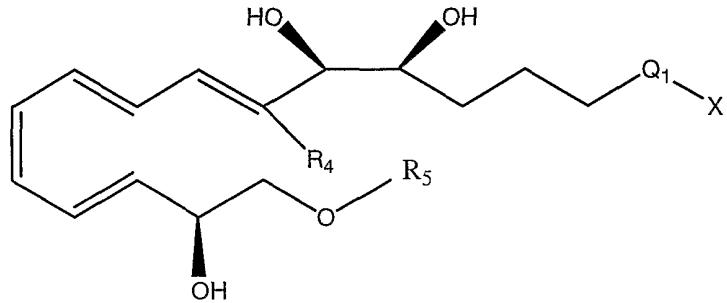
wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from $-\text{NO}_2$, $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, $-\text{OR}_x$, wherein R_x is 1 to 8 carbon atoms, inclusive, which may be a straight chain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

wherein R₆ is

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

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

In one aspect, lipoxins and lipoxin analogs useful as therapeutic agents in
10 the treatment of the maladies, disease states or conditions described throughout
the specification has the formula:

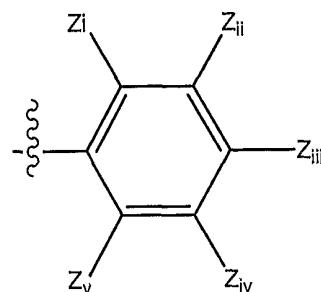


15 wherein X is R₁, OR₁, or SR₁;

wherein R₁ is

- (i) a hydrogen atom;
- (ii) an alkyl of 1 to 8 carbon atoms, inclusive,
20 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

25



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;

5

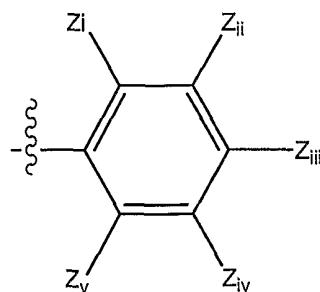
- (vii) a detectable label molecule; or
- (viii) a straight or branched chain alkenyl of 2 to 8 carbon atoms, inclusive;

10 wherein R_4 is

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

15

wherein R_5 is

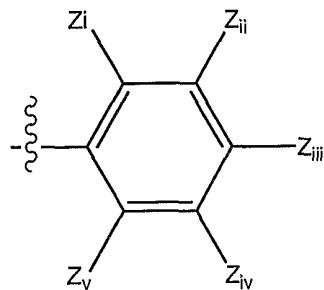


20 wherein Z_i , Z_{ii} , Z_{iii} , Z_{iv} and Z_v are each independently selected from – NO_2 , $-\text{CN}$, $-\text{C}(=\text{O})-\text{R}_1$, $-\text{SO}_3\text{H}$, a hydrogen atom, halogen, methyl, -

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

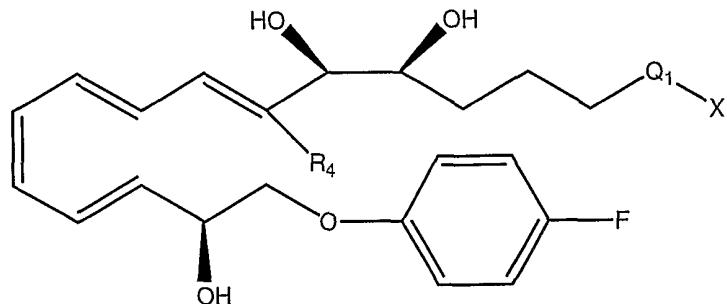
5 pharmaceutically acceptable salts thereof.

In preferred embodiments, X is OR₁ wherein R₁ is a hydrogen atom, an alkyl group of 1 to 4 carbon atoms or a pharmaceutically acceptable salt, Q₁ is C=O, R₂ and R₃, if present, are hydrogen atoms, R₄ is a hydrogen atom or methyl, 10 Q₃ and Q₄, if present, are both O, R₆, if present, is a hydrogen atom, Y₁, if present, is OH, T is O and R₅ is a substituted phenyl, e.g.,



15 wherein Z_i, Z_{ii}, Z_{iii}, Z_{iv} and Z_v 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. In certain embodiments for R₅, para-fluorophenyl and/or unsubstituted phenyl are preferred, e.g., 15-epi-16-(para-fluoro)-phenoxy- 20 LX_A₄, 16-(para-fluoro)-phenoxy-LX_A₄, 15-epi-16-phenoxy-LX_A₄ or 16-phenoxy-LX_A₄.

In still another aspect, the present invention is directed to pharmaceutical compositions including compounds having the formulae described throughout the 25 specification and a pharmaceutically acceptable carrier. In one embodiment, a preferred compound is



In one embodiment, Q₁ is a carbonyl, X is a hydroxyl or an -OR, wherein R is an alkyl group, i.e., methyl or ethyl groups, and R₄ is a hydrogen atom.

5

In other embodiments, Y₁ is a hydroxyl and the carbon bearing the hydroxyl can have an R or S configuration. In most preferred embodiments, the chiral carbon bearing the hydroxyl group, e.g., Y₁, is designated as a 15-*epi*-lipoxin as is known in the art.

10

In certain embodiments the chirality of the carbons bearing the R₂, R₃, Q₃ and Q₄ groups can each independently be either R or S. In preferred embodiments, Q₃ and Q₄ have the chiralities shown in above-referenced structures.

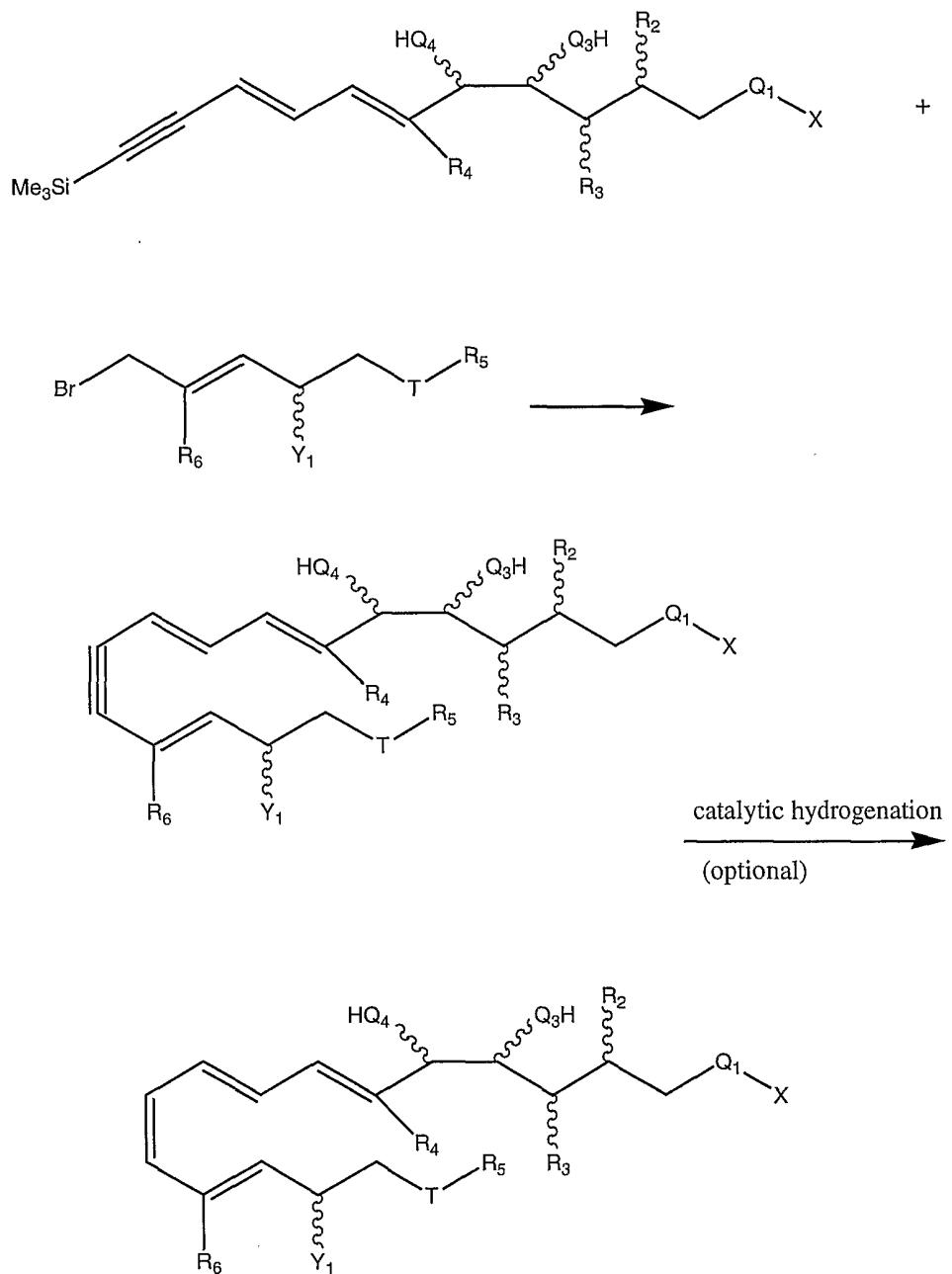
15

In preferred embodiments, R₄ is a hydrogen. In other preferred embodiments, R₆ is a hydrogen.

20

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 halogen atoms, hydroxyl groups, or ether groups.

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



5 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, 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 5 prepared by the methods of Raduchel, B. and Vorbruggen, H. (1985) *Adv. Prostaglandin Thromboxane Leukotriene Res.* 14:263. As a consequence, the acetylenic intermediates are also encompassed by the present invention as being useful for the treatments of the various maladies described herein. Similar approaches can be taken to produce LXB₄ acetylenic compounds as described in, 10 for example, US Patent No. 6,316,648.

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, 15 compounds which share the same receptor recognition site, compounds which share the same or similar lipoxin metabolic transformation region as lipoxin, and 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 20 are present, more than one stereoisomer is possible, and all possible isomeric forms are intended to be included within the structural representations shown. 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 diastereomers as well as the racemic and 25 optically resolved isomers.

The terms “corresponding lipoxin” and “natural lipoxin” refer to a 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 30 receptor on differentiated HL-60 cells, the corresponding lipoxin is LXA₄.

Where an analog has activity as an antagonist to another compound (such as leukotriene C4 and/or leukotriene D4), which is antagonized by a naturally-occurring lipoxin, that natural lipoxin is the corresponding lipoxin.

5 "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 or complex of macromolecules, including an enzyme and its cofactor. For example, lipoxin A₄ analogs have an active region comprising C₅—C₁₅ of natural 10 lipoxin A₄. Similarly, for example, lipoxin B₄ analogs have an active region comprising C₅-C₁₄ of natural lipoxin B₄.

The term "recognition site" or receptor is art-recognized and is intended to refer generally to a functional macromolecule or complex of macromolecules with which certain groups of cellular messengers, such as hormones, 15 leukotrienes, 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 may be cloned. A receptor may normally exist or it may be induced by a disease 20 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.

The term "metabolic transformation region" is intended to refer generally to that portion of a lipoxin, a lipoxin metabolite, or lipoxin analog including a 25 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. Similarly, such regions are possibly located within ATL and LXB₄ analogs. A 30 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.

5 The term “detectable label molecule” is meant to include fluorescent, phosphorescent, and radiolabeled molecules used to trace, track, or identify the 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.

10 The term “labeled analog” is further understood to encompass compounds which are labeled with radioactive isotopes, such as but not limited to 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.

15 The term “inhibits metabolism” means the blocking or reduction of activity of an enzyme which metabolizes a native eicosanoid. 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 20 lipoxin analog, including a lipoxin analog metabolite, a lipoxin, or a lipoxin metabolite. Similarly, this also applies to ATL and LXB₄ analogs.

25 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, ATL or LXB₄ analogs. Two nonlimiting examples of LXA₄ analog that resists metabolism are 1) a structure which can not be oxidized to the 15-oxo 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.

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 lipoxin, lipoxin analogs, ATL or LXB₄ analogs. A nonlimiting example of a 5 LX₄ analog which more slowly undergoes metabolism is a structure which has a higher transition state energy for C-15 dehydrogenation than does LX₄ because the analog is sterically hindered at the C-16.

The term "tissue" is intended to include intact cells, blood, blood 10 preparations such as plasma and serum, bones, joints, muscles, smooth muscles, and organs.

The term "halogen" is meant to include fluorine, chlorine, bromine and iodine, or fluoro, chloro, bromo, and iodo.

The invention is further illustrated by the following examples which in no way should be construed as being further limiting. The contents of all references, 15 pending patent applications and published patent applications, cited throughout this application, including those referenced in the background section, are hereby incorporated by reference. It should be understood that the models used throughout the examples are accepted models and that the demonstration of efficacy in these models is predictive of efficacy in humans.

20

Materials and Methods

Cell Culture

Human umbilical vein endothelial cells (HUVEC) were isolated by 0.1% 25 collagenase digestion (Worthington Biochemical, Bedford, MA) and propagated on gelatin-coated (0.1%) tissue culture plates in medium 199 (Gibco BRL, Grand Island, NY) supplemented with 20% heat-inactivated fetal bovine serum (BioWhittaker, Walkersville, MD), 50 µg/ml of endothelial cell mitogen (Biomedical Technologies, Stoughton, MA), 8 U/ml heparin (APP, Los Angeles, CA), 50 U/ml penicillin and 15 µg/ml streptomycin. Only passages 2 and 3 were 30 used in reported experiments.

Endothelial Cell Proliferation

HUVEC (5×10^3) were plated in 96-well plates coated with 0.1% gelatin for 1h at room temperature. After 24 h, the medium was removed and replaced with fresh medium 199 supplemented with 5% fetal bovine serum and different 5 concentrations of recombinant human VEGF₁₆₅ (R&D Systems, Minneapolis, MN), LTD₄ or LTB₄. Endothelial cells were enumerated after 72 h using the MTT® assay (Sigma, St. Louis, MO) (15). ATL-1 was prepared as in Clish et al. 10 (13). Percent inhibition was evaluated in a similar manner and included a 15 minute incubation (37° C) with 15-epi-lipoxin A₄, 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄ methyl ester (ATL-1), LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ prior to the addition of agonists. All incubations were performed in triplicate. Before each experiment the integrity and concentration of ATL-1, LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ was assessed by physical methods including LC/MS/MS and UV (13).

15

Endothelial Cell Migration

VEGF, ATL-1 or vehicle were added to the lower wells of a 48-well chemotaxis chamber (NeuroProbe, Cabin John, MD). The wells were overlaid with a 10 µm pore size polycarbonate filter coated with 0.1% gelatin. HUVEC 20 (1×10^6) were placed in the upper wells and the chamber was incubated (37°C, 5% CO₂ for 12 h). Following incubations, filters were removed, scraped of cells from the upper surface, fixed and stained with Diff-Quik (Dade Behring, Newark, DE). Cells that migrated across the filter toward the lower surface were 25 enumerated by light microscopy; four fields were counted at high magnification (100X). Incubations were performed in triplicate. To assess inhibition, endothelial cells were suspended in media with vehicle or ATL-1 for 15 min before placement in the chamber.

Quantitative Determination of DNA Fragmentation

DNA fragmentation in individual apoptotic cells was quantitated using a 30 photometric enzyme immunoassay (Apoptosis detection kit: R&D Systems). HUVECs grown in 96-well microtiter plates (5×10^3 cells/well) were incubated for 3 days, fixed with 3.7% formaldehyde, and permeabilized with proteinase K

before the labeling. Biotinylated nucleotides are incorporated onto the DNA fragments and detected by using streptavidin-horseradish peroxidase conjugate followed by the substrate TACS-Sapphire.

Inflammatory angiogenesis

5 Angiogenesis was assessed with murine air pouches that were raised via subcutaneous injection of sterile air (3 ml) beneath the dorsal skin of anesthetized mice (BALB/c, male 6-8 wk). After 24 hrs, ATL-1 (10 µg/pouch) or vehicle was delivered locally, immediately before the injection of VEGF (1 µg/pouch). The vascular content was assessed by the formation of vascular casts (as in 16).
10 Briefly, mice were anesthetized (at 144 h) and peripheral vasodilation was raised by placing the animals in a heated jacket (40° C, 10 min). Vascular casts were formed by the i.v. injection of 1 ml 5% carmine red (Sigma) in 5% gelatin solution warmed to 40° C. Air pouch linings were dissected and weighed. The tissue was then dissolved in 2 ml of 3 N NaOH solution for 0.5 h, 21° C and
15 completely digested in hot water (56° C) for 10 min. Digested samples were centrifuged (2500 rpm, 15 min) and filtered through a 0.45 µm filter. The dye content was quantified employing a 96-well plate spectrophotometer at 530 nm using a calibration curve. The results were expressed as vascular index (VI) as micrograms carmine dye/milligram weight of tissue, for n = 4 animals/group. For
20 visualization of the vasculature, the dorsal surface of the pouches was excised and fixed in formalin for 48 h. The tissues were dehydrated with 100% ethanol (5 days, 4° C) and cleared by immersion in cedar wood oil for 2 weeks. In another set of experiments, mice were anesthetized and injected i.v. with 200 µl of 0.05 g/ml fluorescein isothiocyanate-dextran (Sigma) in PBS at 144 h immediately
25 before sacrifice. Dissected linings were fixed, mounted on glass slides and examined for fluorescence (Nikon Eclipse model E600). In both protocols, the observers were not blinded to the treatments.

Immunohistochemistry

Air pouch membranous tissues were fixed in 10% buffered formalin
30 overnight and processed for paraffin embedding. Five-micrometer paraffin sections of membrane tissue cut on face were used for immunohistochemistry for CD31 expression. Briefly, slides were deparaffinized and pretreated in 0.25% trypsin (Sigma Chemical) for 20 min at 37° C, followed by washing in distilled

water. All further steps were performed at room temperature in a hydrated chamber. Slides were pretreated with Peroxidase Block (DAKO, Carpinteria, CA) for 5 min to quench endogenous peroxidase activity, followed by a 1:5 dilution of goat serum in 50 mM Tris-Cl, pH 7.4, for 20 min to block nonspecific binding sites. Primary rat anti-murine CD31 antibody (BD PharMingen, San Diego, CA) was applied at a 1:100 dilution in 50 mM Tris-Cl, pH 7.4, with 3% goat serum for 1 h. After washing in 50 mM Tris-Cl, pH 7.4, secondary rabbit anti-rat antibody (DAKO) was applied at a 1:200 dilution in 50 mM Tris-Cl, pH 7.4, with 3% goat serum for 30 min. Slides were washed again in 50 mM Tris-Cl, pH 7.4, and goat anti-rabbit horse-radish peroxidase-conjugated antibody (Envision detection kit: DAKO) was applied for 30 min. After further washing, immunoperoxidase staining was developed using a DAB chromogen kit (DAKO) per the manufacturer and counterstained with hematoxylin.

Statistical Analysis

Results are presented as means \pm S.E.M. Statistical evaluation of the results was performed by analysis of variance, and values of $P < 0.05$ were taken to represent statistically significant differences between group means.

Results and Discussion

The aspirin-triggered lipoxin A₄ stable analog (denoted ATL-1) proved a potent inhibitor of VEGF-stimulated proliferation of HUVECs (IC_{50} of ~ 3 nM) (Figure 1). Inhibition was concentration-dependent and maximal plateau at 10 nM and was partially reversed (from 38.0 ± 2.5 to $78.0 \pm 6.21\%$ of VEGF-stimulated proliferation, $P < 0.05$) by incubating the cells with genistein, an inhibitor of tyrosine kinase activity (50 μ M, 5 min, 37° C, n=3). Even at high concentrations (100 nM), the ATL-1 analog alone had no apparent actions on HUVEC proliferation (Figure 1 inset). In sharp contrast, LXB₄ stable analogs increased proliferation. Because these are related structures, the separate actions of ATL and LXB₄ analogs with these cells indicate that the ATL-1 response is highly stereoselective. Also, a direct comparison of ATL-1 with native LXA₄ and 15-epi-LXA₄ at equimolar concentrations (10 nM) in a representative experiment showed that ATL-1 > 15-epi-LXA₄ > LXA₄ in rank order of activity (e.g., 63.3 ± 3.3 , 59.8 ± 1.8 and $38.1 \pm 2.0\%$ inhibition, respectively). After exposure of the cells to ATL-1, $\sim 98\%$ of the HUVEC remained viable, as

determined by trypan blue exclusion assay, indicating that the compound was not cytotoxic. In addition to being an endothelial cell-specific mitogen, VEGF is also an endothelial cell survival factor, thus promoting angiogenesis not only by stimulating cell proliferation but also by inhibiting endothelial cell apoptosis (17).

5 To determine whether this new inhibitory action of ATL-1 on HUVEC proliferation involved apoptosis, DNA fragments were quantitated (under Materials and Methods). Neither the ATL-1 (100 nM) alone nor in combination with VEGF (3 ng/ml) affected DNA fragmentation pattern (n = 2, d = 3), suggesting that the antiproliferative actions of the ATL-1 analog were not a
10 results of induction of apoptosis in endothelial cells. Binding of α v integrins by endothelial cells is accompanied by a decrease in the tumor-suppressor p53 activity and inhibition of apoptotic pathways, thereby facilitating the formation of new blood vessels (18). Along these lines, it was recently shown that p53 can up-regulate human 15-lipoxygenase promoter activity, providing the first link
15 between this enzyme's activity and an established tumor-suppressor gene (19). Also of interest, over-expression of 15-lipoxygenase enhances endogenous LXA₄ formation that in turn inhibits progression of glomerulonephritis (20). Both ATL and LXA₄ share sites of action that are receptor cell type, and tissue-specific, and together with their stable analogs display potent anti-inflammatory actions (21,
20 23 and 14).

25 Endothelial cell migration is an essential component of the angiogenic process, providing directionality for the budding capillary toward the angiogenic stimulus (3). Therefore, endothelial migration was assessed with ATL. VEGF (3 ng/ml) was added to the lower compartment of a chemotaxis chamber and cell migration across a 10- μ m pore-size gelatin-coated filter was quantitated (Figure 2). Results in Figure 2B showed that ATL-1, LXA₄, 15-epi-LXA₄ and 15-R/S-methyl, LXA₄ (ATL-1 shown as representative) gave concentration-dependent inhibition of VEGF-stimulated HUVEC migration with a maximum level of inhibition (~45%) at 10 nM ATL. As observed with proliferation assays, ATL-1,
30 LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ alone, even at higher concentrations (100 nM), did not induce endothelial cell migration (Figure 2A), findings which suggest that ATL, LXA₄, 15-epi-LXA₄ or 15-R/S-methyl, LXA₄ play a role in blocking the early stages of cell migration to sites of neovascularization.

Like VEGF (Figure 1), LTD₄ also stimulated proliferation of HUVEC (42 \pm 1.2%) with a maximum at 10 nM, similar to the response obtained with VEGF (52.7 \pm 1.6%). In contrast, LTB₄ at 10 nM did not give a significant response with these cells (Figure 3B). The mitogenic action of LTD₄ (10 nM) was 5 antagonized by exposure of the cells to ATL-1 (0.1-100 nM) with an IC₅₀ of ~ 3 nM (Figure 3A). LTD₄ did not enhance or inhibit the VEGF-stimulated proliferation.

The antiproliferative actions of native lipoxins were first found with the 10 human lung adenocarcinoma cell line (23) and recently with human renal mesangial cells (24). The present findings, together with endothelial cell results, draw attention to the potential regulatory role for endogenous ATL in proliferative diseases. The actions of LX, ATL and stable analogs are transduced by a high affinity transmembrane receptor (ALXR) identified in several cell types (for a review, see Chiang et al. (25)). In mesangial cells, LXA₄ interacts with its 15 own high affinity receptor (i.e., ALXR) as well as with a subclass of peptido-leukotriene receptors (cysLT₁), where LXA₄ is a partial agonist (24). In this regard, LXA₄ and its bioactive stable analogs effectively displace [³H]LTD₄ specific binding to vascular endothelial cells (26). Also, recent findings provide the first evidence that ATL specifically antagonizes LTD₄ specific binding at 20 recombinant human cysLT₁ cloned from endothelial cells, as well as acts at specific LXA₄ receptors (21). Since ATL-1 proved to be a potent inhibitor of HUVEC proliferation (Figures 1, 2 and 3), it was determined whether ATL affected angiogenesis *in vivo*.

During chronic inflammation, new vessels are required not only for the 25 maintenance of tissue perfusion, but also to allow increased cellular traffic (27). Therefore, to this end angiogenesis was assessed *in vivo* using a well established murine chronic granulomatous air pouch model, and the 6-day time interval was selected because it was shown to give near maximal vascular density (16). ATL-1 injected locally (10 μ g/pouch) immediately before the administration of VEGF 30 (1 μ g/pouch) gave a ~ 48% reduction in the vascular index (Figure 4A). For comparison, ATL-1 (10 μ g/mouse or 0.4 mg/kg mouse) proved to be much more potent than other described anti-angiogenic agents that required much higher doses, including steroids at 1-3 mg/kg (16), or the COX-2 inhibitors, which require 1-6 mg/kg (28).

Figure 4B shows representative vascular casts from typical day 6 air pouch linings. In the mice given VEGF (Figure 4B, bottom left panel), there is an established neovasculature with an extremely high degree of vascular density compared to only slightly dilated capillaries in the ATL-treated animals, where 5 there was routinely clearly reduced vascular density (Figure 4B, bottom right panel). In another set of experiments, fluorescein isothiocyanate-dextran was used to visualize the vessels in this region. In sharp contrast to the actions of ATL-1, when LTB₄, another lipoxygenase pathway product, was given alone at the same dose as ATL-1 (10 µg LTB₄/pouch), LTB₄ stimulated 10 neovascularization (n = 2).

Figure 5 shows photomicrographs of the dorsal linings dissected at day 6. Again, profound angiogenesis was demonstrated with extensive vascular networks in VEGF-treated pouch (Figure 5, bottom left panel). Here too, treatment with ATL-1 (10 µg/pouch) gave striking reduction of VEGF-elicited 15 vasculature, as exemplified by the lack of visible fine capillaries (Figure 5, bottom right panel).

It is important to note that ATL-1 at this dose (10 µg/mouse i.v.) does not evoke apparent changes in mean arterial pressure, excluding a possible action of ATL-1 at the level of vascular tone. This is particularly noteworthy because, at 20 high doses, LXA₄ can stimulate vasodilation in certain vascular beds. The present in vivo experiments were performed in separate series to evaluate histology and the presence of a vascular marker by using immunohistochemical staining of the murine air pouch with the vascular endothelial cell marker CD31 (Figure 6).

25 Platelet/endothelial cell adhesion molecule-1 (or CD31) is a member of the Ig superfamily that is strongly expressed at the endothelial cell-cell junction, is present on platelets as well as leukocytes, and is held to play a role in angiogenesis and in transendothelial migration of leukocytes. Immunohistochemical staining for CD31 in the pouches showed that, in the 30 VEGF-treated mice, strong specific endothelial cell staining was present and identified a prominent vascular network (Figure 6C). In contrast, a marked diminution of vessels was observed in VEGF-treated mice that were also treated with the LXA₄ analog (Figure 6D). The levels of mild nonspecific staining

associated with these air pouches were essentially identical to those of the air pouch sections from mice treated with either vehicle alone (Figure 6A) or with LX analog alone (Figure 6B), namely, mild nonspecific staining of inflammatory cells, predominantly leukocytes and macrophages, that are known to be
5 associated with these air pouches created from murine skin. Taken together, these findings indicate that ATL reduced VEGF-stimulated angiogenesis *in vivo*, suggesting that LXA₄ and 15-epi-LXA₄ can regulate these actions *in vivo*.

Results from many clinical and laboratory studies have demonstrated protective effects of aspirin in several forms of human cancer, including lung, 10 colon and breast cancer, yet its potential anti-cancer mechanism is not clear (see Ref. 9). ASA is thought to act, in part, via reduction of angiogenesis, which might be related to the ability of ASA to inhibit prostanoid biosynthesis (7). More recently, ASA was found to trigger a novel switch in eicosanoid biosynthesis as the acetylation of COX-2 enables the enzyme to produce 15R-HETE that is
15 converted to 15-epi-lipoxins, also known as ATL, during cell-cell interactions *in vitro* and *in vivo* (11, 12, 14). ATLs as well as their stable bioactive analogs are potent inhibitors of several key events in acute inflammation, such as PMN chemotaxis and transmigration across both endothelial and epithelial cells, as well as diapedesis from postcapillary venules (13, 14). The analogs of ATL
20 mimic both endogenous ATL and LX actions and were designed to resist rapid enzymatic inactivation *in vivo*. Bioactive analogs of 15-epi-LXA₄ were also found to compete at both the ALXR on leukocytes and the cysLT₁ receptor present on vascular endothelial cells. In addition, these novel aspirin-triggered mediators inhibit cytokine release and can act at the gene transcriptional level
25 (29) to redirect local cytokine-chemokine axis (30), actions that are both of interest in the angiogenic process (3). It should be noted that most if not all other eicosanoids examined to date are pro-angiogenic including leukotrienes (e.g., LTD₄ and LTB₄, see the following table and Figure 3) (5, 6, 28). In view of this, it was surprising that ATL-1, LXA₄, 15-epi-LXA₄ and 15-R/S-methyl, LXA₄
30 possessed antiangiogenic activity.

LXB₄-induced HUVEC Proliferation

Compound	Percent of Proliferation
14-epi-LXB ₄ 0.1 nM	15.5 ± 1.0
14-epi-LXB ₄ 1.0 nM	18.3 ± 1.6
14-epi-LXB ₄ 10 nM	27.0 ± 5.0
14-epi-LXB ₄ 100 nM	27.7 ± 1.3
15-epi-LXB ₄ 0.1 nM	20.1 ± 1.2
15-epi-LXB ₄ 1.0 nM	26.1 ± 2.8
15-epi-LXB ₄ 10 nM	34.0 ± 2.5
15-epi-LXB ₄ 100 nM	35.2 ± 5.2
15-epi-LXB ₄ -acetylenic 0.1 nM	18.7 ± 1.6
15-epi-LXB ₄ -acetylenic 1.0 nM	23.7 ± 0.4
15-epi-LXB ₄ -acetylenic 10 nM	36.4 ± 3.7
15-epi-LXB ₄ -acetylenic 100 nM	37.4 ± 5.3

VEGF-induced proliferation: 50.4 ± 4.0

In summary, the present results demonstrate that an aspirin-triggered-lipoxin, 15-epi-LXA₄ analog, is a potent inhibitor of angiogenesis and of endothelial cell proliferation *in vivo*. Together these results reveal a novel action of 15-epi-lipoxins and suggest a role for the aspirin-triggered lipoxin circuit (14) 5 as a potential mechanism that can contribute to aspirin's recognized anti-angiogenic and anti-inflammatory properties (2, 7, 10). With increasing insight into the fundamental role of angiogenesis within a broad range of physiological as well as disease processes (1-3), the modulation of vascular growth could be a previously unappreciated and important strategic action for ATL, the natural 10 endogenous lipoxin mimetic and their synthetic analogs.

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25 Those skilled in the art will know, or be able to ascertain, using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. These and all other equivalents are intended to be encompassed by the following claims. All publications and references cited herein, including those in the background section, are expressly incorporated herein by reference in their entirety.

30

CLAIMS

What is claimed is:

1. A method to prevent, reduce or inhibit angiogenesis comprising the step of administering to a subject an effective amount of a lipoxin A₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that angiogenesis is prevented, reduced or inhibited in the subject.
5
2. The method of claim 1, wherein the lipoxin A₄ compound is LXA₄ or 15-R/S-methyl, LXA₄.
3. A method to prevent, reduce or inhibit angiogenesis comprising the step of administering to a subject an effective amount of a 15-epi-lipoxin A₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that angiogenesis is prevented, reduced or inhibited in the subject.
10
4. The method of claim 3, wherein the 15-epi-lipoxin A₄ compound is 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄.
- 15 5. A method for treating a subject for restenosis in a tissue wherein smooth muscle cell migration occurs following angioplasty, comprising administering to said subject a composition comprising an effective amount of a lipoxin A₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that restenosis in a tissue wherein smooth muscle cell migration occurs following angioplasty is prevented, reduced, or inhibited.
20
6. The method of claim 5, wherein the lipoxin A₄ compound is LXA₄ or 15-R/S-methyl, LXA₄.
- 25 7. A method for treating a subject for restenosis in a tissue wherein smooth muscle cell migration occurs following angioplasty, comprising administering to said subject a composition comprising an effective amount of a 15-epi-lipoxin A₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that restenosis in a tissue wherein smooth muscle cell migration occurs following angioplasty is prevented, reduced, or inhibited.

8. The method of claim 15, wherein the 15-epi-lipoxin A₄ compound is 15-epi-16-(*para*-fluoro)-phenoxy-lipoxin A₄.

9. A method to facilitate wound healing in a subject comprising the step of administering to the subject in need thereof, an effective amount of an 5 LXB₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that wound healing in the subject is facilitated.

10. The method of claim 9, wherein the LXB₄ compound is selected from the group consisting of 14-epi-LXB₄, 15-epi-LXB₄ and 15-epi-LXB₄-acetylenic.

11. A method to facilitate angiogenesis in a subject comprising the step of administering to the subject in need thereof, an effective amount of an LXB₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that angiogenesis in the subject is facilitated.

12. The method of claim 11, wherein LXB₄ is excluded.

13. The method of claim 11, wherein the LXB₄ compound is selected from the group consisting of 14-epi-LXB₄, 15-epi-LXB₄ and 15-epi-LXB₄-acetylenic.

14. A method to facilitate neovascularization in a subject comprising the step of administering to the subject in need thereof, an effective amount of an 20 LXB₄ compound and pharmaceutically acceptable salts, esters, amides, and prodrugs thereof, such that neovascularization in the subject is facilitated.

15. The method of claim 14, wherein LXB₄ is excluded.

16. The method of claim 14, wherein the LXB₄ compound is selected from the group consisting of 14-epi-LXB₄, 15-epi-LXB₄ and 15-epi-LXB₄-acetylenic.

17. A method to facilitate cardiac revascularization in a subject comprising the step of administering to the subject in need thereof, an effective amount of an LXB₄ compound and pharmaceutically acceptable salts, esters,

amides, and prodrugs thereof, such that cardiac revascularization in the subject is facilitated.

18. The method of claim 17, wherein the LXB₄ compound is selected from the group consisting of 14-epi-LXB₄, 15-epi-LXB₄ and 15-epi-LXB₄-
5 acetylenic

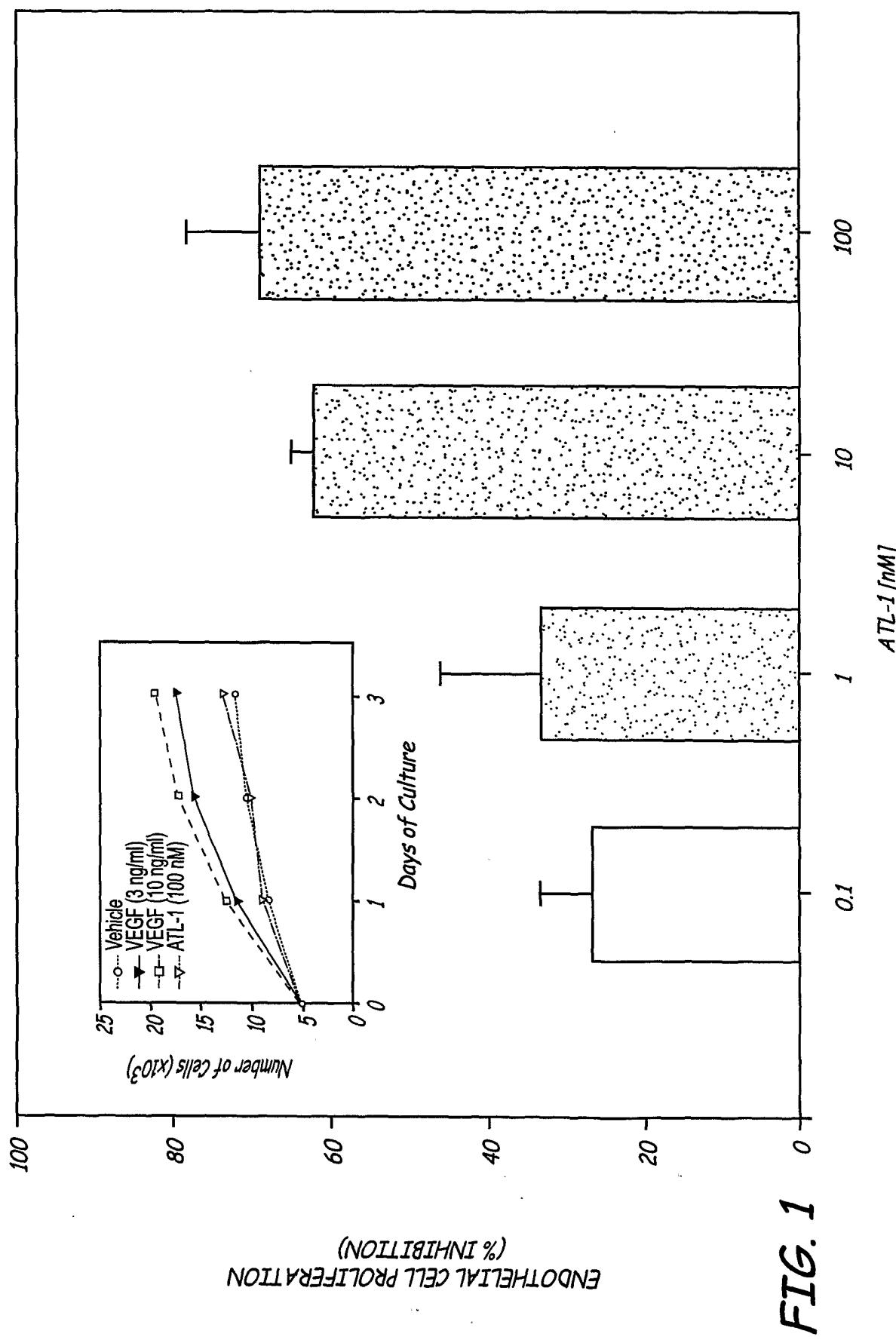
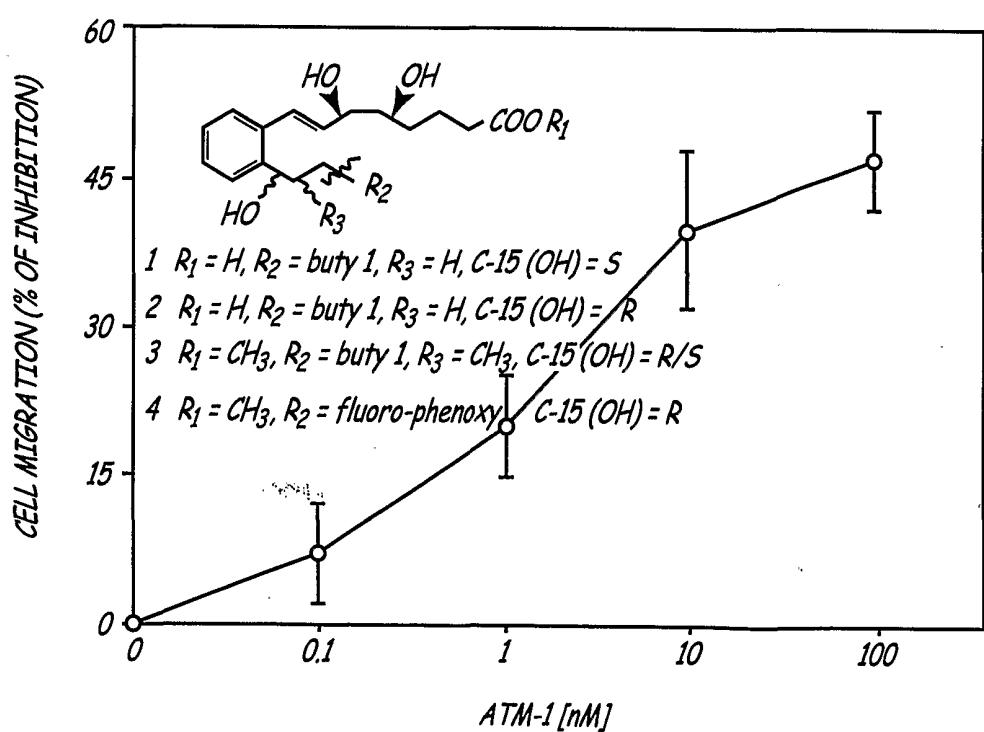
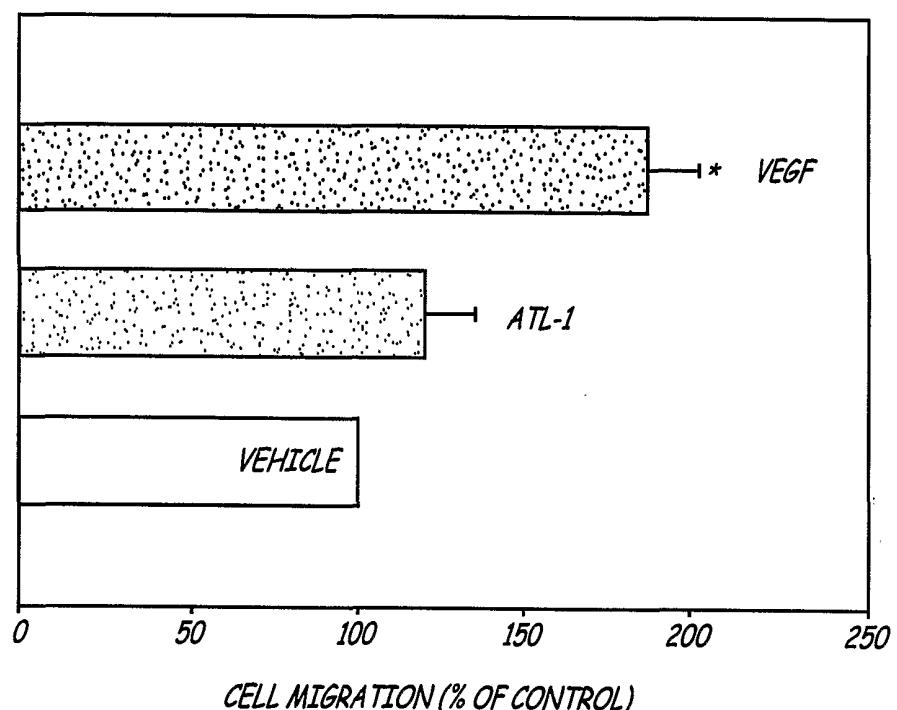


FIG. 2



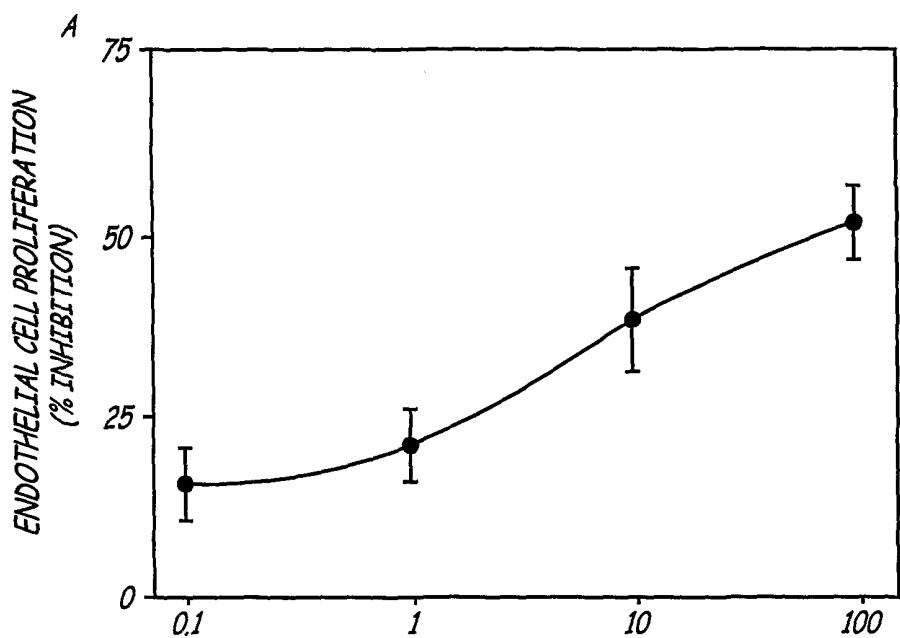


FIG. 3A

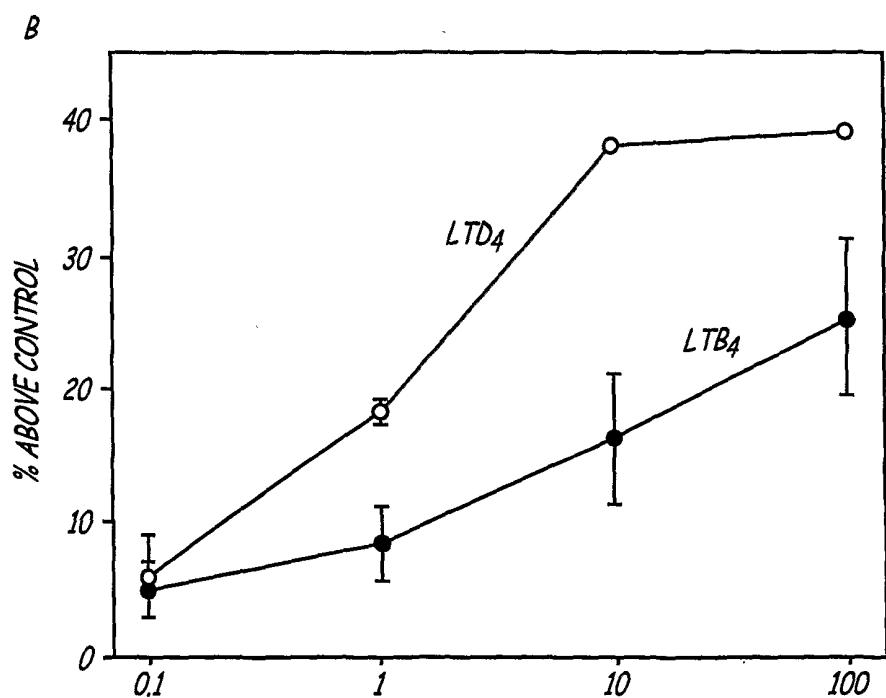


FIG. 3B

FIG. 4A

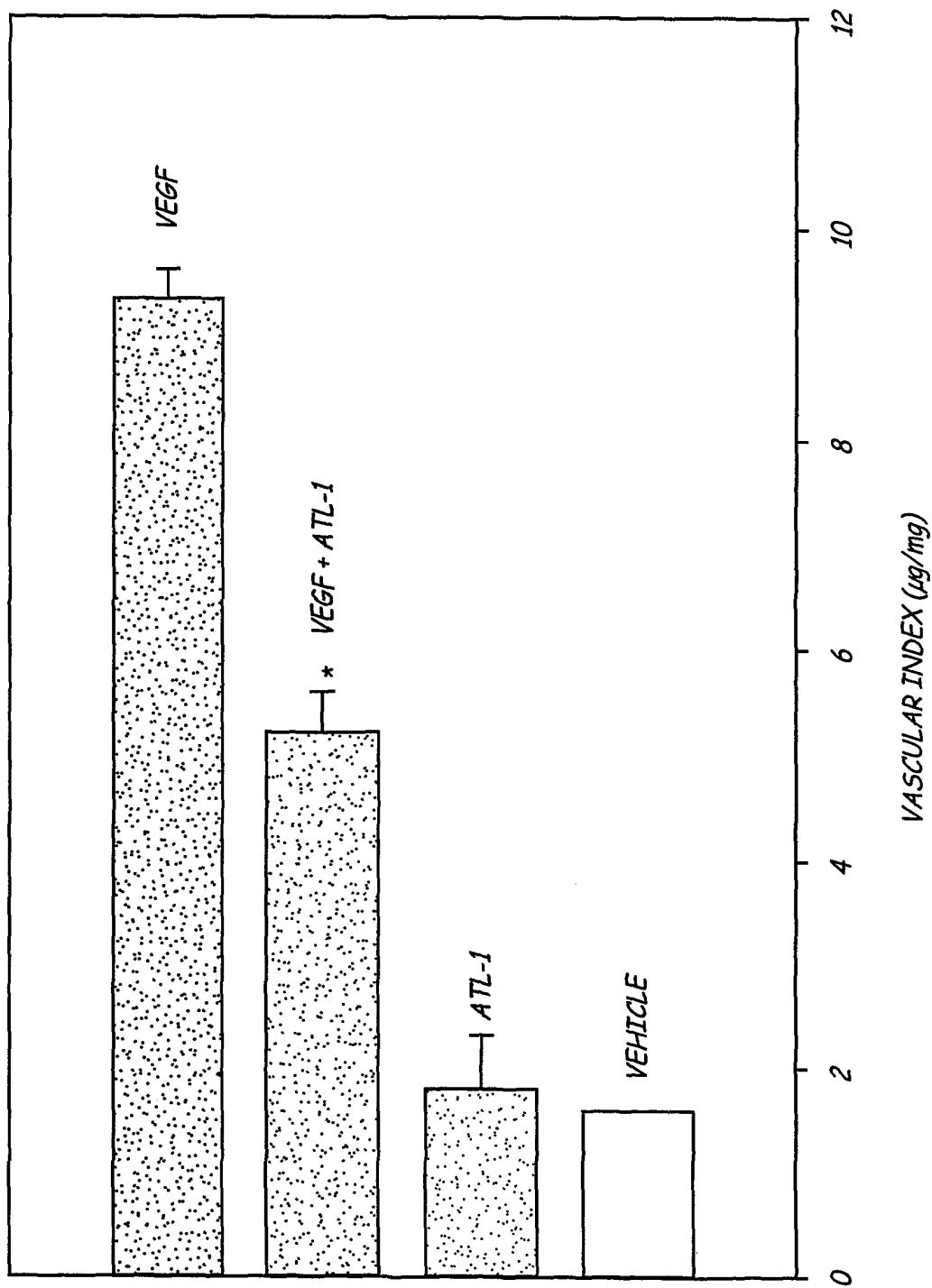


FIG. 4B

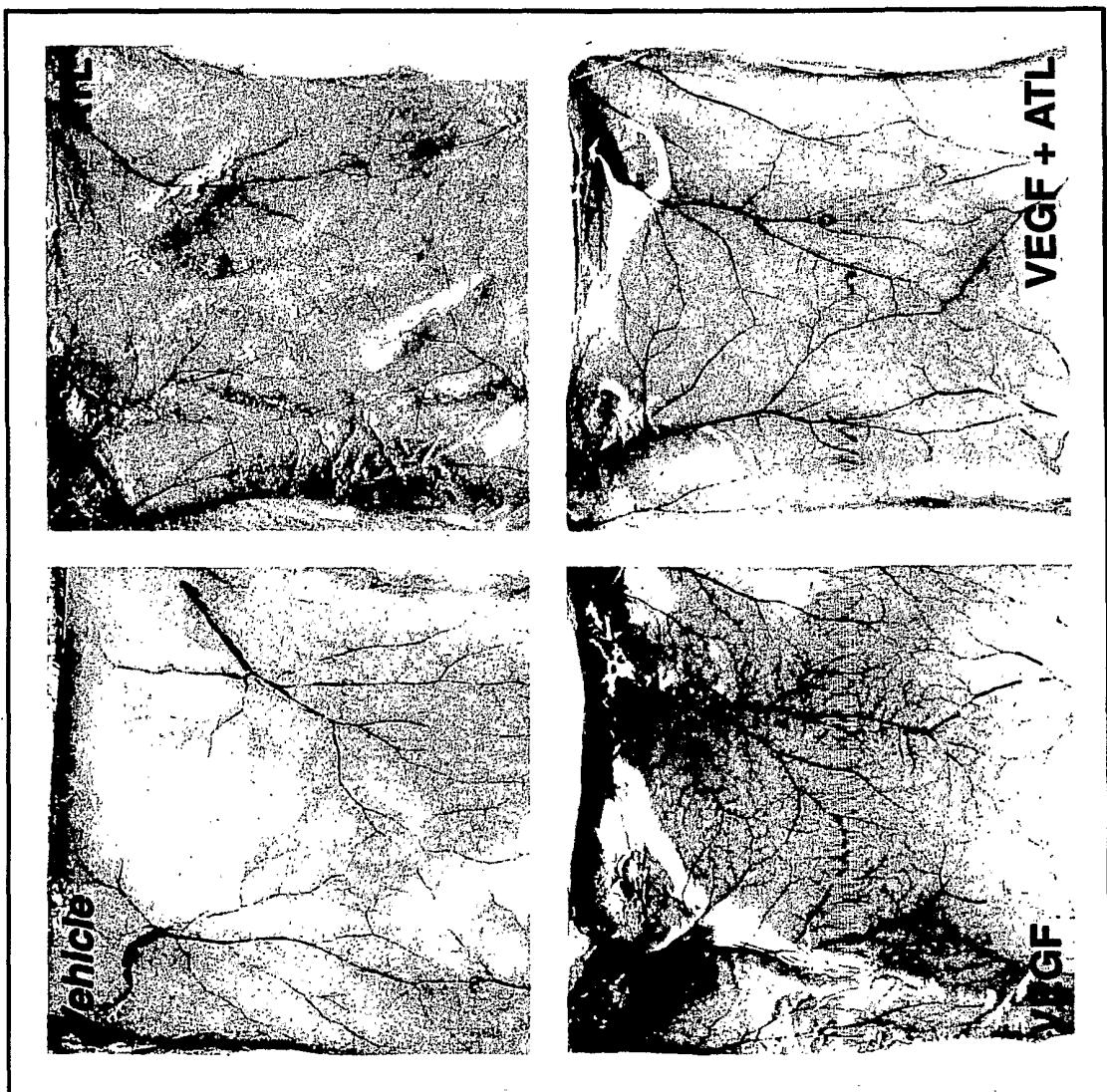


FIG. 5

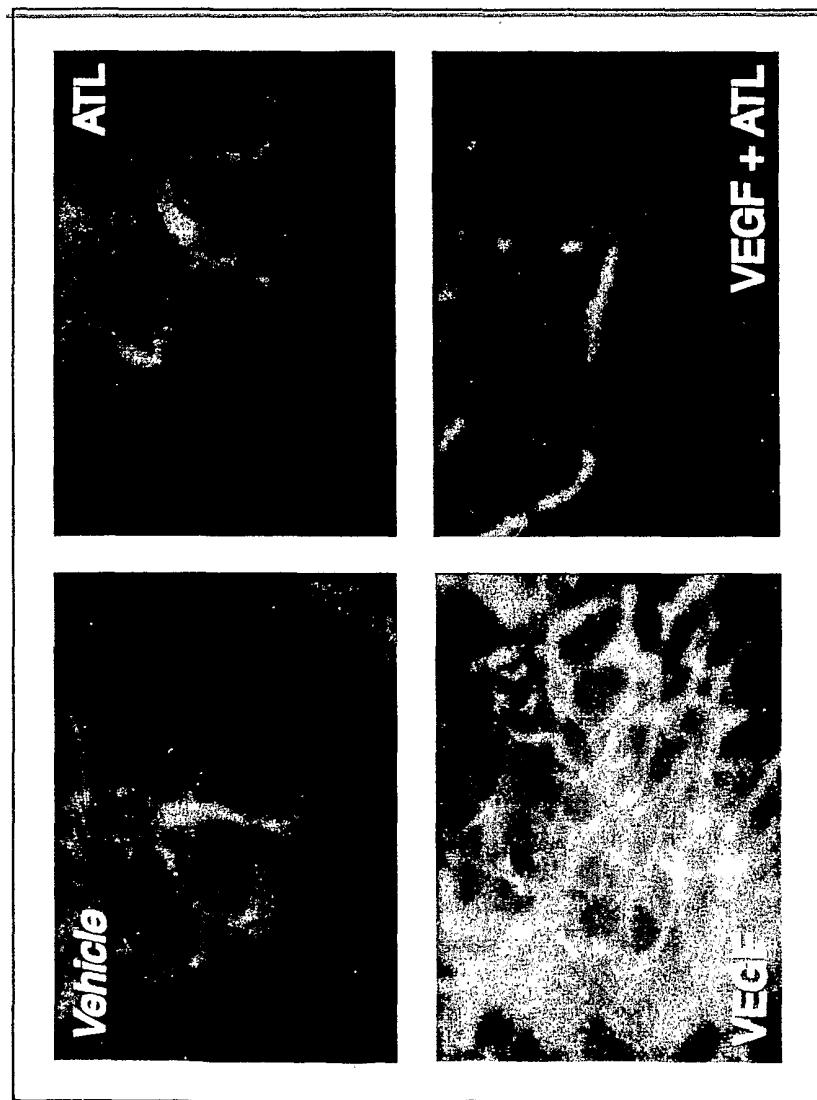


FIG. 6

