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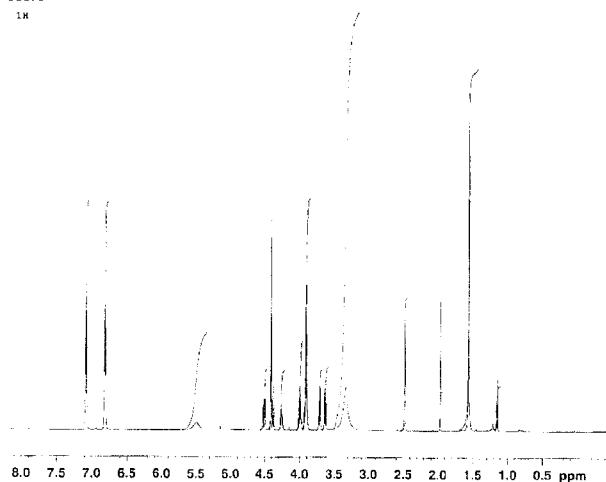
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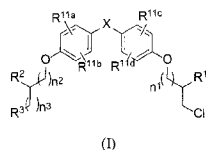
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[Continued on next page]

(54) Title: HETEROCYCLIC COMPOUNDS FOR CANCER IMAGING AND TREATMENT AND METHODS FOR THEIR USE

FIG. 5
1H

(57) Abstract: Compounds having a structure of Formula I: or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein R¹, R², R³, R^{11a}, R^{11b}, R^{11c}, R^{11d}, X, n¹, n², and n³ are as defined herein, are provided. Uses of such compounds for modulating androgen receptor activity, imaging diagnostics in cancer and therapeutics, and methods for treatment of subjects in need thereof, including prostate cancer are also provided.





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HETEROCYCLIC COMPOUNDS FOR CANCER IMAGING AND TREATMENT AND METHODS FOR THEIR USE

CROSS-REFERENCE TO RELATED APPLICATION

This application claims priority to U.S. Provisional Application No. 62/102,719
5 filed January 23, 2015, the disclosure of which is herein incorporated by reference in its entirety for all purposes.

STATEMENT OF GOVERNMENT INTEREST

This invention was made in part with government support under Grant No. 2R01
CA105304 awarded by the National Cancer Institute. The United States Government has
10 certain rights in this invention.

Technical Field

This invention generally relates to bisphenol-related compounds and their use for treatment of various indications. In particular the invention relates to bisphenol ether compounds having novel heterocyclic groups and their use for treatment of various
15 cancers, for example all stages of prostate cancer, including androgen dependent, androgen sensitive and castration-resistant prostate cancers. This invention also relates to bisphenol-related compounds and their use for modulating androgen receptor (AR) activity.

This invention further relates to radiolabeled bisphenol-related compounds and their use in methods for imaging the prostate gland. For example, in certain embodiments the
20 compounds are useful for imaging benign prostate diseases such as benign prostate hyperplasia. In other embodiments, the compounds are useful for imaging cancerous prostate diseases, such as prostate cancer tumors. In certain embodiments the invention relates to radioactive ¹²³I compounds and their use as an imaging tool in prostate cancer and benign prostate diseases. The disclosed compounds find utility in any number of
25 imaging applications, including imaging of androgen receptor (AR) splice variants in prostate cancers, including all stages and androgen dependent, androgen-sensitive and castration-resistant prostate cancers (also referred to as hormone refractory, androgen-

independent, androgen deprivation resistant, androgen ablation resistant, androgen depletion-independent, castration-recurrent, anti-androgen-recurrent).

Description of the Related Art

Androgens mediate their effects through the androgen receptor (AR). Androgens play a role in a wide range of developmental and physiological responses and are involved in male sexual differentiation, maintenance of spermatogenesis, and male gonadotropin regulation (R. K. Ross, G. A. Coetzee, C. L. Pearce, J. K. Reichardt, P. Bretsky, L. N. Kolonel, B. E. Henderson, E. Lander, D. Altshuler & G. Daley, *Eur Urol* 35, 355-361 (1999); A. A. Thomson, *Reproduction* 121, 187-195 (2001); N. Tanji, K. Aoki & M. Yokoyama, *Arch Androl* 47, 1-7 (2001)). Several lines of evidence show that androgens are associated with the development of prostate carcinogenesis. Firstly, androgens induce prostatic carcinogenesis in rodent models (R. L. Noble, *Cancer Res* 37, 1929-1933 (1977); R. L. Noble, *Oncology* 34, 138-141 (1977)) and men receiving androgens in the form of anabolic steroids have a higher incidence of prostate cancer (J. T. Roberts & D. M. Essenhight, *Lancet* 2, 742 (1986); J. A. Jackson, J. Waxman & A. M. Spiekerman, *Arch Intern Med* 149, 2365-2366 (1989); P. D. Guinan, W. Sadoughi, H. Alsheik, R. J. Ablin, D. Alrenga & I. M. Bush, *Am J Surg* 131, 599-600 (1976)). Secondly, prostate cancer does not develop if humans or dogs are castrated before puberty (J. D. Wilson & C. Roehrborn, *J Clin Endocrinol Metab* 84, 4324-4331 (1999); G. Wilding, *Cancer Surv* 14, 113-130 (1992)). Castration of adult males causes involution of the prostate and apoptosis of prostatic epithelium while eliciting no effect on other male external genitalia (E. M. Bruckheimer & N. Kyprianou, *Cell Tissue Res* 301, 153-162 (2000); J. T. Isaacs, *Prostate* 5, 545-557 (1984)). This dependency on androgens provides the underlying rationale for treating prostate cancer with chemical or surgical castration (androgen ablation).

Androgens also play a role in female diseases such as polycystic ovary syndrome as well as cancers. One example is ovarian cancer where elevated levels of androgens are associated with an increased risk of developing ovarian cancer (K. J. Helzlsouer, A. J. Alberg, G. B. Gordon, C. Longcope, T. L. Bush, S. C. Hoffman & G. W. Comstock, *JAMA* 274, 1926-1930 (1995); R. J. Edmondson, J. M. Monaghan & B. R. Davies, *Br J Cancer*

86, 879-885 (2002)). The AR has been detected in a majority of ovarian cancers (H. A. Risch, *J Natl Cancer Inst* 90, 1774-1786 (1998); B. R. Rao & B. J. Slotman, *Endocr Rev* 12, 14-26 (1991); G. M. Clinton & W. Hua, *Crit Rev Oncol Hematol* 25, 1-9 (1997)), whereas estrogen receptor-alpha (ERa) and the progesterone receptor are detected in less
5 than 50% of ovarian tumors.

The only effective treatment available for advanced prostate cancer is the withdrawal of androgens which are essential for the survival of prostate epithelial cells. Androgen ablation therapy causes a temporary reduction in tumor burden concomitant with a decrease in serum prostate-specific antigen (PSA). Unfortunately prostate cancer can
10 eventually grow again in the absence of testicular androgens (castration-resistant disease) (Huber *et al* 1987 *Scand J. Urol Nephrol.* 104, 33-39). Castration-resistant prostate cancer is biochemically characterized before the onset of symptoms by a rising titre of serum PSA (Miller *et al* 1992 *J. Urol.* 147, 956-961). Once the disease becomes castration-resistant most patients succumb to their disease within two years.

15 The AR has distinct functional domains that include the carboxy-terminal ligand-binding domain (LBD), a DNA-binding domain (DBD) comprising two zinc finger motifs, and an N-terminus domain (NTD) that contains one or more transcriptional activation domains. Binding of androgen (ligand) to the LBD of the AR results in its activation such that the receptor can effectively bind to its specific DNA consensus site,
20 termed the androgen response element (ARE), on the promoter and enhancer regions of "normally" androgen regulated genes, such as PSA, to initiate transcription. The AR can be activated in the absence of androgen by stimulation of the cAMP-dependent protein kinase (PKA) pathway, with interleukin-6 (IL-6) and by various growth factors (Culig *et al* 1994 *Cancer Res.* 54, 5474-5478; Nazareth *et al* 1996 *J. Biol. Chem.* 271, 19900-19907; Sadar
25 1999 *J. Biol. Chem.* 274, 7777-7783; Ueda *et al* 2002 A *J. Biol. Chem.* 277, 7076-7085; and Ueda *et al* 2002 B *J. Biol. Chem.* 277, 38087-38094). The mechanism of ligand-independent transformation of the AR has been shown to involve: 1) increased nuclear AR protein suggesting nuclear translocation; 2) increased AR/ARE complex formation; and 3) the AR-NTD (Sadar 1999 *J. Biol. Chem.* 274, 7777-7783; Ueda *et al*
30 2002 A *J. Biol. Chem.* 277, 7076-7085; and Ueda *et al* 2002 B *J. Biol. Chem.* 277,

38087-38094). The AR can be activated in the absence of testicular androgens by alternative signal transduction pathways in castration-resistant disease, which is consistent with the finding that nuclear AR protein is present in secondary prostate cancer tumors (Kim *et al* 2002 *Am. J. Pathol.* 160, 219-226; and van der Kwast *et al* 1991 *Inter. J. Cancer* 5 48, 189-193).

Available inhibitors of the AR include nonsteroidal antiandrogens such as bicalutamide (Casodex™), nilutamide, flutamide, enzlutamide and investigational drug ARN-509 and steroidal antiandrogens, such as cyproterone acetate. These antiandrogens target the LBD of the AR and predominantly fail presumably due to poor affinity and mutations that lead to activation of the AR by these same antiandrogens (Taplin, M.E., Bubley, G.J., Kom Y.J., Small E.J., Uptonm M., Rajeshkumarm B., Balkm S.P., *Cancer Res.*, 59, 2511-2515 (1999)). These antiandrogens would also have no effect on the recently discovered AR splice variants that lack the ligand-binding domain (LBD) to result in a constitutively active receptor which promotes progression of castration recurrent prostate cancer (Dehm SM, Schmidt LJ, Heemers HV, Vessella RL, Tindall DJ., *Cancer Res* 68, 5469-77, 2008; Guo Z, Yang X, Sun F, Jiang R, Linn DE, Chen H, Chen H, Kong X, Melamed J, Tepper CG, Kung HJ, Brodie AM, Edwards J, Qiu Y., *Cancer Res.* 69, 2305-13, 2009; Hu *et al* 2009 *Cancer Res.* 69, 16-22; Sun *et al* 2010 *J Clin Invest.* 2010 120, 2715-30).

Conventional therapy has concentrated on androgen-dependent activation of the AR through its C-terminal domain. Studies developing antagonists to the AR have concentrated on the C-terminus and specifically: 1) the allosteric pocket and AF-2 activity (Estébanez-Perpiñá *et al* 2007, *PNAS* 104, 16074-16079); 2) *in silico* "drug repurposing" procedure for identification of nonsteroidal antagonists (Bisson *et al* 2007, *PNAS* 104, 11927 – 11932); and coactivator or corepressor interactions (Chang *et al* 2005, *Mol Endocrinology* 19, 2478-2490; Hur *et al* 2004, *PLoS Biol* 2, E274; Estébanez-Perpiñá *et al* 2005, *JBC* 280, 8060-8068; He *et al* 2004, *Mol Cell* 16, 425-438).

The AR-NTD is also a target for drug development (e.g. WO 2000/001813), since the NTD contains Activation-Function-1 (AF-1) which is the essential region required for AR transcriptional activity (Jenster *et al* 1991. *Mol Endocrinol.* 5, 1396-404). The AR-

NTD importantly plays a role in activation of the AR in the absence of androgens (Sadar, M.D. 1999 *J. Biol. Chem.* 274, 7777-7783; Sadar MD *et al* 1999 *Endocr Relat Cancer.* 6, 487-502; Ueda *et al* 2002 *J. Biol. Chem.* 277, 7076-7085; Ueda 2002 *J. Biol. Chem.* 277, 38087-38094; Blaszczyk *et al* 2004 *Clin Cancer Res.* 10, 1860-9; Dehm *et al* 2006 *J Biol Chem.* 28, 27882-93; Gregory *et al* 2004 *J Biol Chem.* 279, 7119-30). The AR-NTD is important in hormonal progression of prostate cancer as shown by application of decoy molecules (Quayle *et al* 2007, *Proc Natl Acad Sci U S A.* 104,1331-1336).

While the crystal structure has been resolved for the AR C-terminus LBD, this has not been the case for the NTD due to its high flexibility and intrinsic disorder in solution (Reid *et al* 2002 *J. Biol. Chem.* 277, 20079-20086) thereby hampering virtual docking drug discovery approaches. Compounds that modulate AR include the bis-phenol compounds disclosed in published PCT Nos: WO 2010/000066, WO 2011/082487; WO 2011/082488; WO 2012/145330; WO 2012/139039; WO 2012/145328; WO 2013/028572; WO 2013/028791; and WO 2014/179867, which are hereby incorporated by reference in their entireties, to the British Columbia Cancer Agency Branch and The University of British Columbia.

In addition to compounds which modulate AR, compounds and methods for imaging the prostate are useful research, diagnostic and prognostic tools. Such compounds are useful in many applications, including imaging of benign and/or malignant prostate cells and tissue. In this regard, positron emission tomography (PET) is an often used imaging technique for non-invasive identification of pathological state and tumors. In PET imaging, the distribution of a radioisotope (*e.g.*, ^{18}F) in the body can be determined. Thus incorporating ^{18}F into compounds which concentrate in tumor sites (see *e.g.*, WO 2013/028791) offers potential for diagnosis, staging, and monitoring treatment of cancers. However, improved methods for imaging are needed, for example methods which employ ^{123}I and single photon emission coupled tomography (SPECT) techniques have potential to improve methods for imaging AR-rich tissues such as the benign prostate, and in particular prostate cancers and AR splice variants in castrate recurrent prostate cancers.

While significant advances have been made in this field, there remains a need for improved imaging agents. In particular, methods and compounds suitable for imaging

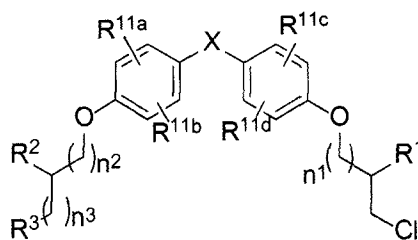
benign and malignant prostate tissues and cells are needed. The present invention fulfills these needs, and provides other related advantages.

BRIEF SUMMARY

Some embodiments of the compounds described herein can be used for diagnostic purposes to investigate diseases of the prostate, including cancer. In particular embodiments, the compounds are useful for imaging diagnostics in cancer. In some embodiments, such imaging allows for the detection and/or location of cancer sites (e.g., tumor sites). Furthermore, these compounds can be used individually or as part of a kit for such purposes.

The present disclosure is based in part on the surprising discovery that the compounds described herein, can be used to modulate AR activity either *in vivo* or *in vitro* for both research and therapeutic uses. Accordingly, embodiments of the compounds are useful for imaging the prostate. The imaging can be for any number of diagnostic purposes. For example, in certain embodiments the compounds are useful for imaging benign prostate cancer diseases. In other embodiments, the compounds find utility for imaging of certain cancers, including prostate cancer since certain embodiments of the compounds localize in prostate tumor sites. Other imaging agents are androgen mimics; however, in one embodiment, the compounds are useful for imaging AR splice variants or any AR species (i.e., those mutated in other domains or regions). The AR can be mammalian. For example, the AR can be human. The prostate cancer can be castration-resistant prostate cancer. The prostate cancer can be androgen-dependent prostate cancer.

In accordance with one embodiment, there is provided a compound having a structure of Formula I:



(I)

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein R^1 , R^2 , R^3 , R^{11a} , R^{11b} , R^{11c} , R^{11d} , X , n^1 , n^2 and n^3 are as defined herein. In one embodiment, the compound of Formula (I) comprising at least one F, Cl, Br, I, ^{18}F or ^{123}I moiety, are provided.

5 In other embodiments pharmaceutical compositions comprising a compound of Formula (I) are provided. Methods employing such pharmaceutical compositions for imaging cancer are also provided. Methods for modulating AR activity employing the present compounds and pharmaceutical compositions are also provided.

10 These and other aspects of the invention will be apparent upon reference to the following detailed description. To this end, various references are set forth herein which describe in more detail certain background information, procedures, compounds and/or compositions, and are each hereby incorporated by reference in their entirety.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 is a ^1H NMR spectrum of Compound **i-H**.
15 **Figure 2** is a ^{13}C NMR spectrum of Compound **i-H**.
Figure 3 is a ^1H NMR spectrum of Compound **ii-H**.
Figure 4 is a ^{13}C NMR spectrum of Compound **ii-H**.
Figure 5 is a ^1H NMR spectrum of Compound **1a**.
Figure 6 is a ^{13}C NMR spectrum of Compound **1a**.
20 **Figure 7** is a ^1H NMR spectrum of Compound **25a**.
Figure 8 is a ^{13}C NMR spectrum of Compound **25a**.
Figure 9 is a ^1H NMR spectrum of Compound **i-I**.
Figure 10 is a ^{13}C NMR spectrum of Compound **i-I**.
Figure 11 is a ^1H NMR spectrum of Compound **ii-I**.
25 **Figure 12** is a ^{13}C NMR spectrum of Compound **ii-I**.
Figure 13 is a ^1H NMR spectrum of Compound **4a**.
Figure 14 is a ^{13}C NMR spectrum of Compound **5a**.
Figure 15 is a ^1H NMR spectrum of Compound **28a**.
Figure 16 is a ^{13}C NMR spectrum of Compound **28a**.

Figure 17 is a ^1H NMR spectrum of Compound 13.

DETAILED DESCRIPTION

I. Definitions

In the following description, certain specific details are set forth in order to provide
5 a thorough understanding of various embodiments. However, one skilled in the art will
understand that the invention can be practiced without these details. In other instances,
well-known structures have not been shown or described in detail to avoid unnecessarily
obscuring descriptions of the embodiments. Unless the context requires otherwise,
throughout the specification and claims which follow, the word “comprise” and variations
10 thereof, such as, “comprises” and “comprising” are to be construed in an open, inclusive
sense, that is, as “including, but not limited to.” Further, headings provided herein are for
convenience only and do not interpret the scope or meaning of the claimed invention.

Reference throughout this specification to “one embodiment” or “an embodiment”
means that a particular feature, structure or characteristic described in connection with the
15 embodiment is included in at least one embodiment. Thus, the appearances of the phrases
“in one embodiment” or “in an embodiment” in various places throughout this specification
are not necessarily all referring to the same embodiment. Furthermore, the particular
features, structures, or characteristics can be combined in any suitable manner in one or
more embodiments. Also, as used in this specification and the appended claims, the
20 singular forms “a,” “an,” and “the” include plural referents unless the content clearly
dictates otherwise. It should also be noted that the term “or” is generally employed in its
sense including “and/or” unless the content clearly dictates otherwise.

The terms below, as used herein, have the following meanings, unless indicated
otherwise:

- 25 “Amino” refers to the $-\text{NH}_2$ radical.
“Cyano” refers to the $-\text{CN}$ radical.
“Halo” or “halogen” refers to bromo, chloro, fluoro or iodo radical.
“Hydroxy” or “hydroxyl” refers to the $-\text{OH}$ radical.
“Imino” refers to the $=\text{NH}$ substituent.

“Nitro” refers to the -NO₂ radical.

“Oxo” refers to the =O substituent.

“Thioxo” refers to the =S substituent.

“Alkyl” or “alkyl group” refers to a fully saturated, straight or branched hydrocarbon chain radical having from one to twelve carbon atoms, and which is attached to the rest of the molecule by a single bond. Alkyls comprising any number of carbon atoms from 1 to 12 are included. An alkyl comprising up to 12 carbon atoms is a C₁-C₁₂ alkyl, an alkyl comprising up to 10 carbon atoms is a C₁-C₁₀ alkyl, an alkyl comprising up to 6 carbon atoms is a C₁-C₆ alkyl and an alkyl comprising up to 5 carbon atoms is a C₁-C₅ alkyl. A C₁-C₅ alkyl includes C₅ alkyls, C₄ alkyls, C₃ alkyls, C₂ alkyls and C₁ alkyl (*i.e.*, methyl). A C₁-C₆ alkyl includes all moieties described above for C₁-C₅ alkyls but also includes C₆ alkyls. A C₁-C₁₀ alkyl includes all moieties described above for C₁-C₅ alkyls and C₁-C₆ alkyls, but also includes C₇, C₈, C₉ and C₁₀ alkyls. Similarly, a C₁-C₁₂ alkyl includes all the foregoing moieties, but also includes C₁₁ and C₁₂ alkyls. Non-limiting examples of C₁-C₁₂ alkyl include methyl, ethyl, *n*-propyl, *i*-propyl, *sec*-propyl, *n*-butyl, *i*-butyl, *sec*-butyl, *t*-butyl, *n*-pentyl, *t*-amyl, *n*-hexyl, *n*-heptyl, *n*-octyl, *n*-nonyl, *n*-decyl, *n*-undecyl, and *n*-dodecyl. Unless stated otherwise specifically in the specification, an alkyl group can be optionally substituted.

“Alkylene” or “alkylene chain” refers to a fully saturated, straight or branched divalent hydrocarbon chain radical, and having from one to twelve carbon atoms. Non-limiting examples of C₁-C₁₂ alkylene include methylene, ethylene, propylene, *n*-butylene, ethenylene, propenylene, *n*-butenylene, propynylene, *n*-butynylene, and the like. The alkylene chain is attached to the rest of the molecule through a single bond and to the radical group through a single bond. The points of attachment of the alkylene chain to the rest of the molecule and to the radical group can be through one carbon or any two carbons within the chain. Unless stated otherwise specifically in the specification, an alkylene chain can be optionally substituted.

“Alkenyl” or “alkenyl group” refers to a straight or branched hydrocarbon chain radical having from two to twelve carbon atoms, and having one or more carbon-carbon double bonds. Each alkenyl group is attached to the rest of the molecule by a single bond.

Alkenyl group comprising any number of carbon atoms from 2 to 12 are included. An alkenyl group comprising up to 12 carbon atoms is a C₂-C₁₂ alkenyl, an alkenyl comprising up to 10 carbon atoms is a C₂-C₁₀ alkenyl, an alkenyl group comprising up to 6 carbon atoms is a C₂-C₆ alkenyl and an alkenyl comprising up to 5 carbon atoms is a C₂-C₅ alkenyl. A C₂-C₅ alkenyl includes C₅ alkenyls, C₄ alkenyls, C₃ alkenyls, and C₂ alkenyls. A C₂-C₆ alkenyl includes all moieties described above for C₂-C₅ alkenyls but also includes C₆ alkenyls. A C₂-C₁₀ alkenyl includes all moieties described above for C₂-C₅ alkenyls and C₂-C₆ alkenyls, but also includes C₇, C₈, C₉ and C₁₀ alkenyls. Similarly, a C₂-C₁₂ alkenyl includes all the foregoing moieties, but also includes C₁₁ and C₁₂ alkenyls. Non-limiting examples of C₂-C₁₂ alkenyl include ethenyl (vinyl), 1-propenyl, 2-propenyl (allyl), isopropenyl, 2-methyl-1-propenyl, 1-butenyl, 2-butenyl, 3-butenyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 1-heptenyl, 2-heptenyl, 3-heptenyl, 4-heptenyl, 5-heptenyl, 6-heptenyl, 1-octenyl, 2-octenyl, 3-octenyl, 4-octenyl, 5-octenyl, 6-octenyl, 7-octenyl, 1-nonenyl, 2-nonenyl, 3-nonenyl, 4-nonenyl, 5-nonenyl, 6-nonenyl, 7-nonenyl, 8-nonenyl, 1-decenyl, 2-decenyl, 3-decenyl, 4-decenyl, 5-decenyl, 6-decenyl, 7-decenyl, 8-decenyl, 9-decenyl, 1-undecenyl, 2-undecenyl, 3-undecenyl, 4-undecenyl, 5-undecenyl, 6-undecenyl, 7-undecenyl, 8-undecenyl, 9-undecenyl, 10-undecenyl, 1-dodecenyl, 2-dodecenyl, 3-dodecenyl, 4-dodecenyl, 5-dodecenyl, 6-dodecenyl, 7-dodecenyl, 8-dodecenyl, 9-dodecenyl, 10-dodecenyl, and 11-dodecenyl. Unless stated otherwise specifically in the specification, an alkyl group can be optionally substituted.

“Alkenylene” or “alkenylene chain” refers to a straight or branched divalent hydrocarbon chain radical, having from two to twelve carbon atoms, and having one or more carbon-carbon double bonds. Non-limiting examples of C₂-C₁₂ alkenylene include ethene, propene, butene, and the like. The alkenylene chain is attached to the rest of the molecule through a single bond and to the radical group through a single bond. The points of attachment of the alkenylene chain to the rest of the molecule and to the radical group can be through one carbon or any two carbons within the chain. Unless stated otherwise specifically in the specification, an alkenylene chain can be optionally substituted.

“Alkynyl” or “alkynyl group” refers to a straight or branched hydrocarbon chain radical having from two to twelve carbon atoms, and having one or more carbon-carbon triple bonds. Each alkynyl group is attached to the rest of the molecule by a single bond. Alkynyl group comprising any number of carbon atoms from 2 to 12 are included. An
5 alkynyl group comprising up to 12 carbon atoms is a C₂-C₁₂ alkynyl, an alkynyl comprising up to 10 carbon atoms is a C₂-C₁₀ alkynyl, an alkynyl group comprising up to 6 carbon atoms is a C₂-C₆ alkynyl and an alkynyl comprising up to 5 carbon atoms is a C₂-C₅ alkynyl. A C₂-C₅ alkynyl includes C₅ alkynyls, C₄ alkynyls, C₃ alkynyls, and C₂ alkynyls. A C₂-C₆ alkynyl includes all moieties described above for C₂-C₅ alkynyls but also includes
10 C₆ alkynyls. A C₂-C₁₀ alkynyl includes all moieties described above for C₂-C₅ alkynyls and C₂-C₆ alkynyls, but also includes C₇, C₈, C₉ and C₁₀ alkynyls. Similarly, a C₂-C₁₂ alkynyl includes all the foregoing moieties, but also includes C₁₁ and C₁₂ alkynyls. Non-limiting examples of C₂-C₁₂ alkenyl include ethynyl, propynyl, butynyl, pentynyl and the like. Unless stated otherwise specifically in the specification, an alkyl group can be optionally
15 substituted.

“Alkynylene” or “alkynylene chain” refers to a straight or branched divalent hydrocarbon chain radical, having from two to twelve carbon atoms, and having one or more carbon-carbon triple bonds. Non-limiting examples of C₂-C₁₂ alkynylene include ethynylene, propargylene and the like. The alkynylene chain is attached to the rest of the
20 molecule through a single bond and to the radical group through a single bond. The points of attachment of the alkynylene chain to the rest of the molecule and to the radical group can be through one carbon or any two carbons within the chain. Unless stated otherwise specifically in the specification, an alkynylene chain can be optionally substituted.

“Alkoxy” refers to a radical of the formula -OR_a where R_a is an alkyl, alkenyl or
25 alkynyl radical as defined above containing one to twelve carbon atoms. Unless stated otherwise specifically in the specification, an alkoxy group can be optionally substituted.

“Alkylamino” refers to a radical of the formula -NHR_a or -NR_aR_a where each R_a is, independently, an alkyl, alkenyl or alkynyl radical as defined above containing one to
30 twelve carbon atoms. Unless stated otherwise specifically in the specification, an alkylamino group can be optionally substituted.

“Alkylcarbonyl” refers to the $-C(=O)R_a$ moiety, wherein R_a is an alkyl, alkenyl or alkynyl radical as defined above. A non-limiting example of an alkyl carbonyl is the methyl carbonyl (“acetal”) moiety. Alkylcarbonyl groups can also be referred to as “C_w-C_z acyl” where w and z depicts the range of the number of carbon in R_a , as defined above. For example, “C₁-C₁₀ acyl” refers to alkylcarbonyl group as defined above, where R_a is C₁-C₁₀ alkyl, C₁-C₁₀ alkenyl, or C₁-C₁₀ alkynyl radical as defined above. Unless stated otherwise specifically in the specification, an alkyl carbonyl group can be optionally substituted.

“Aryl” refers to a hydrocarbon ring system radical comprising hydrogen, 6 to 18 carbon atoms and at least one aromatic ring. For purposes of this invention, the aryl radical can be a monocyclic, bicyclic, tricyclic or tetracyclic ring system, which can include fused or bridged ring systems. Aryl radicals include, but are not limited to, aryl radicals derived from aceanthrylene, acenaphthylene, acephenanthrylene, anthracene, azulene, benzene, chrysene, fluoranthene, fluorene, *as*-indacene, *s*-indacene, indane, indene, naphthalene, phenalene, phenanthrene, pleiadene, pyrene, and triphenylene. Unless stated otherwise specifically in the specification, the term “aryl” is meant to include aryl radicals that are optionally substituted.

“Aralkyl” refers to a radical of the formula $-R_b-R_c$ where R_b is an alkylene, alkenylene or alkynylene group as defined above and R_c is one or more aryl radicals as defined above, for example, benzyl, diphenylmethyl and the like. Unless stated otherwise specifically in the specification, an aralkyl group can be optionally substituted.

“Carbocyclyl,” “carbocyclic ring” or “carbocycle” refers to a rings structure, wherein the atoms which form the ring are each carbon. Carbocyclic rings can comprise from 3 to 20 carbon atoms in the ring. Carbocyclic rings include aryls and cycloalkyl, cycloalkenyl and cycloalkynyl as defined herein. Unless stated otherwise specifically in the specification, a carbocyclyl group can be optionally substituted.

“Cycloalkyl” refers to a stable non-aromatic monocyclic or polycyclic fully saturated hydrocarbon radical consisting solely of carbon and hydrogen atoms, which can include fused or bridged ring systems, having from three to twenty carbon atoms, preferably having from three to ten carbon atoms, and which is attached to the rest of the molecule by a single bond. Monocyclic cycloalkyl radicals include, for example,

cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, and cyclooctyl. Polycyclic cycloalkyl radicals include, for example, adamantyl, norbornyl, decalanyl, 7,7-dimethyl-bicyclo[2.2.1]heptanyl, and the like. Unless otherwise stated specifically in the specification, a cycloalkyl group can be optionally substituted.

5 “Cycloalkenyl” refers to a stable non-aromatic monocyclic or polycyclic hydrocarbon radical consisting solely of carbon and hydrogen atoms, having one or more carbon-carbon double bonds, which can include fused or bridged ring systems, having from three to twenty carbon atoms, preferably having from three to ten carbon atoms, and which is attached to the rest of the molecule by a single bond. Monocyclic cycloalkenyl radicals
10 include, for example, cyclopentenyl, cyclohexenyl, cycloheptenyl, cyclooctenyl, and the like. Polycyclic cycloalkenyl radicals include, for example, bicyclo[2.2.1]hept-2-enyl and the like. Unless otherwise stated specifically in the specification, a cycloalkenyl group can be optionally substituted.

 “Cycloalkynyl” refers to a stable non-aromatic monocyclic or polycyclic
15 hydrocarbon radical consisting solely of carbon and hydrogen atoms, having one or more carbon-carbon triple bonds, which can include fused or bridged ring systems, having from three to twenty carbon atoms, preferably having from three to ten carbon atoms, and which is attached to the rest of the molecule by a single bond. Monocyclic cycloalkynyl radicals include, for example, cycloheptynyl, cyclooctyny, and the like. Unless otherwise stated
20 specifically in the specification, a cycloalkynyl group can be optionally substituted.

 “Cycloalkylalkyl” refers to a radical of the formula $-R_b-R_d$ where R_b is an alkylene, alkenylene, or alkynylene group as defined above and R_d is a cycloalkyl, cycloalkenyl, cycloalkynyl radical as defined above. Unless stated otherwise specifically in the specification, a cycloalkylalkyl group can be optionally substituted.

25 “Haloalkyl” refers to an alkyl radical, as defined above, that is substituted by one or more halo radicals, as defined above, *e.g.*, trifluoromethyl, difluoromethyl, trichloromethyl, 2,2,2-trifluoroethyl, 1,2-difluoroethyl, 3-bromo-2-fluoropropyl, 1,2-dibromoethyl, and the like. Unless stated otherwise specifically in the specification, a haloalkyl group can be optionally substituted.

“Haloalkenyl” refers to an alkenyl radical, as defined above, that is substituted by one or more halo radicals, as defined above, *e.g.*, 1-fluoropropenyl, 1,1-difluorobutenyl, and the like. Unless stated otherwise specifically in the specification, a haloalkenyl group can be optionally substituted.

5 “Haloalkynyl” refers to an alkynyl radical, as defined above, that is substituted by one or more halo radicals, as defined above, *e.g.*, 1-fluoropropynyl, 1-fluorobutynyl, and the like. Unless stated otherwise specifically in the specification, a haloalkenyl group can be optionally substituted.

“Heterocyclyl,” “heterocyclic ring” or “heterocycle” refers to a stable 3- to
10 20-membered non-aromatic ring radical which consists of two to twelve carbon atoms and from one to six heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur. Heterocyclyl or heterocyclic rings include heteroaryls as defined below. Unless stated otherwise specifically in the specification, the heterocyclyl radical can be a monocyclic, bicyclic, tricyclic or tetracyclic ring system, which can include fused or
15 bridged ring systems; and the nitrogen, carbon or sulfur atoms in the heterocyclyl radical can be optionally oxidized; the nitrogen atom can be optionally quaternized; and the heterocyclyl radical can be partially or fully saturated. Examples of such heterocyclyl radicals include, but are not limited to, dioxolanyl, thienyl[1,3]dithianyl, decahydroisoquinolyl, imidazolyl, imidazolidinyl, isothiazolidinyl, isoxazolidinyl,
20 morpholyl, octahydroindolyl, octahydroisoindolyl, 2-oxopiperazinyl, 2-oxopiperidinyl, 2-oxopyrrolidinyl, oxazolidinyl, piperidinyl, piperazinyl, 4-piperidonyl, pyrrolidinyl, pyrazolidinyl, quinuclidinyl, thiazolidinyl, tetrahydrofuryl, trithianyl, tetrahydropyranlyl, thiomorpholyl, thiamorpholyl, 1-oxo-thiomorpholyl, and 1,1-dioxo-thiomorpholyl. Unless stated otherwise specifically in the specification, a heterocyclyl group can be
25 optionally substituted.

“*N*-heterocyclyl” refers to a heterocyclyl radical as defined above containing at least one nitrogen and where the point of attachment of the heterocyclyl radical to the rest of the molecule is through a nitrogen atom in the heterocyclyl radical. Unless stated otherwise specifically in the specification, a *N*-heterocyclyl group can be optionally
30 substituted.

“Heterocyclalkyl” refers to a radical of the formula $-R_b-R_e$ where R_b is an alkylene, alkenylene, or alkynylene chain as defined above and R_e is a heterocycl radical as defined above, and if the heterocycl is a nitrogen-containing heterocycl, the heterocycl can be attached to the alkyl, alkenyl, alkynyl radical at the nitrogen atom.

- 5 Unless stated otherwise specifically in the specification, a heterocyclalkyl group can be optionally substituted.

“Heteroaryl” refers to a 5- to 20-membered ring system radical comprising hydrogen atoms, one to thirteen carbon atoms, one to six heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur, and at least one aromatic ring. For
 10 purposes of this invention, the heteroaryl radical can be a monocyclic, bicyclic, tricyclic or tetracyclic ring system, which can include fused or bridged ring systems; and the nitrogen, carbon or sulfur atoms in the heteroaryl radical can be optionally oxidized; the nitrogen atom can be optionally quaternized. Examples include, but are not limited to, azepinyl, acridinyl, benzimidazolyl, benzothiazolyl, benzindolyl, benzodioxolyl, benzofuranlyl,
 15 benzooxazolyl, benzothiazolyl, benzothiadiazolyl, benzo[*b*][1,4]dioxepinyl, 1,4-benzodioxanyl, benzonaphthofuranlyl, benzoxazolyl, benzodioxolyl, benzodioxinyl, benzopyranlyl, benzopyranonyl, benzofuranlyl, benzofuranonyl, benzothieryl (benzothiophenyl), benzotriazolyl, benzo[4,6]imidazo[1,2-*a*]pyridinyl, carbazolyl, cinnolinyl, dibenzofuranlyl, dibenzothiophenyl, furanyl, furanonyl, isothiazolyl,
 20 imidazolyl, indazolyl, indolyl, indazolyl, isoindolyl, indolinyl, isoindolinyl, isoquinolyl, indoliziny, isoxazolyl, naphthyridinyl, oxadiazolyl, 2-oxoazepinyl, oxazolyl, oxiranyl, 1-oxidopyridinyl, 1-oxidopyrimidinyl, 1-oxidopyrazinyl, 1-oxidopyridazinyl, 1-phenyl-1*H*-pyrrolyl, phenazinyl, phenothiazinyl, phenoxazinyl, phthalazinyl, pteridinyl, purinyl, pyrrolyl, pyrazolyl, pyridinyl, pyrazinyl, pyrimidinyl, pyridazinyl, quinazolinyl,
 25 quinoxalinyl, quinolinyl, quinuclidinyl, isoquinolinyl, tetrahydroquinolinyl, thiazolyl, thiadiazolyl, triazolyl, tetrazolyl, triazinyl, and thiophenyl (i.e. thienyl). Unless stated otherwise specifically in the specification, a heteroaryl group can be optionally substituted.

“*N*-heteroaryl” refers to a heteroaryl radical as defined above containing at least one nitrogen and where the point of attachment of the heteroaryl radical to the rest of the

molecule is through a nitrogen atom in the heteroaryl radical. Unless stated otherwise specifically in the specification, an *N*-heteroaryl group can be optionally substituted.

“Heteroarylalkyl” refers to a radical of the formula $-R_b-R_f$ where R_b is an alkylene, alkenylene, or alkynylene chain as defined above and R_f is a heteroaryl radical as defined
5 above. Unless stated otherwise specifically in the specification, a heteroarylalkyl group can be optionally substituted.


“ ^{123}I ” refers to the radioactive isotope of iodine having atomic mass 123. The compounds of Formula I can comprise at least one ^{123}I moiety. Throughout the present application, where structures depict a ^{123}I moiety at a certain position it is meant that the I
10 moiety at this position is enriched for ^{123}I . In other words, the compounds contain more than the natural abundance of ^{123}I at the indicated position(s). It is not required that the compounds comprise 100% ^{123}I at the indicated positions, provided ^{123}I is present in more than the natural abundance. Typically the ^{123}I isotope is enriched to greater than 50%, greater than 60%, greater than 70%, greater than 80% or greater than 90%, relative to ^{127}I .



“ ^{18}F ” refers to the radioactive isotope of fluorine having atomic mass 18. “ F ” or “ ^{19}F ” refers to the abundant, non-radioactive fluorine isotope having atomic mass 19. The compounds of Formula I can comprise at least one ^{18}F moiety. Throughout the present application, where structures depict a ^{18}F moiety at a certain position it is meant that the F
15 moiety at this position is enriched for ^{18}F . In other words, the compounds contain more than the natural abundance of ^{18}F at the indicated position(s). It is not required that the compounds comprise 100% ^{18}F at the indicated positions, provided ^{18}F is present in more than the natural abundance. Typically the ^{18}F isotope is enriched to greater than 50%, greater than 60%, greater than 70%, greater than 80% or greater than 90%, relative to ^{19}F .
20

“Thioalkyl” refers to a radical of the formula $-\text{SR}_a$ where R_a is an alkyl, alkenyl, or alkynyl radical as defined above containing one to twelve carbon atoms. Unless stated
25 otherwise specifically in the specification, a thioalkyl group can be optionally substituted.

The term “substituted” used herein means any of the above groups (*i.e.*, alkyl, alkylene, alkenyl, alkenylene, alkynyl, alkynylene, alkoxy, alkylamino, alkylcarbonyl, thioalkyl, aryl, aralkyl, carbocyclyl, cycloalkyl, cycloalkenyl, cycloalkynyl,
30 cycloalkylalkyl, haloalkyl, heterocyclyl, *N*-heterocyclyl, heterocyclylalkyl, heteroaryl, *N*-

heteroaryl and/or heteroarylalkyl) wherein at least one hydrogen atom is replaced by a bond to a non-hydrogen atoms such as, but not limited to: a halogen atom such as F, Cl, Br, and I; an oxygen atom in groups such as hydroxyl groups, alkoxy groups, and ester groups; a sulfur atom in groups such as thiol groups, thioalkyl groups, sulfone groups, sulfonyl groups, and sulfoxide groups; a nitrogen atom in groups such as amines, amides, alkylamines, dialkylamines, arylamines, alkylarylamines, diarylamines, N-oxides, imides, and enamines; a silicon atom in groups such as trialkylsilyl groups, dialkylarylsilyl groups, alkyl-diarylsilyl groups, and triarylsilyl groups; and other heteroatoms in various other groups. "Substituted" also means any of the above groups in which one or more hydrogen atoms are replaced by a higher-order bond (e.g., a double- or triple-bond) to a heteroatom such as oxygen in oxo, carbonyl, carboxyl, and ester groups; and nitrogen in groups such as imines, oximes, hydrazones, and nitriles. For example, "substituted" includes any of the above groups in which one or more hydrogen atoms are replaced with $-NR_gR_h$, $-NR_gC(=O)R_h$, $-NR_gC(=O)NR_gR_h$, $-NR_gC(=O)OR_h$, $-NR_gSO_2R_h$, $-OC(=O)NR_gR_h$, $-OR_g$, $-SR_g$, $-SOR_g$, $-SO_2R_g$, $-OSO_2R_g$, $-SO_2OR_g$, $=NSO_2R_g$, and $-SO_2NR_gR_h$. "Substituted" also means any of the above groups in which one or more hydrogen atoms are replaced with $-C(=O)R_g$, $-C(=O)OR_g$, $-C(=O)NR_gR_h$, $-CH_2SO_2R_g$, $-CH_2SO_2NR_gR_h$. In the foregoing, R_g and R_h are the same or different and independently hydrogen, alkyl, alkenyl, alkynyl, alkoxy, alkylamino, thioalkyl, aryl, aralkyl, cycloalkyl, cycloalkenyl, cycloalkynyl, cycloalkylalkyl, haloalkyl, haloalkenyl, haloalkynyl, heterocyclyl, *N*-heterocyclyl, heterocyclylalkyl, heteroaryl, *N*-heteroaryl and/or heteroarylalkyl. "Substituted" further means any of the above groups in which one or more hydrogen atoms are replaced by a bond to an amino, cyano, hydroxyl, imino, nitro, oxo, thioxo, halo, alkyl, alkenyl, alkynyl, alkoxy, alkylamino, thioalkyl, aryl, aralkyl, cycloalkyl, cycloalkenyl, cycloalkynyl, cycloalkylalkyl, haloalkyl, haloalkenyl, haloalkynyl, heterocyclyl, *N*-heterocyclyl, heterocyclylalkyl, heteroaryl, *N*-heteroaryl and/or heteroarylalkyl group. In addition, each of the foregoing substituents can also be optionally substituted with one or more of the above substituents.

As used herein, the symbol “” (hereinafter can be referred to as “a point of attachment bond”) denotes a bond that is a point of attachment between two chemical entities, one of which is depicted as being attached to the point of attachment bond and the other of which is not depicted as being attached to the point of attachment bond. For

5 example, “ $XY\text{---}\text{$ ” indicates that the chemical entity “XY” is bonded to another chemical entity via the point of attachment bond. Furthermore, the specific point of attachment to the non-depicted chemical entity can be specified by inference. For example, the compound $\text{CH}_3\text{---}R^3$, wherein R^3 is H or “ $XY\text{---}\text{$ ” infers that when R^3 is “XY”, the point of attachment bond is the same bond as the bond by which R^3 is depicted as being bonded to CH_3 .

10 “Fused” refers to any ring structure described herein which is fused to an existing ring structure in the compounds of the invention. When the fused ring is a heterocyclyl ring or a heteroaryl ring, any carbon atom on the existing ring structure which becomes part of the fused heterocyclyl ring or the fused heteroaryl ring can be replaced with a nitrogen atom.

15 The invention disclosed herein is also meant to encompass the *in vivo* metabolic products of the disclosed compounds. Such products can result from, for example, the oxidation, reduction, hydrolysis, amidation, esterification, and the like of the administered compound, primarily due to enzymatic processes. Accordingly, the invention includes compounds produced by a process comprising administering a compound of this invention
20 to a mammal for a period of time sufficient to yield a metabolic product thereof. Such products are typically identified by administering a radiolabelled compound of the invention in a detectable dose to an animal, such as rat, mouse, guinea pig, monkey, or to human, allowing sufficient time for metabolism to occur, and isolating its conversion products from the urine, blood or other biological samples.

25 “Stable compound” and “stable structure” are meant to indicate a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into an efficacious therapeutic agent.

As used herein, a “subject” can be a human, non-human primate, mammal, rat, mouse, cow, horse, pig, sheep, goat, dog, cat and the like. The subject can be suspected of having or at risk for having a cancer, such as prostate cancer, breast cancer, ovarian cancer, salivary gland carcinoma, or endometrial cancer, or suspected of having or at risk for having acne, hirsutism, alopecia, benign prostatic hyperplasia, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, or age-related macular degeneration. Diagnostic methods for various cancers, such as prostate cancer, breast cancer, ovarian cancer, salivary gland carcinoma, or endometrial cancer, and diagnostic methods for acne, hirsutism, alopecia, benign prostatic hyperplasia, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, or age-related macular degeneration and the clinical delineation of cancer, such as prostate cancer, breast cancer, ovarian cancer, salivary gland carcinoma, or endometrial cancer, diagnoses and the clinical delineation of acne, hirsutism, alopecia, benign prostatic hyperplasia, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, or age-related macular degeneration are known to those of ordinary skill in the art.

“Mammal” includes humans and both domestic animals such as laboratory animals and household pets (*e.g.*, cats, dogs, swine, cattle, sheep, goats, horses, rabbits), and non-domestic animals such as wildlife and the like.

“Optional” or “optionally” means that the subsequently described event of circumstances can or can not occur, and that the description includes instances where said event or circumstance occurs and instances in which it does not. For example, “optionally substituted aryl” means that the aryl radical can or can not be substituted and that the description includes both substituted aryl radicals and aryl radicals having no substitution.

“Pharmaceutically acceptable carrier, diluent or excipient” includes without limitation any adjuvant, carrier, excipient, glidant, sweetening agent, diluent, preservative, dye/colorant, flavor enhancer, surfactant, wetting agent, dispersing agent, suspending agent, stabilizer, isotonic agent, solvent, or emulsifier which has been approved by the United States Food and Drug Administration as being acceptable for use in humans or domestic animals.

“Pharmaceutically acceptable salt” includes both acid and base addition salts.

“Pharmaceutically acceptable acid addition salt” refers to those salts which retain the biological effectiveness and properties of the free bases, which are not biologically or otherwise undesirable, and which are formed with inorganic acids such as, but are not limited to, hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid and the like, and organic acids such as, but not limited to, acetic acid, 2,2-dichloroacetic acid, adipic acid, alginic acid, ascorbic acid, aspartic acid, benzenesulfonic acid, benzoic acid, 4-acetamidobenzoic acid, camphoric acid, camphor-10-sulfonic acid, capric acid, caproic acid, caprylic acid, carbonic acid, cinnamic acid, citric acid, cyclamic acid, dodecylsulfuric acid, ethane-1,2-disulfonic acid, ethanesulfonic acid, 2-hydroxyethanesulfonic acid, formic acid, fumaric acid, galactaric acid, gentisic acid, glucoheptonic acid, gluconic acid, glucuronic acid, glutamic acid, glutaric acid, 2-oxo-glutaric acid, glycerophosphoric acid, glycolic acid, hippuric acid, isobutyric acid, lactic acid, lactobionic acid, lauric acid, maleic acid, malic acid, malonic acid, mandelic acid, methanesulfonic acid, mucic acid, naphthalene-1,5-disulfonic acid, naphthalene-2-sulfonic acid, 1-hydroxy-2-naphthoic acid, nicotinic acid, oleic acid, orotic acid, oxalic acid, palmitic acid, pamoic acid, propionic acid, pyroglutamic acid, pyruvic acid, salicylic acid, 4-aminosalicylic acid, sebacic acid, stearic acid, succinic acid, tartaric acid, thiocyanic acid, *p*-toluenesulfonic acid, trifluoroacetic acid, undecylenic acid, and the like.

“Pharmaceutically acceptable base addition salt” refers to those salts which retain the biological effectiveness and properties of the free acids, which are not biologically or otherwise undesirable. These salts are prepared from addition of an inorganic base or an organic base to the free acid. Salts derived from inorganic bases include, but are not limited to, the sodium, potassium, lithium, ammonium, calcium, magnesium, iron, zinc, copper, manganese, aluminum salts and the like. Preferred inorganic salts are the ammonium, sodium, potassium, calcium, and magnesium salts. Salts derived from organic bases include, but are not limited to, salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as ammonia, isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, diethanolamine, ethanolamine, deanol,

2-dimethylaminoethanol, 2-diethylaminoethanol, dicyclohexylamine, lysine, arginine, histidine, caffeine, procaine, hydrabamine, choline, betaine, benethamine, benzathine, ethylenediamine, glucosamine, methylglucamine, theobromine, triethanolamine, tromethamine, purines, piperazine, piperidine, *N*-ethylpiperidine, polyamine resins and the like. Particularly preferred organic bases are isopropylamine, diethylamine, ethanolamine, trimethylamine, dicyclohexylamine, choline and caffeine.

Often crystallizations produce a solvate of the compound of the invention. As used herein, the term "solvate" refers to an aggregate that comprises one or more molecules of a compound of the invention with one or more molecules of solvent. The solvent can be water, in which case the solvate can be a hydrate. Alternatively, the solvent can be an organic solvent. Thus, the compounds of the present invention can exist as a hydrate, including a monohydrate, dihydrate, hemihydrate, sesquihydrate, trihydrate, tetrahydrate and the like, as well as the corresponding solvated forms. The compound of the invention can be true solvates, while in other cases, the compound of the invention can merely retain adventitious water or be a mixture of water plus some adventitious solvent.

A "pharmaceutical composition" refers to a formulation of a compound of the invention and a medium generally accepted in the art for the delivery of the biologically active compound to mammals, *e.g.*, humans. Such a medium includes all pharmaceutically acceptable carriers, diluents or excipients therefor.

"An "effective amount" refers to a therapeutically effective amount or a prophylactically effective amount. A "therapeutically effective amount" refers to an amount effective, at dosages and for periods of time necessary, to achieve the desired therapeutic result, such as reduced tumor size, increased life span or increased life expectancy. A therapeutically effective amount of a compound can vary according to factors such as the disease state, age, sex, and weight of the subject, and the ability of the compound to elicit a desired response in the subject. Dosage regimens can be adjusted to provide the optimum therapeutic response. A therapeutically effective amount is also one in which any toxic or detrimental effects of the compound 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, such as smaller tumors, increased life span, increased life expectancy or prevention of the progression of prostate cancer to a castration-resistant form. Typically, a prophylactic dose is used in subjects prior to or at an earlier stage of disease, so that a prophylactically effective amount can be less than a therapeutically effective amount.

5 “Treating” or “treatment” as used herein covers the treatment of the disease or condition of interest in a mammal, preferably a human, having the disease or condition of interest, and includes:

 (i) preventing the disease or condition from occurring in a mammal, in particular, when such mammal is predisposed to the condition but has not yet been
10 diagnosed as having it;

 (ii) inhibiting the disease or condition, i.e., arresting its development;

 (iii) relieving the disease or condition, i.e., causing regression of the disease or condition; or

 (iv) relieving the symptoms resulting from the disease or condition, i.e.,
15 relieving pain without addressing the underlying disease or condition. As used herein, the terms “disease” and “condition” can be used interchangeably or can be different in that the particular malady or condition can not have a known causative agent (so that etiology has not yet been worked out) and it is therefore not yet recognized as a disease but only as an undesirable condition or syndrome, wherein a more or less specific set of symptoms have
20 been identified by clinicians.

 The compounds of the invention, or their pharmaceutically acceptable salts can contain one or more asymmetric centers and can thus give rise to enantiomers, diastereomers, and other stereoisomeric forms that can be defined, in terms of absolute stereochemistry, as (*R*)- or (*S*)- or, as (*D*)- or (*L*)- for amino acids. The present invention is
25 meant to include all such possible isomers, as well as their racemic and optically pure forms whether or not they are specifically depicted herein. Optically active (+) and (-), (*R*)- and (*S*)-, or (*D*)- and (*L*)- isomers can be prepared using chiral synthons or chiral reagents, or resolved using conventional techniques, for example, chromatography and fractional crystallization. Conventional techniques for the preparation/isolation of individual
30 enantiomers include chiral synthesis from a suitable optically pure precursor or resolution

of the racemate (or the racemate of a salt or derivative) using, for example, chiral high pressure liquid chromatography (HPLC). When the compounds described herein contain olefinic double bonds or other centers of geometric asymmetry, and unless specified otherwise, it is intended that the compounds include both E and Z geometric isomers.

5 Likewise, all tautomeric forms are also intended to be included.

A “stereoisomer” refers to a compound made up of the same atoms bonded by the same bonds but having different three-dimensional structures, which are not interchangeable. The present invention contemplates various stereoisomers and mixtures thereof and includes “enantiomers”, which refers to two stereoisomers whose molecules are
10 nonsuperimposable mirror images of one another.

A “tautomer” refers to a proton shift from one atom of a molecule to another atom of the same molecule. The present invention includes tautomers of any said compounds.

The chemical naming protocol and structure diagrams used herein are a modified form of the I.U.P.A.C. nomenclature system, using the ACD/Name Version 9.07 software
15 program, ChemDraw Ultra Version 11.0.1 and/or ChemDraw Ultra Version 14.0 software naming program (CambridgeSoft). For complex chemical names employed herein, a substituent group is named before the group to which it attaches. For example, cyclopropylethyl comprises an ethyl backbone with cyclopropyl substituent. Except as described below, all bonds are identified in the chemical structure diagrams herein, except
20 for some carbon atoms, which are assumed to be bonded to sufficient hydrogen atoms to complete the valency.

Throughout the present specification, the terms “about” and/or “approximately” can be used in conjunction with numerical values and/or ranges. The term “about” is understood to mean those values near to a recited value. For example, “about 40 [units]”
25 can mean within $\pm 25\%$ of 40 (*e.g.*, from 30 to 50), within $\pm 20\%$, $\pm 15\%$, $\pm 10\%$, $\pm 9\%$, $\pm 8\%$, $\pm 7\%$, $\pm 6\%$, $\pm 5\%$, $\pm 4\%$, $\pm 3\%$, $\pm 2\%$, $\pm 1\%$, less than $\pm 1\%$, or any other value or range of values therein or therebelow. Furthermore, the phrases “less than about [a value]” or “greater than about [a value]” should be understood in view of the definition of the term “about” provided herein. The terms “about” and “approximately” can be used
30 interchangeably.

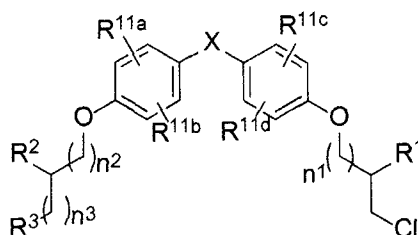
Throughout the present specification, numerical ranges are provided for certain quantities. It is to be understood that these ranges comprise all subranges therein. Thus, the range "from 50 to 80" includes all possible ranges therein (e.g., 51-79, 52-78, 53-77, 54-76, 55-75, 60-70, *etc.*). Furthermore, all values within a given range can be an endpoint
5 for the range encompassed thereby (e.g., the range 50-80 includes the ranges with endpoints such as 55-80, 50-75, *etc.*).

II. Compounds and Pharmaceutical Compositions

As noted above, certain embodiments of the present invention are directed to compounds useful for treatment of various cancers, including various types of prostate
10 cancers. While not wishing to be bound by theory, it is believed that binding of the compounds to the androgen receptor (for example at the N-terminal domain) can contribute to the activity of the disclosed compounds. The compounds of the present invention include novel heteroaryl, aryl, heterocyclyl, or carbocyclyl groups (i.e., R^3 in Formula I) which impart improved properties to the compounds compared to compounds lacking the
15 described R^3 moiety. For example, the improved properties include improved drug-like properties such as improved activity (e.g., androgen receptor (AR) modulation), longer half-life (e.g., *in vivo*); decreased toxicity; better solubility, improved formulation, better bioavailability, better pharmacokinetic profile; reduction in unwanted metabolites and the like.

20 In one embodiment the invention includes compounds which form covalent bonds with the androgen receptor (AR) (e.g., at the N-terminal domain), thus resulting in irreversible (or substantially irreversible) inhibition of the same. In this regard, the certain compounds of the present invention are designed to include functional groups capable of forming covalent bonds with a nucleophile under certain *in vivo* conditions. For example,
25 in some embodiments the reactivity of compounds of the present invention is such that they will not substantially react with various nucleophiles (e.g., glutathione) when the compounds are free in solution. However, when the free mobility of the compounds is restricted, and an appropriate nucleophile is brought into close proximity to the compound, for example when the compounds associate with, or bind to, the androgen receptor, the
30 compounds are capable of forming covalent bonds with certain nucleophiles (e.g., thiols).

The present invention includes all compounds which have the above described properties (i.e., binding to androgen receptor (AR)). In one embodiment, the present invention is directed to a compound having a structure of Formula I:



5

(I)

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-, -C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-

R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

10 R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally substituted with one or more R⁶;

R⁵ is each independently H, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

15 R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I, hydroxyl, oxo, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₆-C₁₂ aryl, wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -OS(O)₂-aryl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

20 R⁸ and R⁹ are each independently H, halogen, -S(O)₀₋₂R⁵, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, aryl, aralkyl, C₁-C₁₀ acyl, or -NR⁵R⁵, or R⁸ and R⁹ can join to form an unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

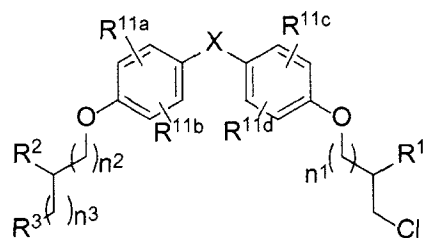
R^{11a}, R^{11b}, R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ¹²³I, hydroxyl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl; -OR⁵, -OC(=O)R¹³, C₁-C₁₀ acyl, -S(O)₀₋₂R⁵, -NO₂-, CN-, -NH₂-, -NHR⁵-, or -N(R⁵)₂;

R^{13} is each independently C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl;

n^1 and n^2 are each independently 0, 1, or 2; and

n^3 is 0, 1, 2, 3, 4, or 5.

Accordingly, certain embodiments of the present invention are directed to
 5 compounds that bind to the AR NTD and are useful for imaging of tumors with splice
 variants using SPECT and/or methods of modulating AR NTD activity. Other
 embodiments are directed to compound and methods useful for imaging and/or treating
 benign prostate conditions or diseases. In one embodiment, the present disclosure provides
 a compound having a structure of Formula I:



(I)

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-,
 -C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-;

15 R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally
 substituted with one or more R⁶;

R⁵ is each independently H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl;

R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I,
 20 hydroxyl, oxo, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_1 - C_6 alkoxy, C_6 - C_{12} aryl,
 wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -
 OS(O)₂-aryl, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl;

R^8 and R^9 are each independently H, halogen, $-S(O)_{0-2}R^5$, C_1 - C_{10} alkyl, C_2 - C_{10} alkenyl, C_2 - C_{10} alkynyl, aryl, aralkyl, C_1 - C_{10} acyl, or $-NR^5R^5$, or R^8 and R^9 can join to form a unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

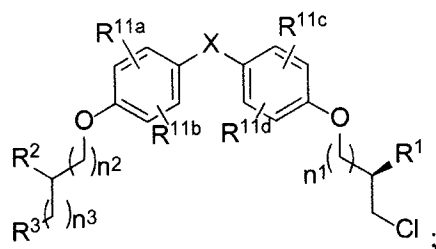
- 5 R^{11a} , R^{11b} , R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ^{123}I , hydroxyl, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl; $-OR^5$, $-OC(=O)R^{13}$, C_1 - C_{10} acyl, $-S(O)_{0-2}R^5$, $-NO_2$, $-CN$, $-NH_2$, $-NHR^5$, or $-N(R^5)_2$;

R^{13} is each independently C_1 - C_6 alkyl, C_2 - C_6 alkenyl, or C_2 - C_6 alkynyl;

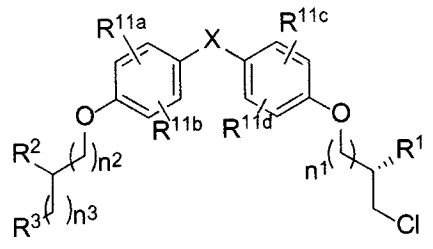
n^1 and n^2 are each independently 0, 1, or 2; and

- 10 n^3 is 0, 1, 2, 3, 4, or 5.

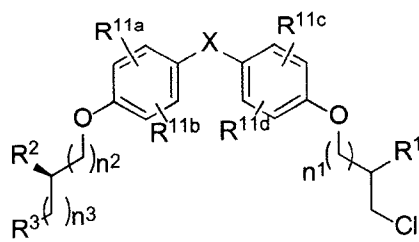
In various embodiments, different stereoisomers of the compound of structure (I) are provided, for example in some embodiments the compound has one of the following structures (Ia), (Ib), (Ic) or (Id):



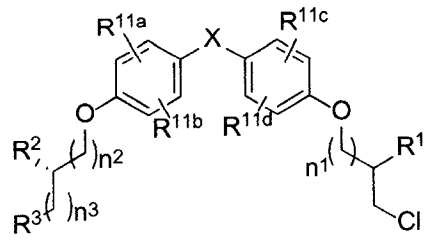
(Ia)



(Ib)

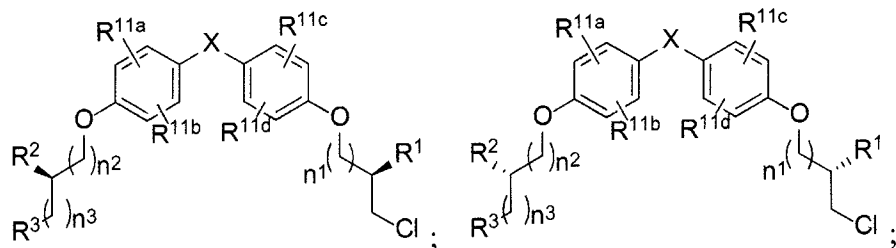


(Ic)



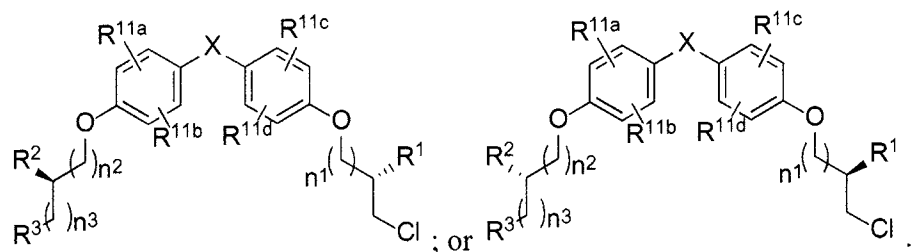
(Id)

In still other embodiments, the compound has one of the following structures (Ie), (If), (Ig) or (Ih):



(Ie)

(If)



(Ig)

(Ih)

In some embodiments, X is -O-. In other embodiments, X is -S(O)₀₋₂-. In some embodiments, X is -C(=O)-. In one embodiment, X is -C(OR⁵)₂-. In one embodiment, X is -C(OR⁵)(OC(=O)R¹³)-. In some embodiments, X is -C(R⁸R⁹)-. In some embodiments, X is -C(=CR⁸R⁹)-. In other embodiments, X is -N(R⁹)-. In one embodiment, X is -N(COR⁹)-. In one embodiment, X is -CHNR⁸R⁹-. In another embodiment, X is -C(=NR⁹)-. In some embodiments, X is -C(=NOR⁵)-. In other embodiments, X is -C(=N-NHR⁵)-.

In some embodiments, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently H, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, or C₂-C₁₀ alkynyl. In some embodiments, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently H or C₁-C₁₀ alkyl. In other embodiments, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently C₁-C₁₀ alkyl. In some embodiments, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently C₁-C₅ alkyl. In one embodiment, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently C₁ alkyl. In another embodiment, X is -C(R⁸R⁹)- wherein R⁸ and R⁹ are each independently a methyl.

In certain of the foregoing embodiments, R³ is 5-6 membered heteroaryl or 3-7 membered heterocyclyl, wherein said heteroaryl or said heterocyclyl comprises at least one N atom in the ring.

In another embodiment, R³ is selected from the group consisting of pyrrole, furan, thiophene, pyrazole, pyridine, pyridazine, pyrimidine, imidazole, thiazole, isoxazole, oxadiazole, thiadiazole, oxazole, triazole, isothiazole, oxazine, triazine, azepine, pyrrolidine, pyrroline, imidazoline, imidazolidine, pyrazoline, pyrazolidine, piperidine, dioxane, morpholine, dithiane, thiomorpholine, piperazine, and tetrazine. In one embodiment R³ is selected from the group consisting of triazole, morpholine, thiomorpholine, pyrazole, and imidazole.

In one embodiment, R³ is substituted with at least one R⁶. In some embodiments, R³ is substituted with at least one ¹²³I, I, Br, Cl, or F. In another embodiment, R³ is substituted with at least one ¹²³I or I.

In one embodiment, R³ is substituted with at least one R⁶ which is further substituted with at least one ¹²³I, I, or F. In another embodiment, R³ is substituted with at least one R⁶ which is C₁-C₆ alkyl, and said C₁-C₆ alkyl is further substituted with at least one ¹²³I, I, or F. In some embodiments, R³ is substituted with at least one R⁶ which is C₁-C₆ alkyl, and said C₁-C₆ alkyl is further substituted with at least one ¹²³I. In other embodiments, R³ is substituted with at least one R⁶ which is C₁-C₆ alkyl, and said C₁-C₆ alkyl is further substituted with at least one ¹⁸F.

In one embodiment, at least one of R⁵, R^{11a}, R^{11b}, R^{11c} and R^{11d} is I. In another embodiment, at least one of R⁵, R^{11a}, R^{11b}, R^{11c} and R^{11d} is ¹²³I. In some embodiments, at least one of R⁵, R^{11a}, R^{11b}, R^{11c} and R^{11d} is I and at least one of R⁵, R^{11a}, R^{11b}, R^{11c} and R^{11d} is ¹²³I.

In some embodiments, R¹³ is C₁-C₃ alkyl. In other embodiments, R¹³ is methyl, ethyl, or propyl. In one embodiment, R¹³ is a methyl.

In one embodiment, n³ is 0, 1, or 2.

The compounds for use in the imaging and treatment methods described herein. In some embodiments, the compounds comprise one F, Cl, Br, I, ^{18}F or ^{123}I substitution. For example in certain other embodiments, three of R^{11a} , R^{11b} , R^{11c} and R^{11d} are H, and the remaining R^{11a} , R^{11b} , R^{11c} or R^{11d} is F, Cl, Br, I or ^{123}I . In some embodiments, the compounds comprise two F, Cl, Br, I or ^{123}I substitutions on the phenyl rings (*i.e.*, two of R^{11a} , R^{11b} , R^{11c} and R^{11d} are H, and the other two of R^{11a} , R^{11b} , R^{11c} or R^{11d} are F, Cl, Br, I or ^{123}I). In other embodiments, the compounds comprise three F, Cl, Br, I or ^{123}I substitutions (*i.e.*, one of R^{11a} , R^{11b} , R^{11c} and R^{11d} is H, and the remaining R^{11a} , R^{11b} , R^{11c} or R^{11d} is F, Cl, Br, I or ^{123}I) and in other embodiments the compounds comprise four F, Cl, Br, I or ^{123}I substitutions (*i.e.*, each of R^{11a} , R^{11b} , R^{11c} and R^{11d} are F, Cl, Br, I or ^{123}I).

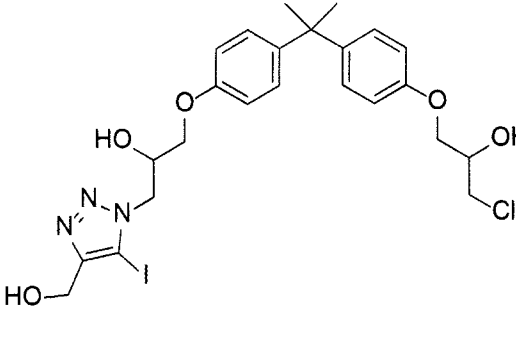
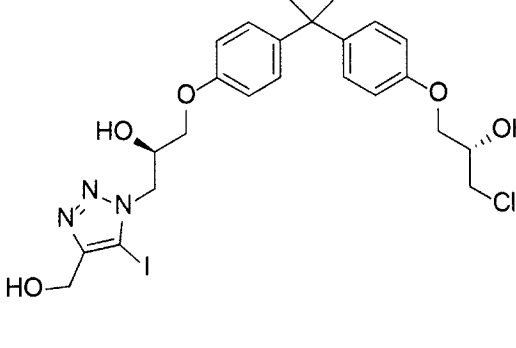
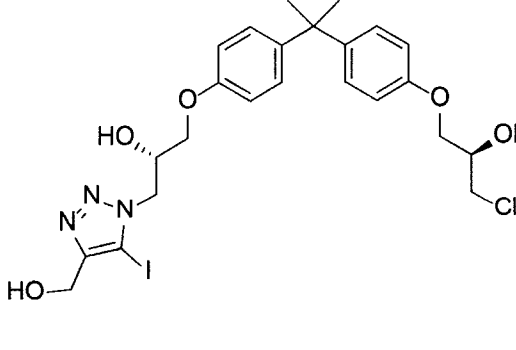
In another embodiment, the compound comprise one or more F, Cl, Br, I or ^{123}I substitutions for R^3 . In one embodiment, the compound comprise one or more I or ^{123}I substitutions for R^3 .

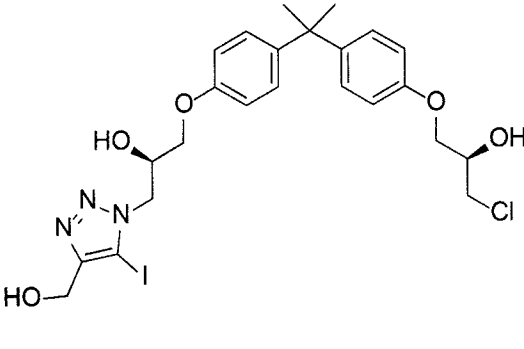
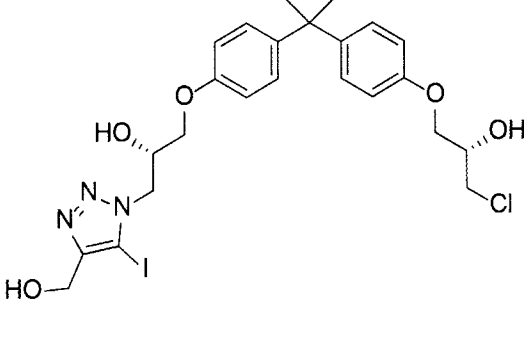
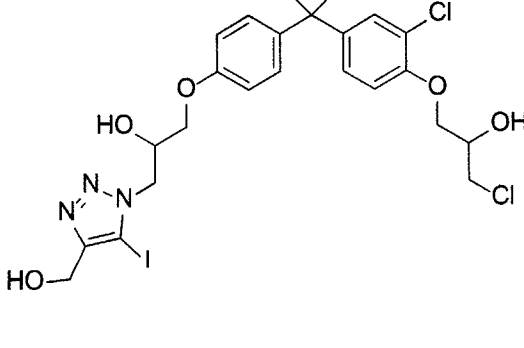
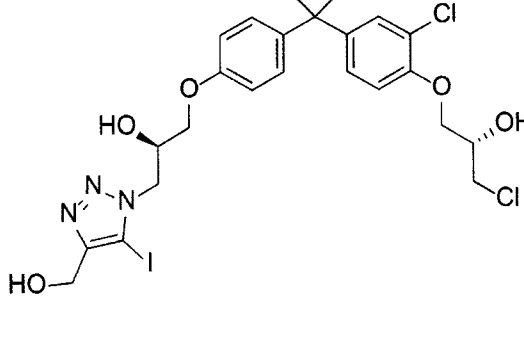
In some embodiments, the compound comprises at least one R^6 substituent on R^3 , wherein at least one R^6 is further substituted with at least one of ^{123}I , I, or ^{18}F .

Favorable imaging and/or AR NTD modulating results can be obtained by substitution with F, Cl, Br, I or ^{123}I at any of the " R^{11} " positions. In some of the foregoing embodiments, R^{11a} is ^{123}I . In other embodiments, R^{11c} is ^{123}I .

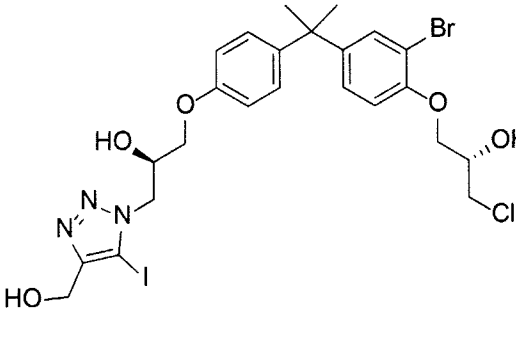
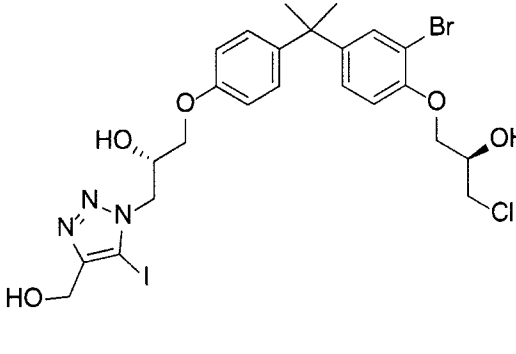
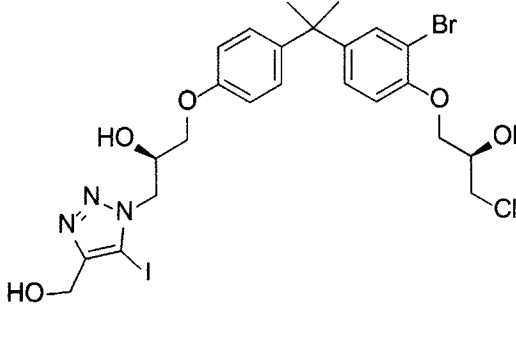
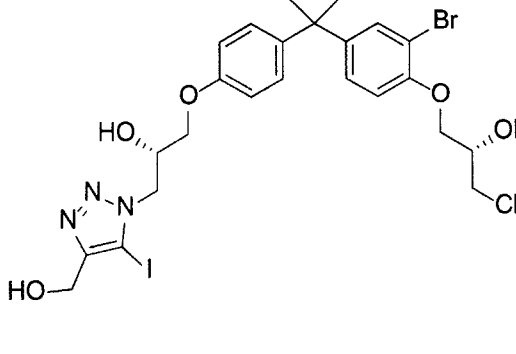
In some more specific embodiments of the compound of Formula I, the compound has one of the following structures from Table 1, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof:

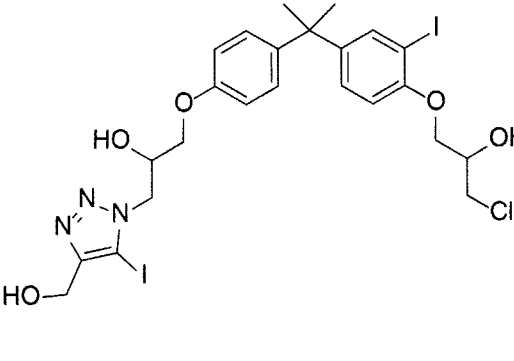
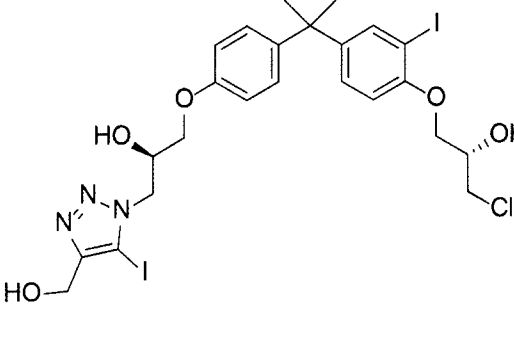
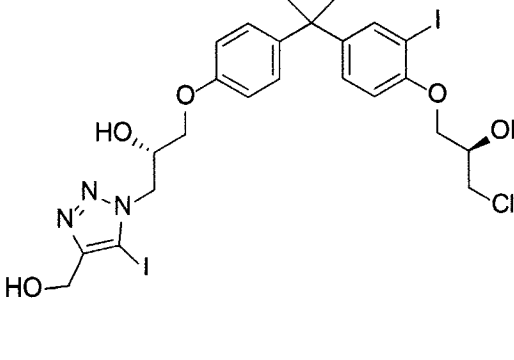
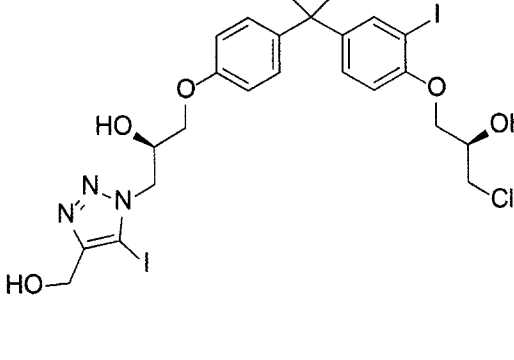
Table 1. Compounds

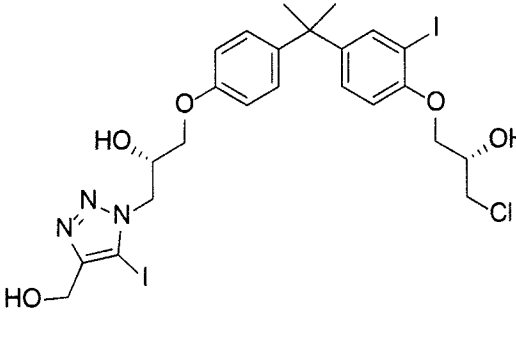
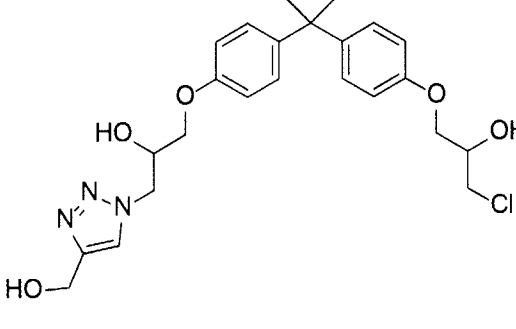
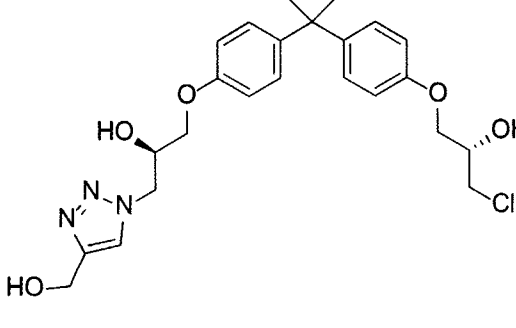
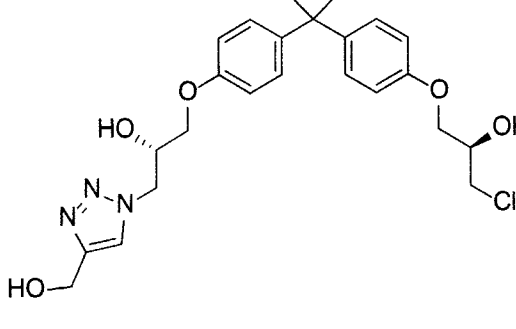
No.	Structure	Name
1		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
1a		<i>(S)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
1b		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

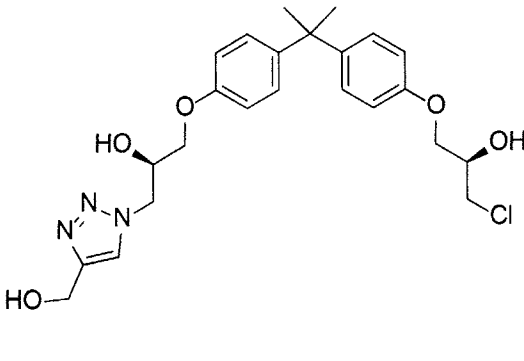
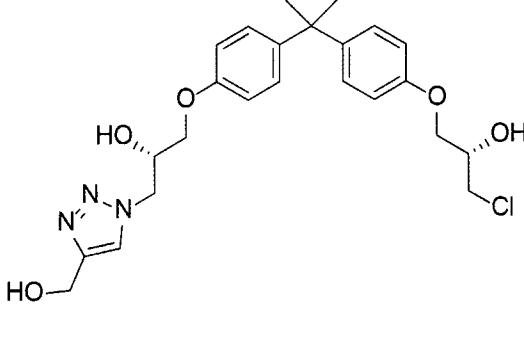
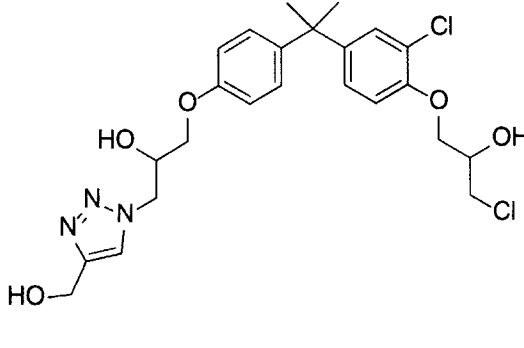
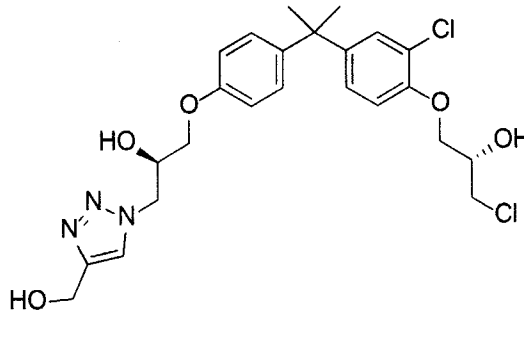
No.	Structure	Name
1c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
1d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
2		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
2a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
2b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
2c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
2d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
3		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

No.	Structure	Name
3a		(R)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
3b		(S)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
3c		(R)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
3d		(S)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol

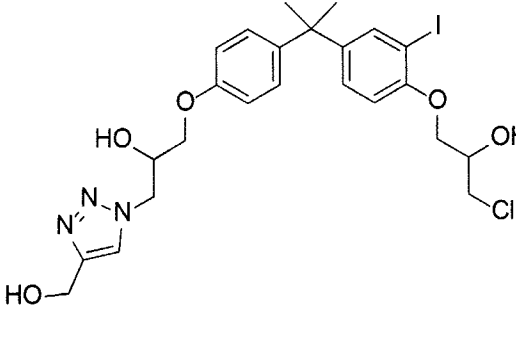
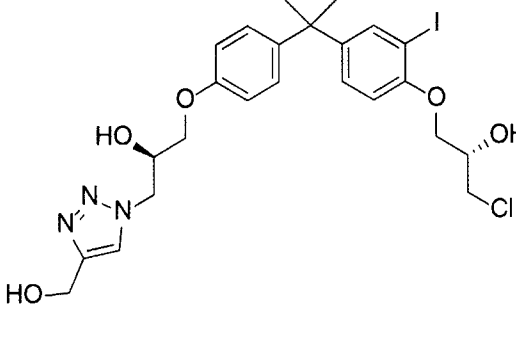
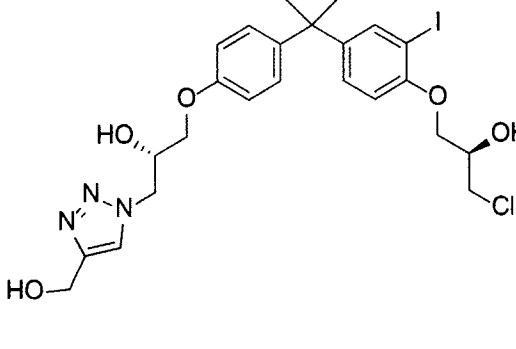
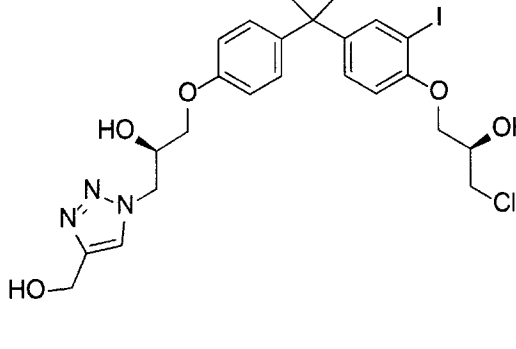
No.	Structure	Name
4		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
4a		(S)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
4b		(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
4c		(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

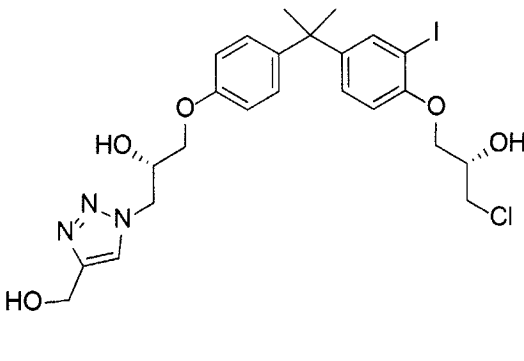
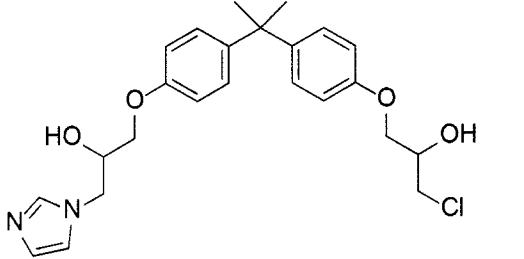
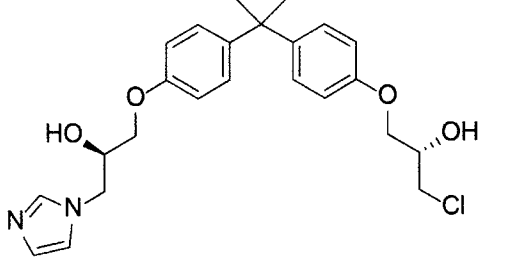
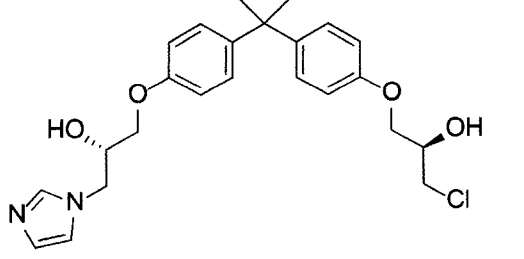
No.	Structure	Name
4d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
5		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
5a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
5b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
5c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
5d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
6		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
6a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
6b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
6c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
6d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
7		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

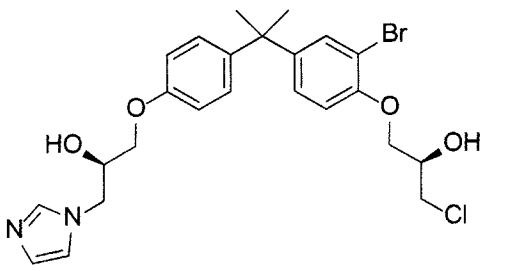
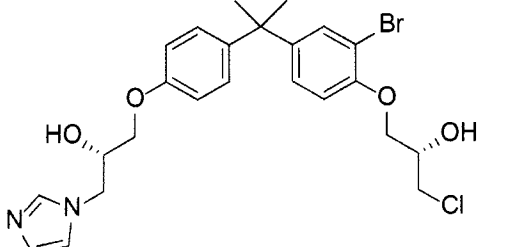
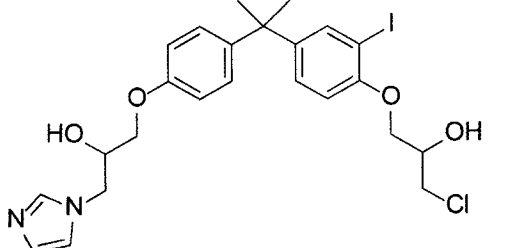
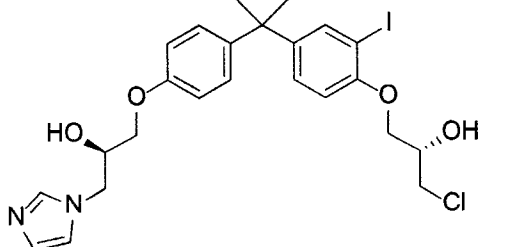
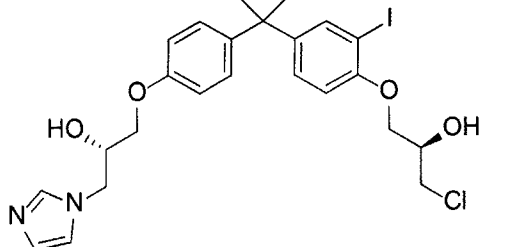
No.	Structure	Name
7a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
7b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
7c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
7d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol

No.	Structure	Name
8		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
8a		<i>(S)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
8b		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>SR</i>)-2-hydroxy-3-(4-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
8c		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

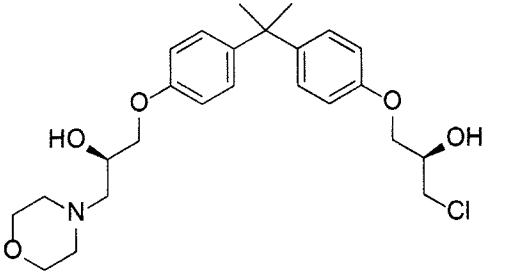
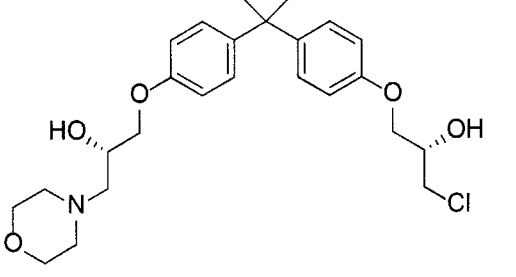
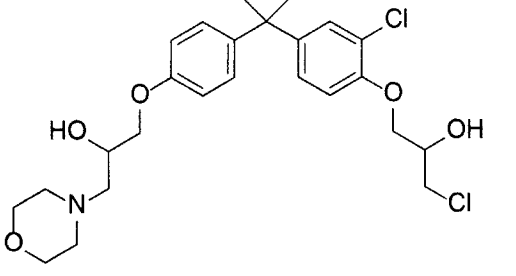
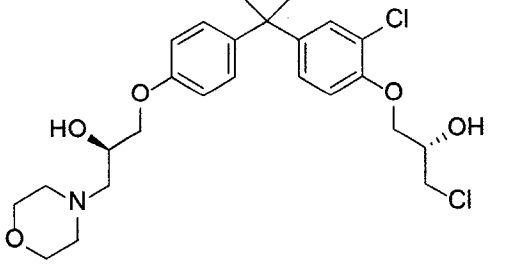
No.	Structure	Name
8d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
9		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
9a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
9b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
9c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
9d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
10		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
10a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
10b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
10c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
10d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
11		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol
11a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(1H-imidazol-1-yl)propan-2-ol
11b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(1H-imidazol-1-yl)propan-2-ol

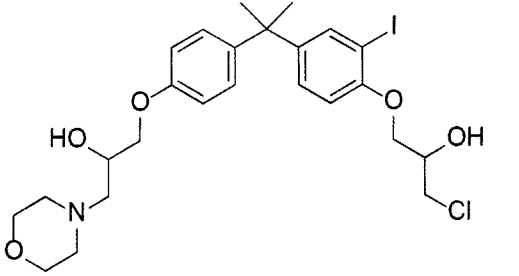
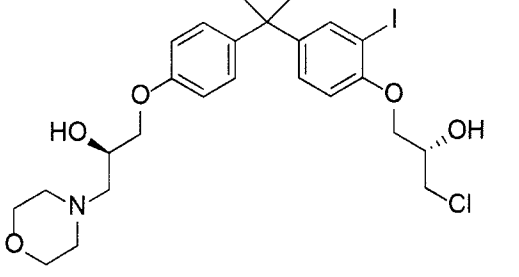
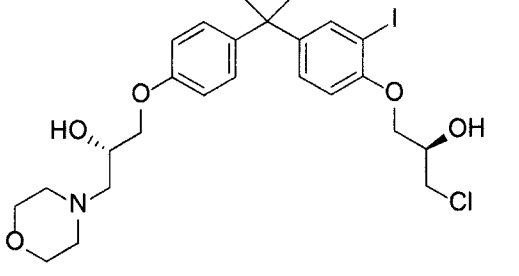
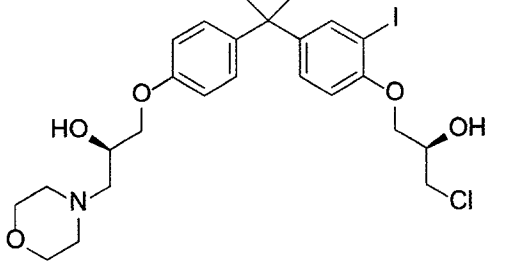
No.	Structure	Name
11c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(1H-imidazol-1-yl)propan-2-ol
11d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(1H-imidazol-1-yl)propan-2-ol
12		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
12a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
12b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

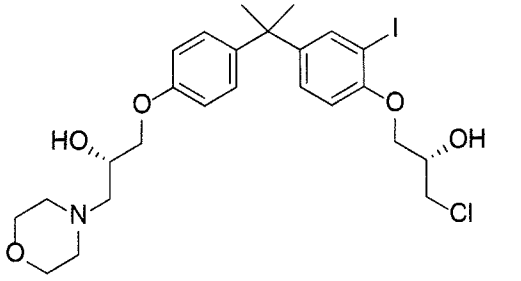
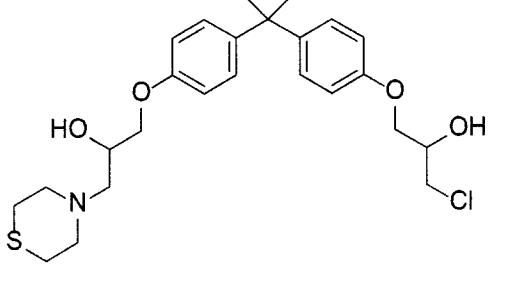
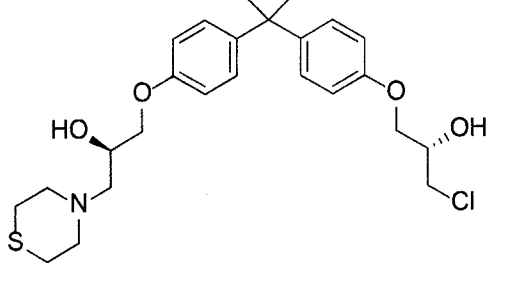
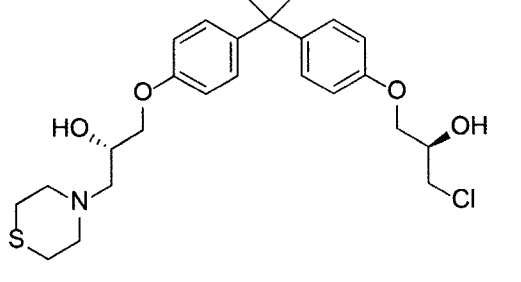
No.	Structure	Name
12c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
12d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
13		1-chloro-3-(4-(2-(4-(2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
13a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
13b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

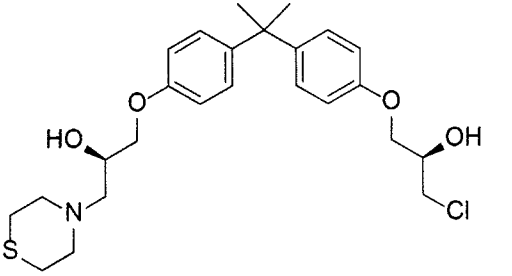
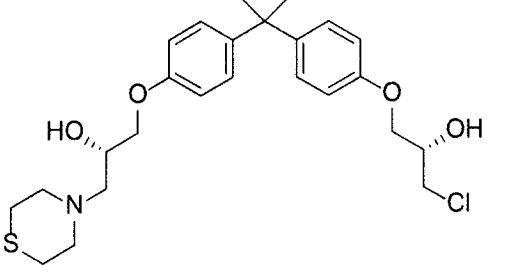
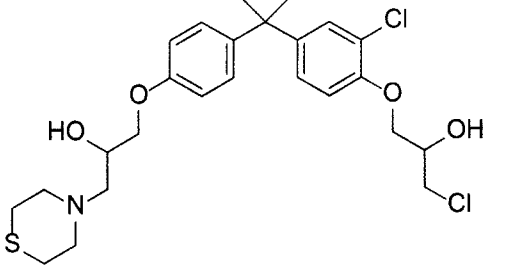
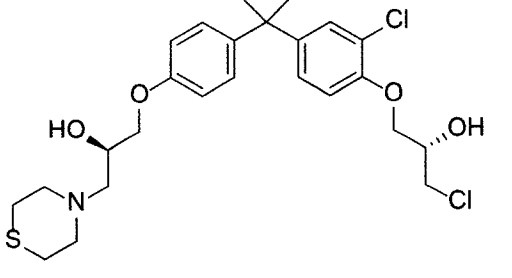
No.	Structure	Name
13c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
13d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
14		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
14a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
14b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
14c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
14d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
15		1-(2-bromo-4-(2-(4-(2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

No.	Structure	Name
15a		(R)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-morpholinopropan-2-ol
15b		(<i>S</i>)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-morpholinopropan-2-ol
15c		(R)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-morpholinopropan-2-ol
15d		(<i>S</i>)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-morpholinopropan-2-ol

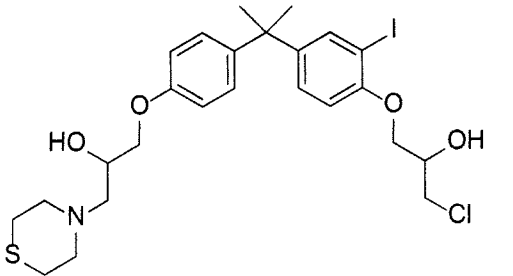
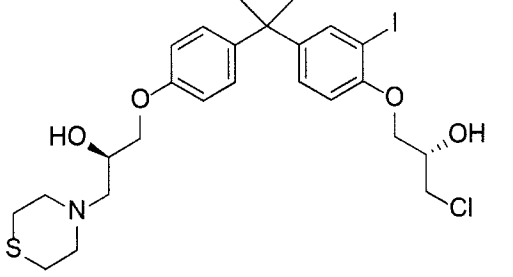
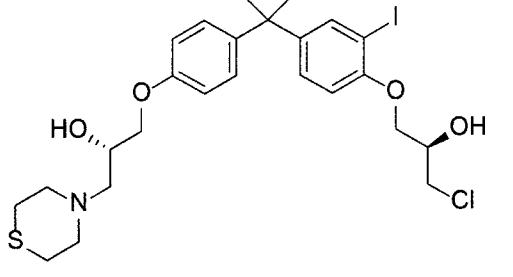
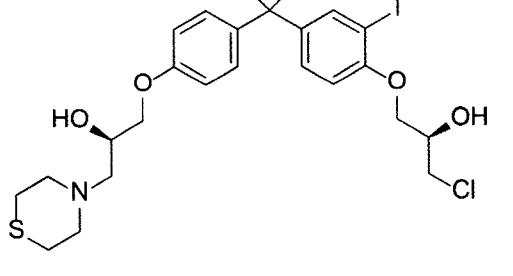
No.	Structure	Name
16		1-chloro-3-(4-(2-(4-(2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
16a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
16b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
16c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
16d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
17		1-chloro-3-(4-(2-(4-(2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
17a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
17b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
17c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
17d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
18		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
18a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
18b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
18c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
18d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
19		1-(2-bromo-4-(2-(4-(2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

No.	Structure	Name
19a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-thiomorpholinopropan-2-ol
19b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-thiomorpholinopropan-2-ol
19c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-thiomorpholinopropan-2-ol
19d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-thiomorpholinopropan-2-ol

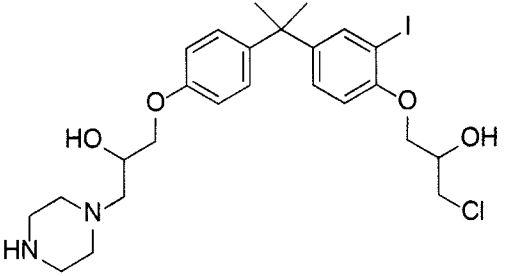
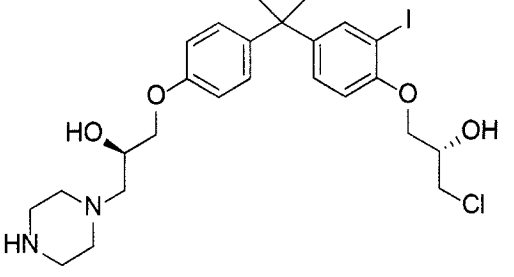
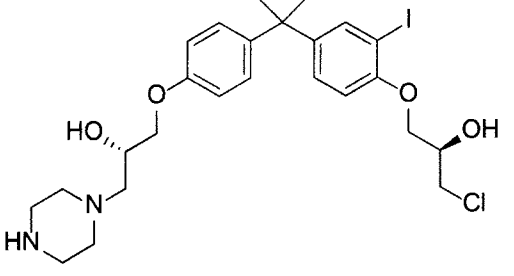
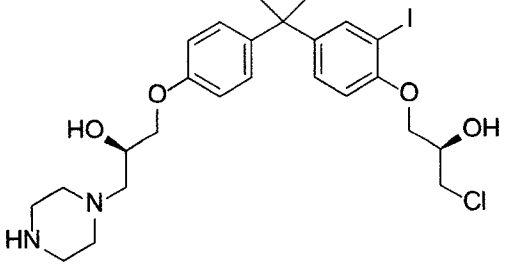
No.	Structure	Name
20		1-chloro-3-(4-(2-(4-(2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
20a		<i>(S)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
20b		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>RS</i>)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
20c		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

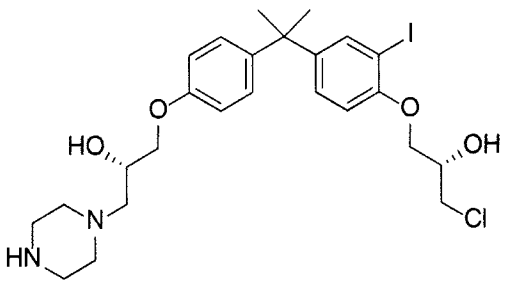
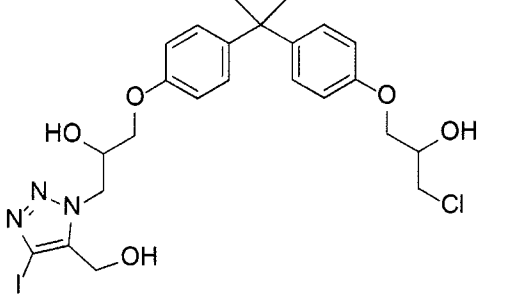
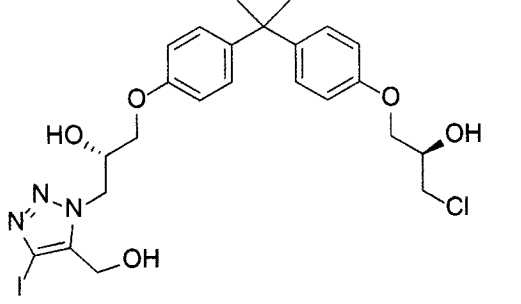
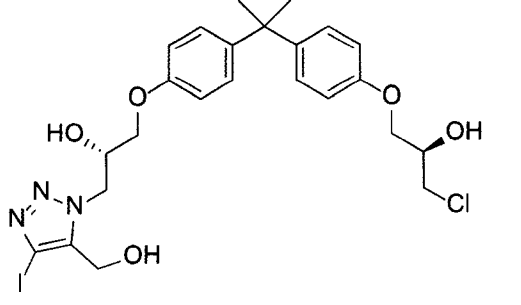
No.	Structure	Name
20d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
21		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
21a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
21b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
21c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
21d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
22		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
22a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
22b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
22c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
22d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
23		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

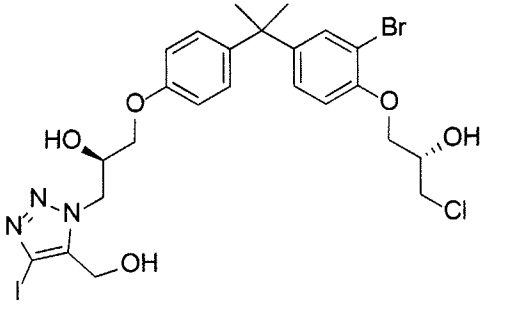
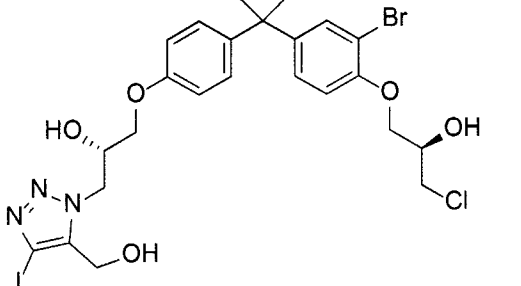
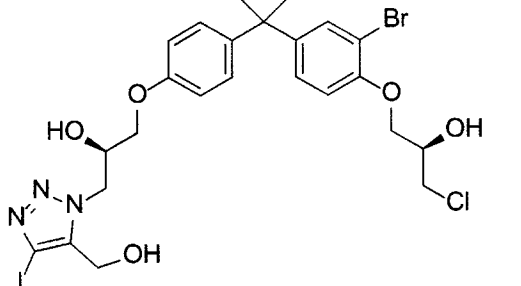
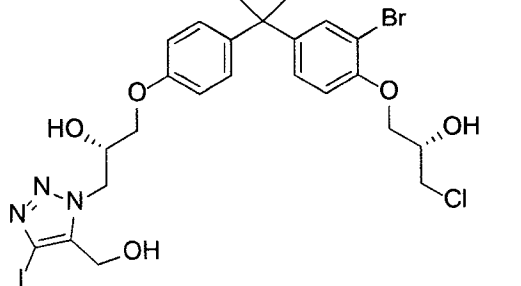
No.	Structure	Name
23a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(piperazin-1-yl)propan-2-ol
23b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(piperazin-1-yl)propan-2-ol
23c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(piperazin-1-yl)propan-2-ol
23d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(piperazin-1-yl)propan-2-ol

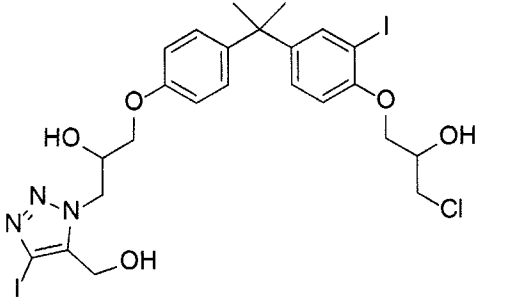
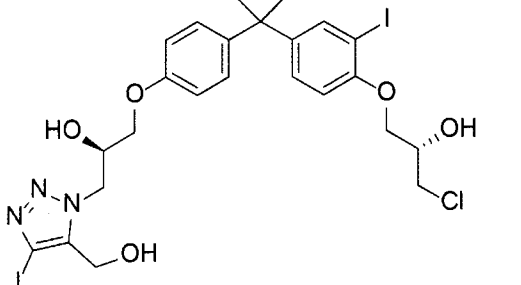
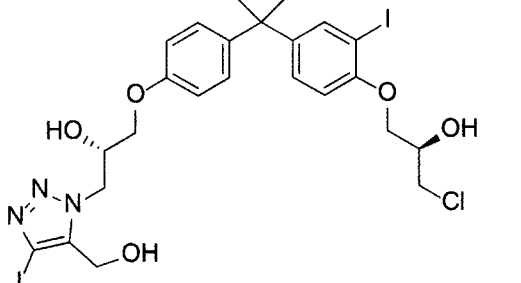
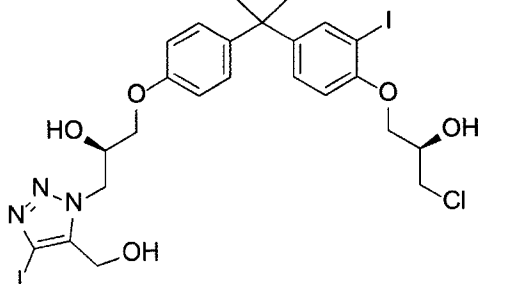
No.	Structure	Name
24		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
24a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
24b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
24c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
24d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
25		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
25a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
25b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

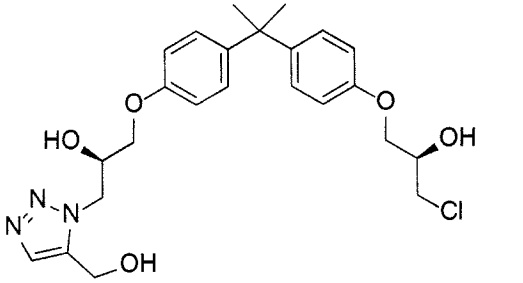
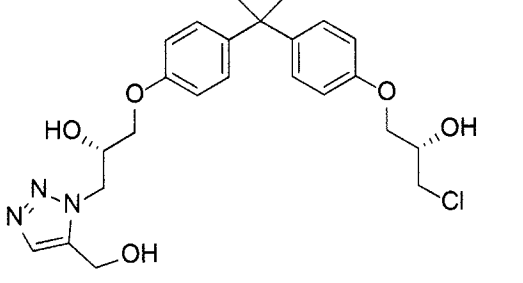
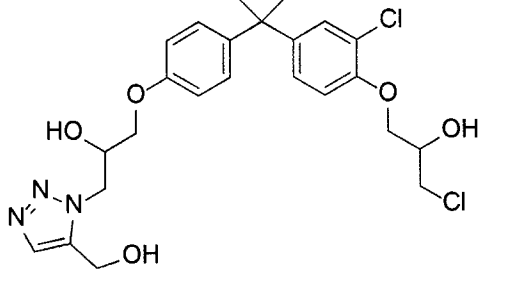
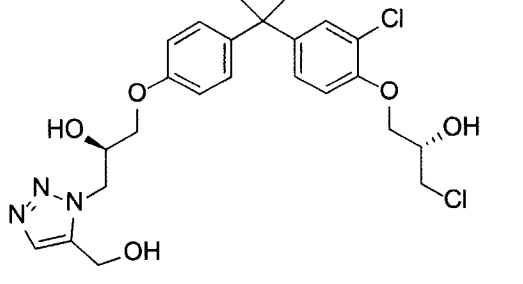
No.	Structure	Name
25c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
25d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
26		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
26a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
26b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
26c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
26d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
27		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

No.	Structure	Name
27a		(R)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
27b		(S)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
27c		(R)-1-(4-(2-(3-bromo-4-((<i>R</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol
27d		(S)-1-(4-(2-(3-bromo-4-((<i>S</i>)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propan-2-ol

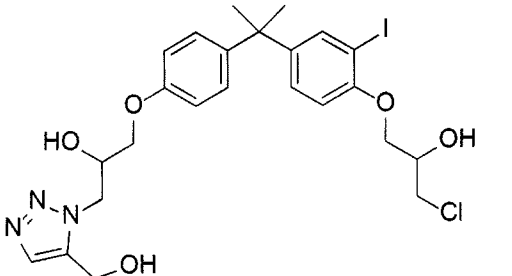
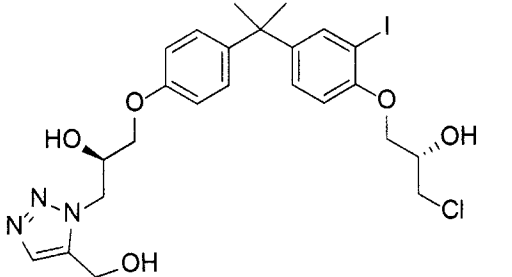
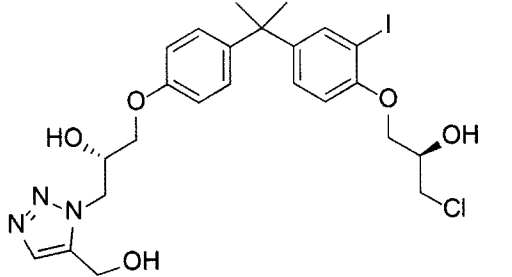
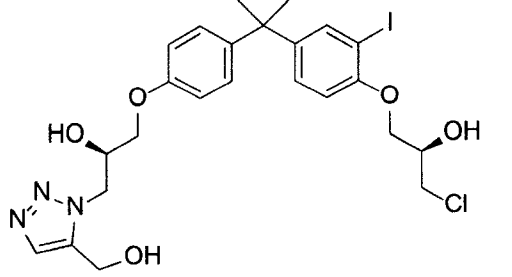
No.	Structure	Name
28		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
28a		(S)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
28b		(R)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
28c		(R)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

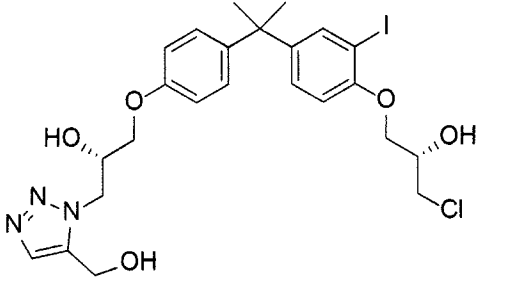
No.	Structure	Name
28d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
29		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
29a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
29b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
29c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
29d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
30		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
30a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
30b		<p>(<i>R</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1<i>H</i>-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
30c		<p>(<i>R</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1<i>H</i>-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
30d		<p>(<i>S</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1<i>H</i>-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
31		<p>1-(2-bromo-4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-1<i>H</i>-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol</p>

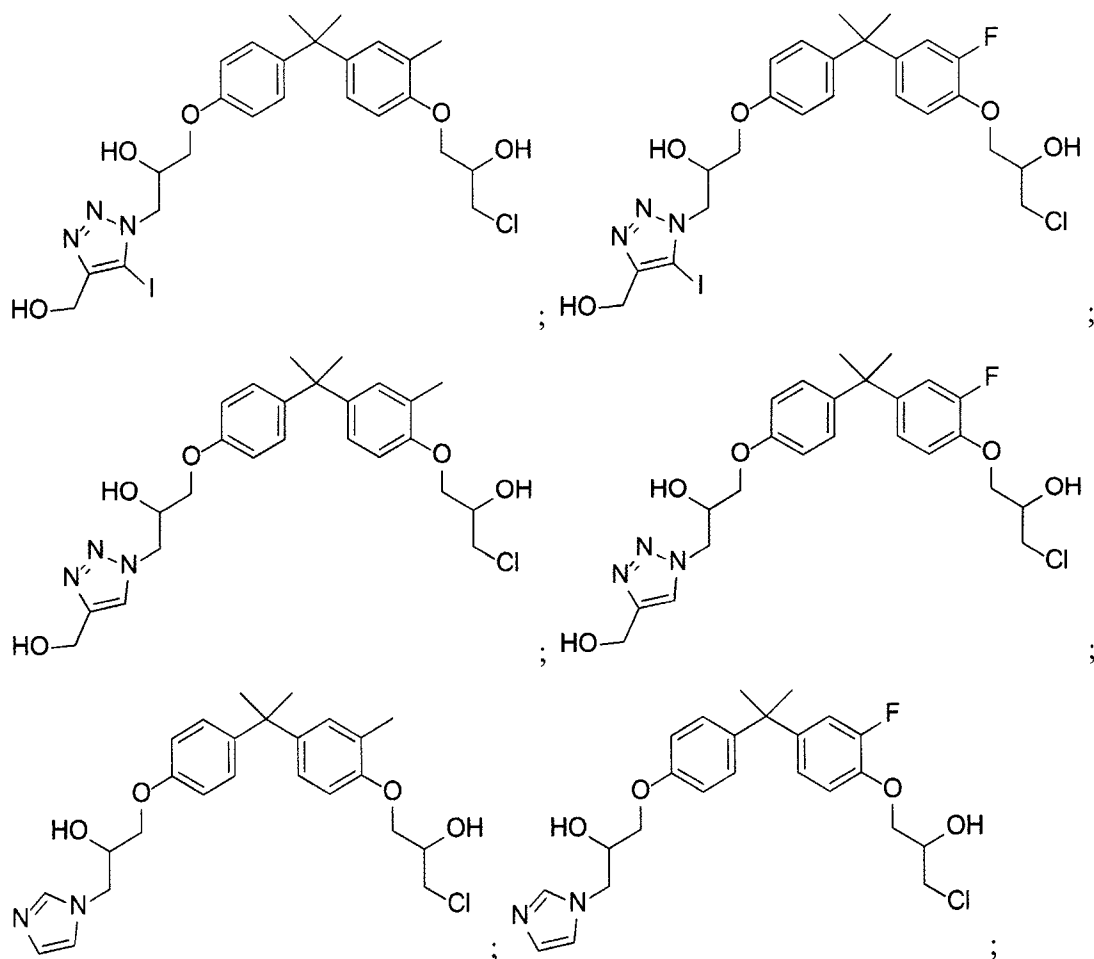
No.	Structure	Name
31a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
31b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
31c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
31d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propan-2-ol

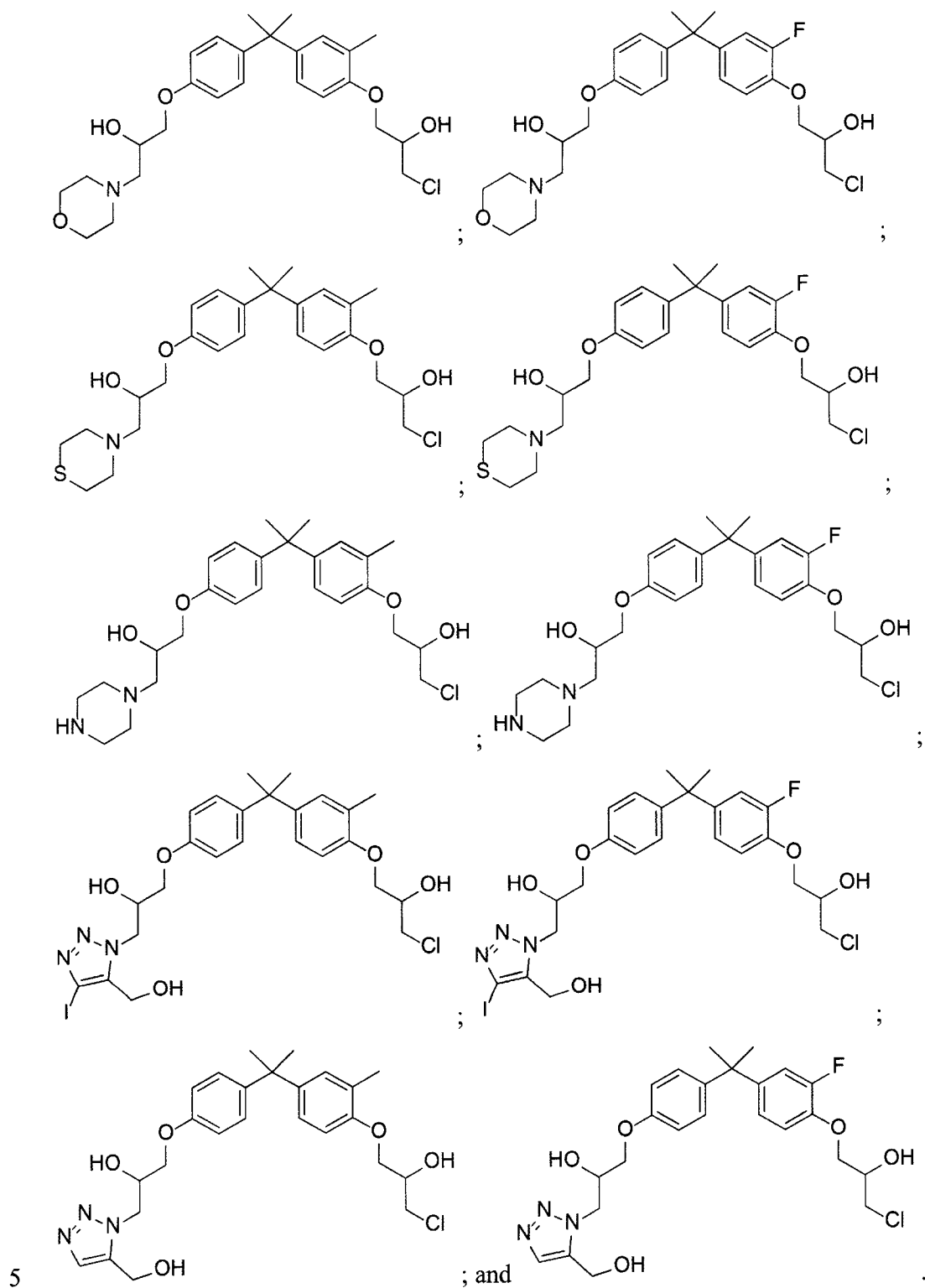
No.	Structure	Name
32		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
32a		<i>(S)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
32b		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
32c		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
32d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

In some more specific embodiments of the compound of Formula I, the compound has one of the following structures, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof including all 4 distinct stereoisomers e.g., (*S,R*), (*R,S*), (*R,R*), and

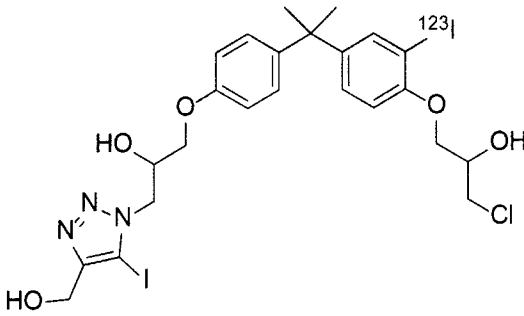
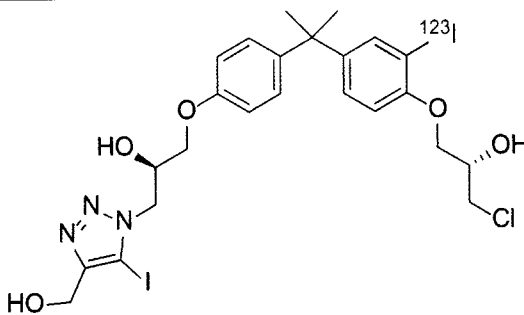
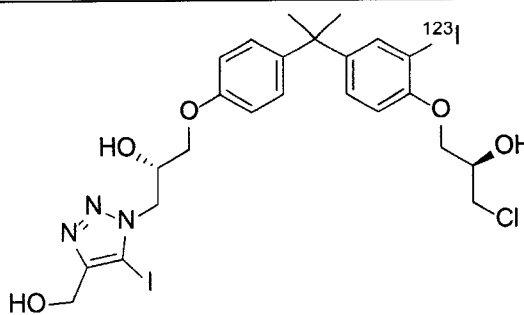
5 (*S,S*):





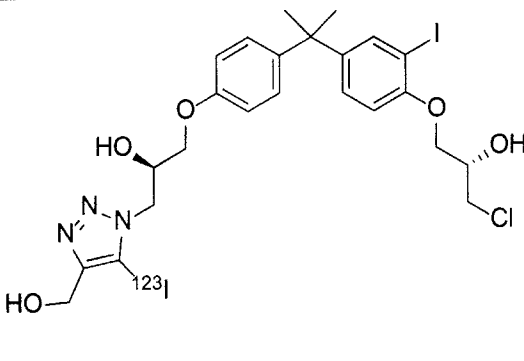
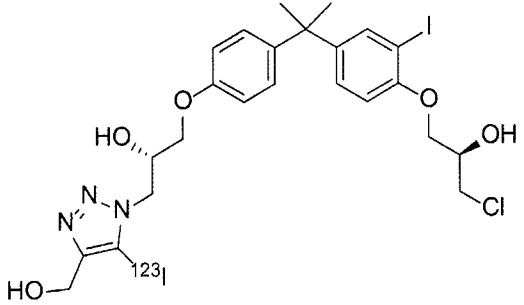
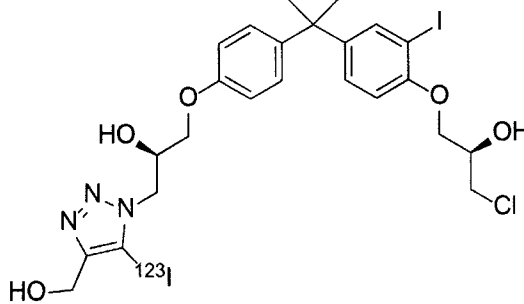
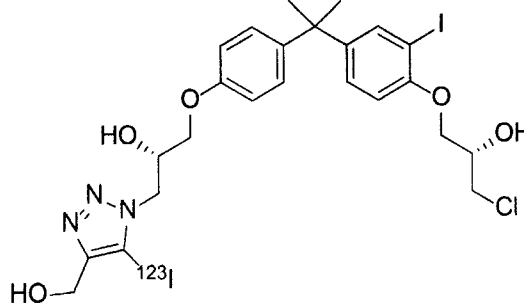
In some more specific embodiments of the compound of Formula I, the compound has one of the following structures from Table 2, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof:

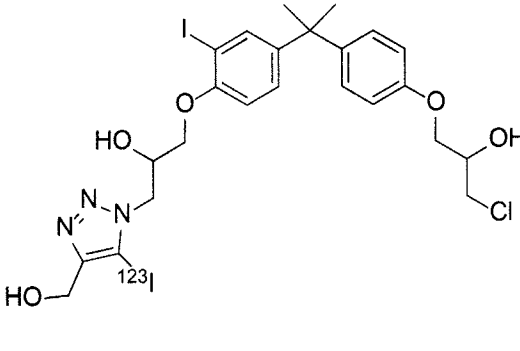
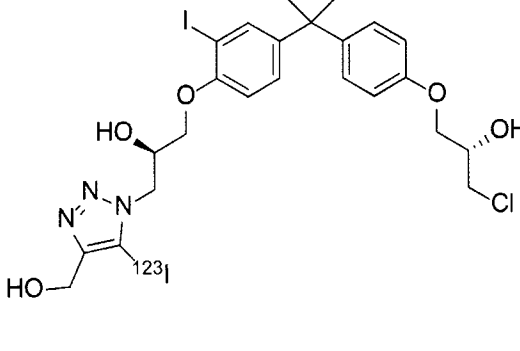
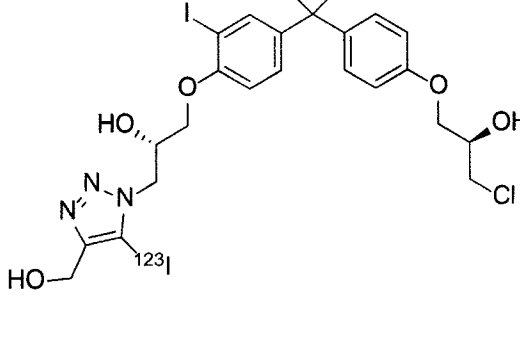
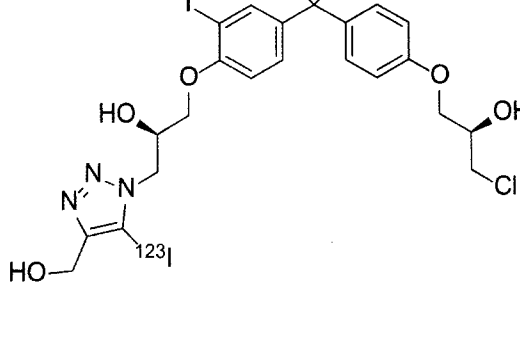
Table 2. ^{123}I Compounds

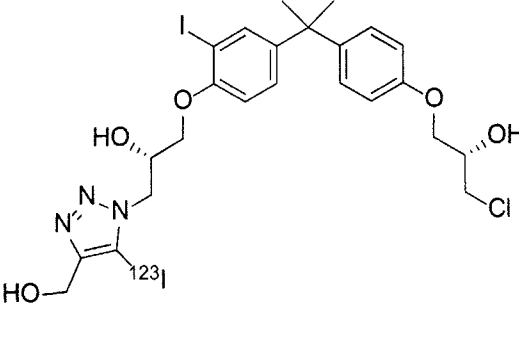
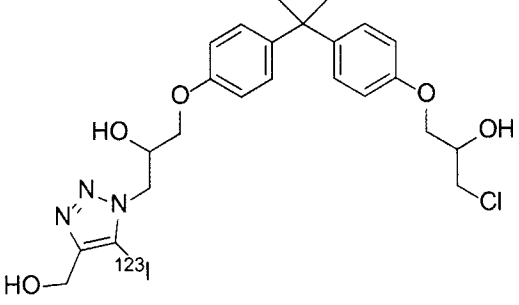
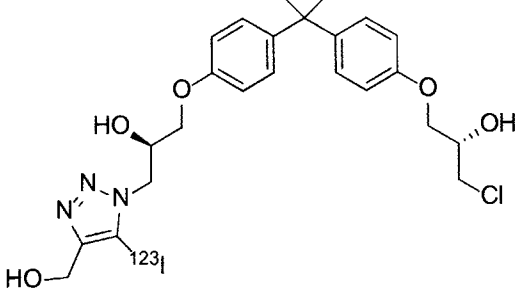
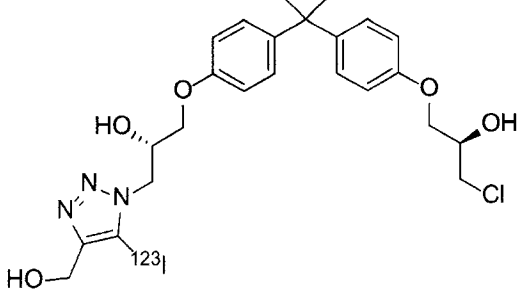
No.	Structure	Name
33		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ^{123}I)phenoxy)propan-2-ol
33a		(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ^{123}I)phenoxy)propan-2-ol
33b		(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ^{123}I)phenoxy)propan-2-ol

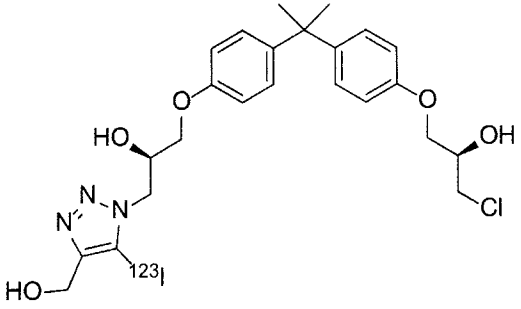
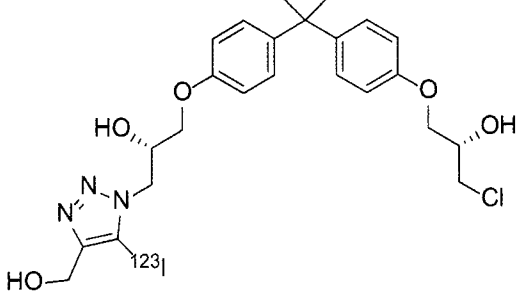
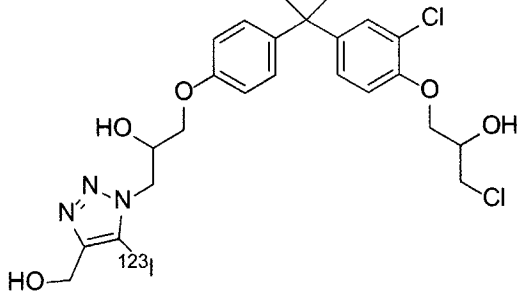
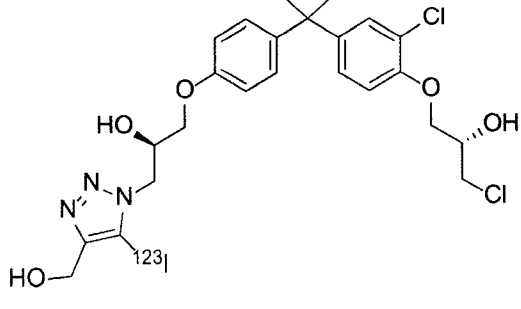
No.	Structure	Name
33c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
33d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
34		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)-3-(iodo- ¹²³ I)phenyl)propan-2-yl)phenoxy)propan-2-ol
34a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)-3-(iodo- ¹²³ I)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
34b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)-3-(iodo- ¹²³ I)phenyl)propan-2-yl)phenoxy)propan-2-ol
34c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)-3-(iodo- ¹²³ I)phenyl)propan-2-yl)phenoxy)propan-2-ol
34d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-iodo-1H-1,2,3-triazol-1-yl)propoxy)-3-(iodo- ¹²³ I)phenyl)propan-2-yl)phenoxy)propan-2-ol
35		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
35a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
35b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
35c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
35d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

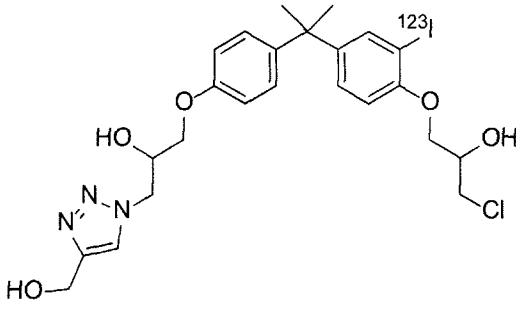
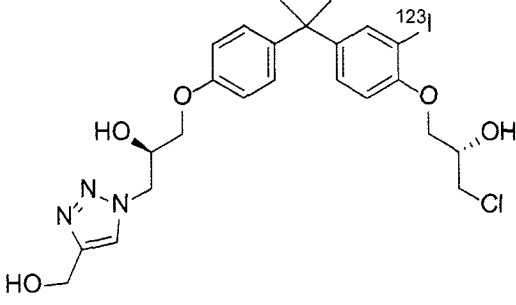
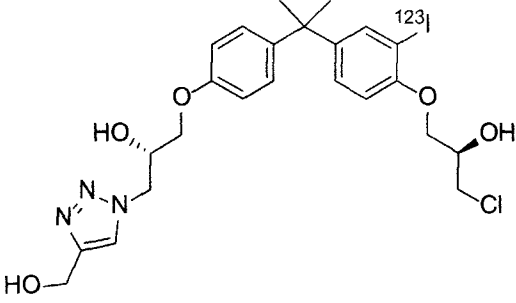
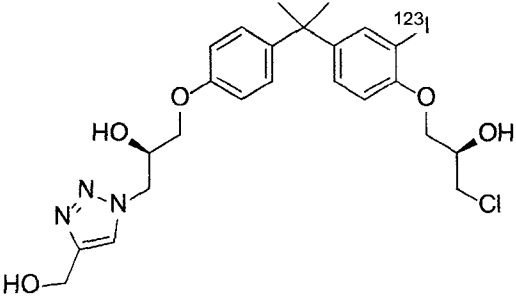
No.	Structure	Name
36		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)-3-iodophenyl)propan-2-yl)phenoxy)propan-2-ol
36a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)-3-iodophenyl)propan-2-yl)phenoxy)propan-2-ol
36b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)-3-iodophenyl)propan-2-yl)phenoxy)propan-2-ol
36c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)-3-iodophenyl)propan-2-yl)phenoxy)propan-2-ol

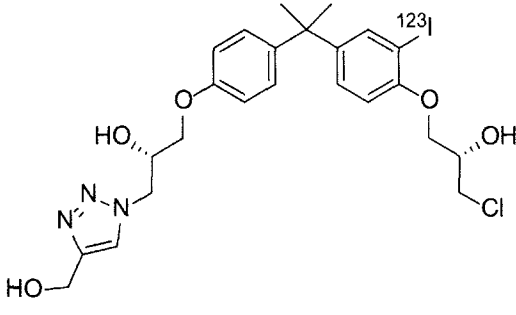
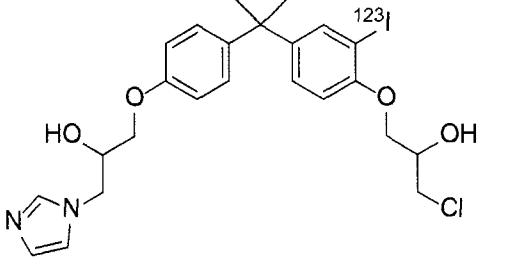
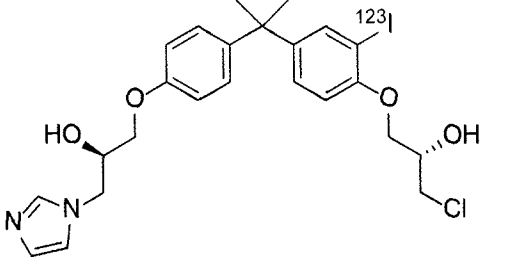
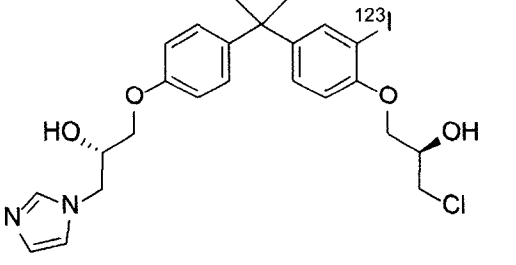
No.	Structure	Name
36d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)-3-iodophenyl)propan-2-yl)phenoxy)propan-2-ol
37		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
37a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
37b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
37c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
37d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
38		1-chloro-3-(2-chloro-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
38a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

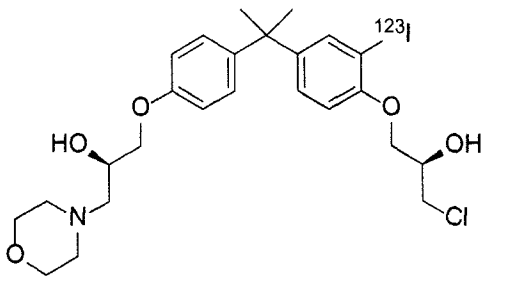
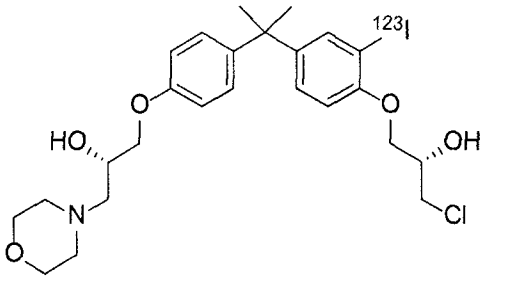
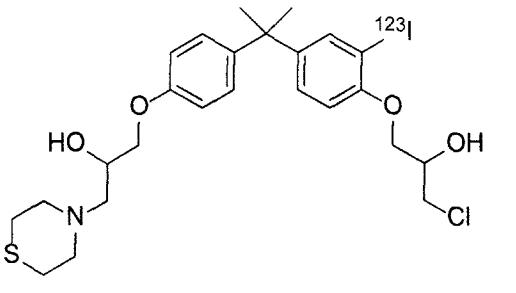
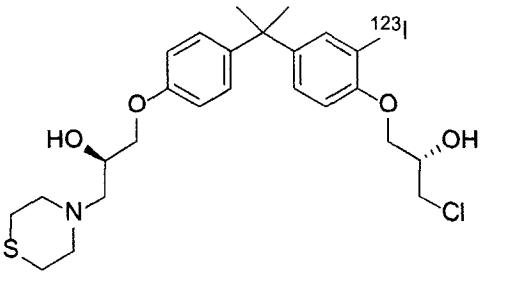
No.	Structure	Name
38b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
38c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
38d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
39		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol

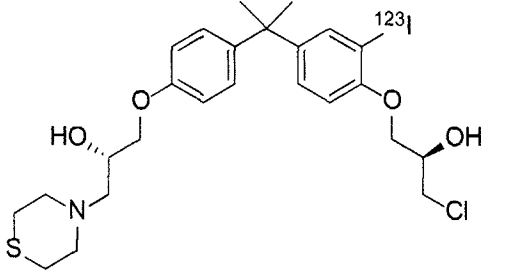
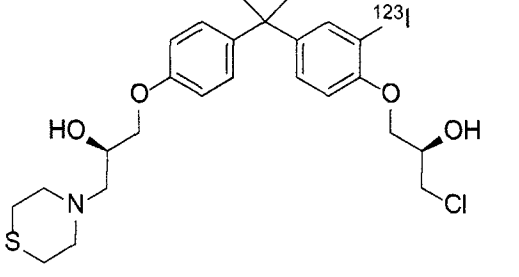
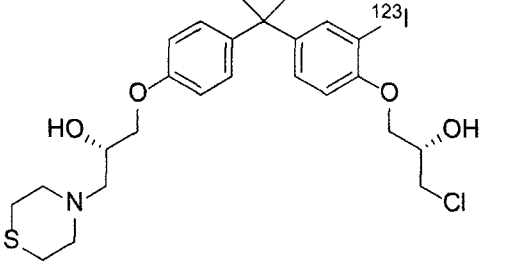
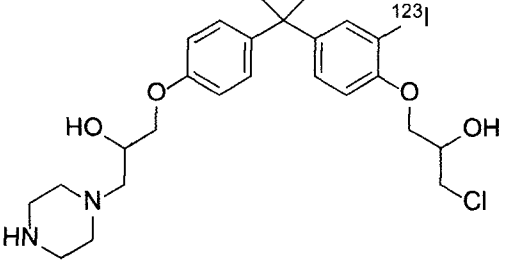
No.	Structure	Name
39a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
39b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
39c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
39d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-(hydroxymethyl)-5-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol

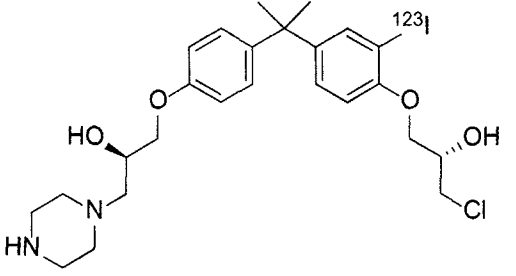
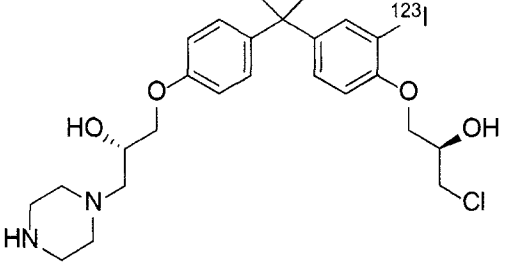
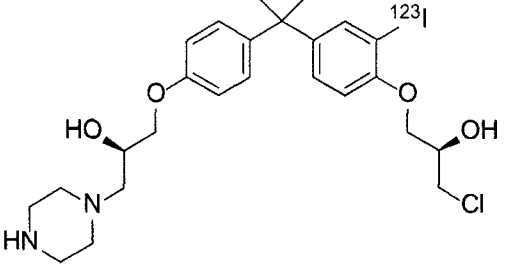
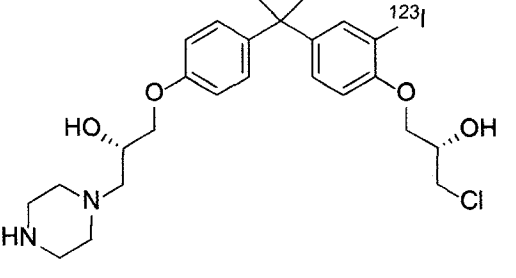
No.	Structure	Name
40		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
40a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
40b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
40c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

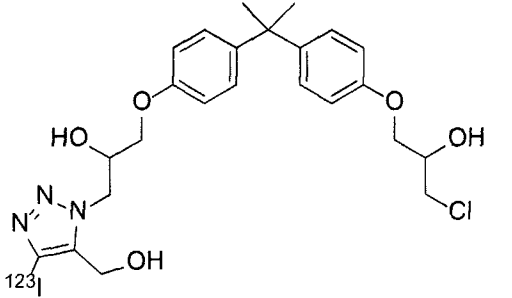
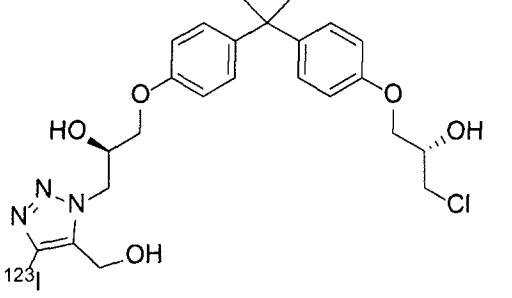
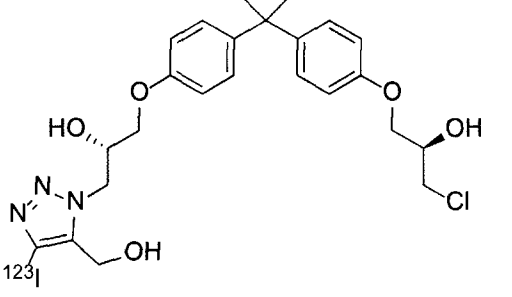
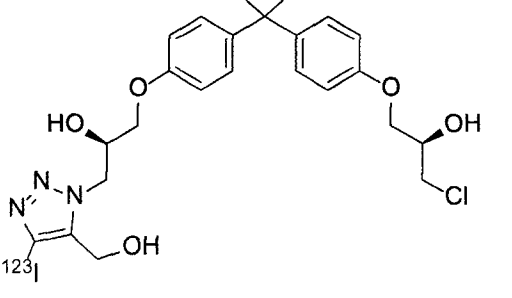
No.	Structure	Name
40d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(4-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
41		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
41a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
41b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(1H-imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

No.	Structure	Name
41c		(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(1 <i>H</i> -imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
41d		(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(1 <i>H</i> -imidazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
42		1-chloro-3-(4-(2-(4-(2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
42a		(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
42b		(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol

No.	Structure	Name
42c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
42d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-morpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
43		1-chloro-3-(4-(2-(4-(2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
43a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

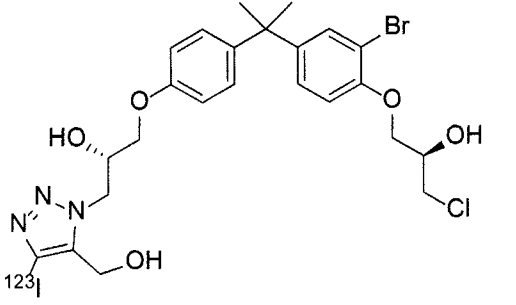
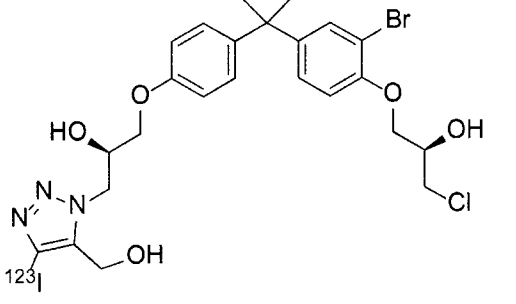
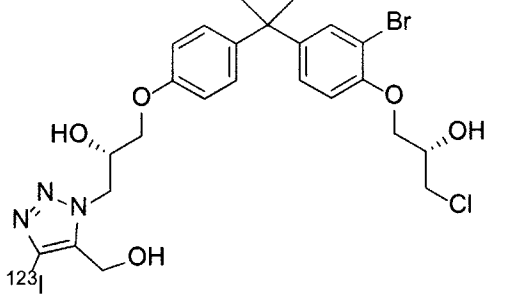
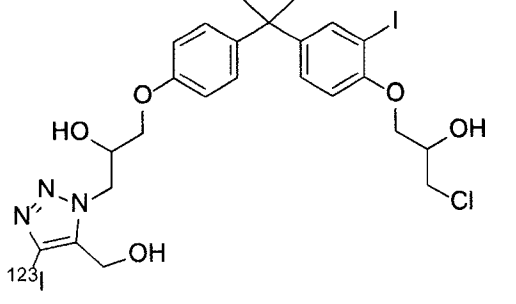
No.	Structure	Name
43b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
43c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
43d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-thiomorpholinopropoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
44		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

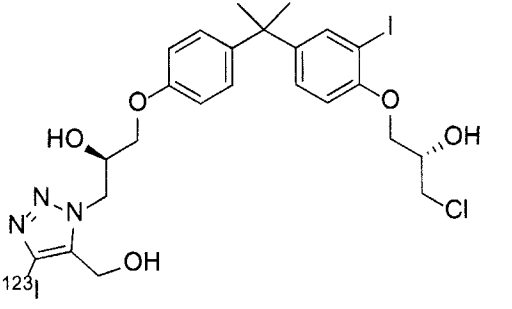
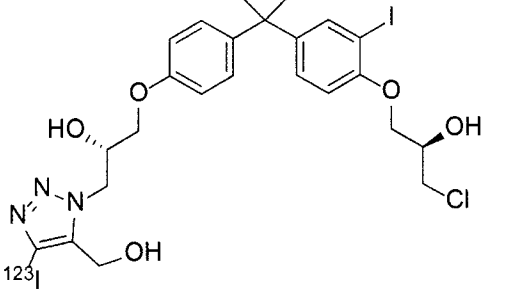
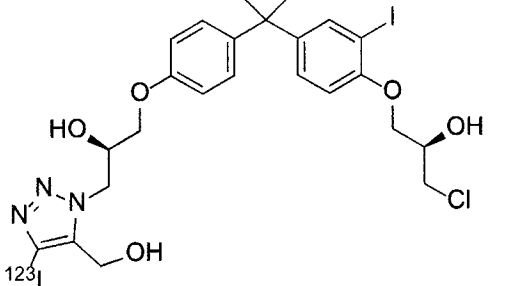
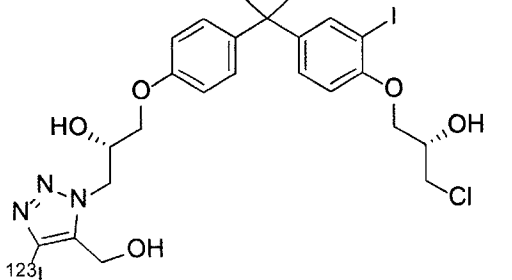
No.	Structure	Name
44a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
44b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
44c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
44d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(piperazin-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

No.	Structure	Name
45		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
45a		(S)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
45b		(R)-1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
45c		(R)-1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
45d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
46		1-chloro-3-(2-chloro-4-(2-(4(2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
46a		(S)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
46b		(R)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
46c		(R)-1-chloro-3-(2-chloro-4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
46d		(S)-1-chloro-3-(2-chloro-4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
47		1-(2-bromo-4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol
47a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol

No.	Structure	Name
47b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
47c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
47d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propan-2-ol
48		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

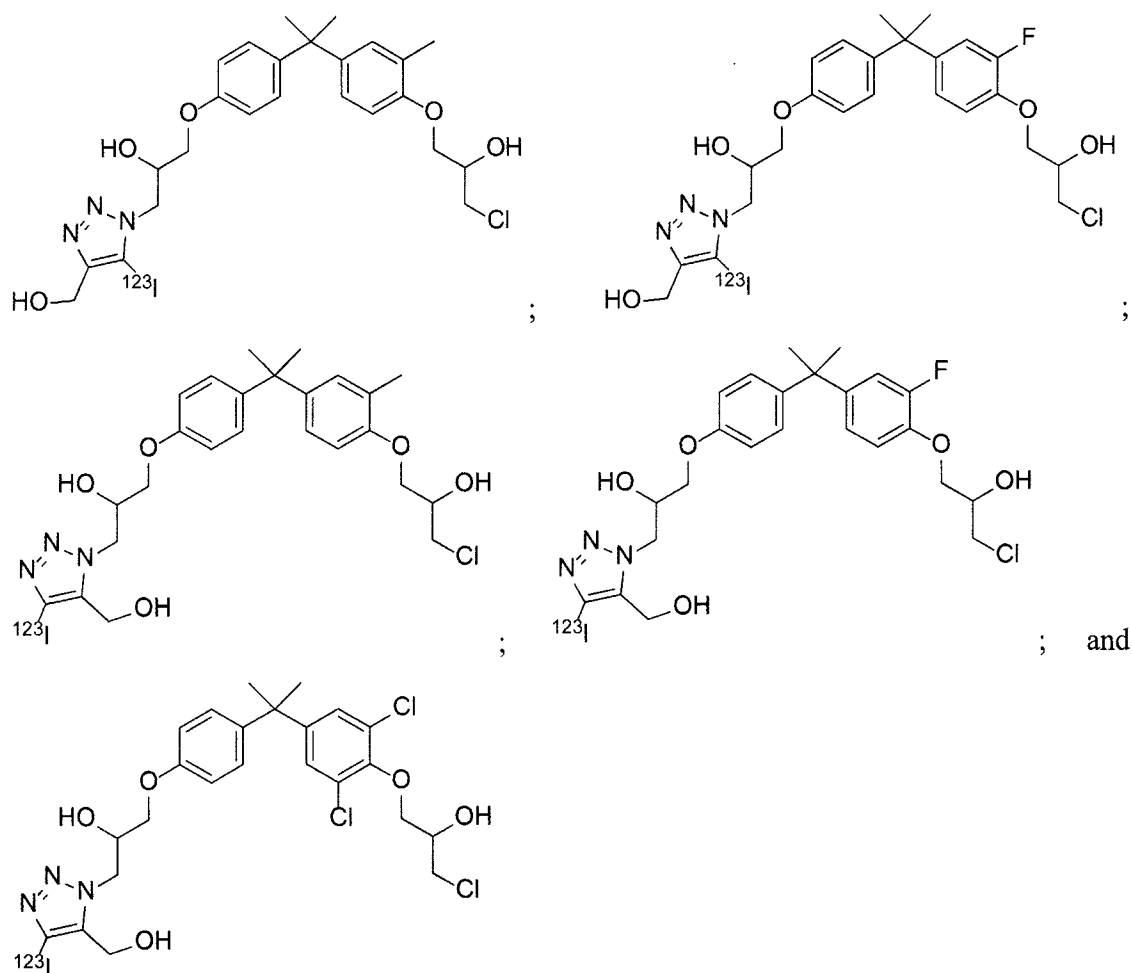
No.	Structure	Name
48a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
48b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
48c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
48d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-(iodo- ¹²³ I)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
49		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
49a		<i>(S)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
49b		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>S</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol
49c		<i>(R)</i> -1-chloro-3-(4-(2-(4-((<i>R</i>)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1 <i>H</i> -1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ <i>I</i>)phenoxy)propan-2-ol

No.	Structure	Name
49d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-4-iodo-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
50		1-chloro-3-(4-(2-(4-(2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
50a		(S)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
50b		(R)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

No.	Structure	Name
50c		(R)-1-chloro-3-(4-(2-(4-((R)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol
50d		(S)-1-chloro-3-(4-(2-(4-((S)-2-hydroxy-3-(5-(hydroxymethyl)-1H-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-(iodo- ¹²³ I)phenoxy)propan-2-ol

In some embodiments of the compound of Formula I, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof including all 4 distinct stereoisomers e.g., (*S,R*), (*R,S*), (*R,R*), and (*S,S*), has one of the following structures,:



In some embodiments of the compound of Formula I, the compound has one of the following structures from Table 3, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof:

Table 3. ^{18}F Compounds

No.	Structure	Name
51		1-chloro-3-(4-(2-(4-(3-(4-((fluoro- ^{18}F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
51a		(S)-1-chloro-3-(4-(2-(4-((R)-3-(4-((fluoro- ^{18}F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol
51b		(R)-1-chloro-3-(4-(2-(4-((S)-3-(4-((fluoro- ^{18}F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

No.	Structure	Name
51c		<p>(<i>R</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
51d		<p>(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
52		<p>1-chloro-3-(2-chloro-4-(2-(4-(3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
52a		<p>(<i>S</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>

No.	Structure	Name
52b		<p>(<i>R</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
52c		<p>(<i>R</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
52d		<p>(<i>S</i>)-1-chloro-3-(2-chloro-4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
53		<p>1-(2-bromo-4-(2-(4-(3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-chloropropan-2-ol</p>

No.	Structure	Name
53a		(R)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-((fluoro- ¹⁸ F)methyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
53b		(S)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-((fluoro- ¹⁸ F)methyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
53c		(R)-1-(4-(2-(3-bromo-4-((R)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-((fluoro- ¹⁸ F)methyl)-1H-1,2,3-triazol-1-yl)propan-2-ol
53d		(S)-1-(4-(2-(3-bromo-4-((S)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)-3-(4-((fluoro- ¹⁸ F)methyl)-1H-1,2,3-triazol-1-yl)propan-2-ol

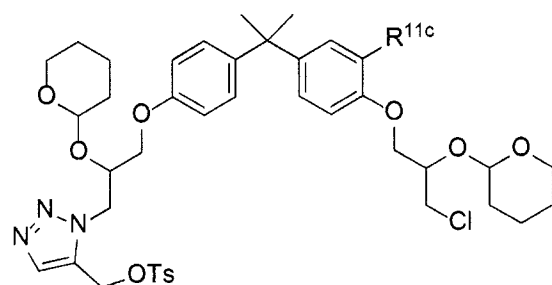
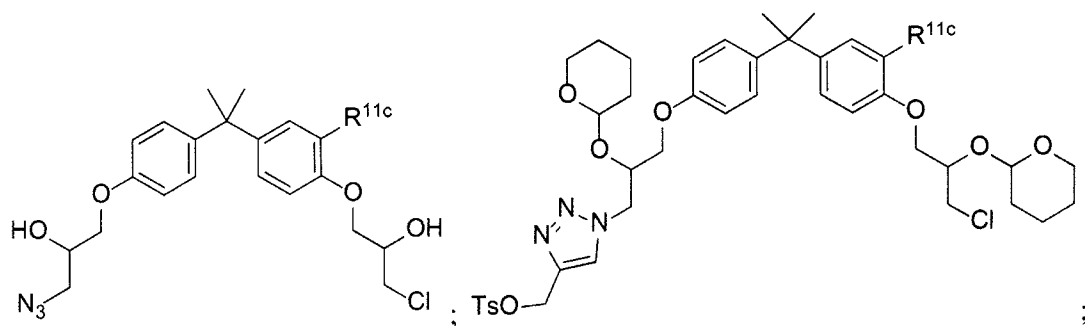
No.	Structure	Name
54		1-chloro-3-(4-(2-(4-(3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
54a		(S)-1-chloro-3-(4-(2-(4-((<i>R</i>)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
54b		(R)-1-chloro-3-(4-(2-(4-((<i>S</i>)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol
54c		(R)-1-chloro-3-(4-(2-(4-((<i>R</i>)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol

No.	Structure	Name
54d		<p>(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol</p>
55		<p>1-chloro-3-(2-fluoro-4-(2-(4-(3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
55a		<p>(<i>S</i>)-1-chloro-3-(2-fluoro-4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
55b		<p>(<i>R</i>)-1-chloro-3-(2-fluoro-4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>

No.	Structure	Name
55c		<p>(<i>R</i>)-1-chloro-3-(2-fluoro-4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
55d		<p>(<i>S</i>)-1-chloro-3-(2-fluoro-4-(2-(4-((<i>S</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol</p>
56		<p>1-chloro-3-(4-(2-(4-(3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-methylphenoxy)propan-2-ol</p>
56a		<p>(<i>S</i>)-1-chloro-3-(4-(2-(4-((<i>R</i>)-3-(4-((fluoro-¹⁸F)methyl)-1<i>H</i>-1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-methylphenoxy)propan-2-ol</p>

No.	Structure	Name
56b		(R)-1-chloro-3-(4-(2-(4-((S)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-methylphenoxy)propan-2-ol
56c		(R)-1-chloro-3-(4-(2-(4-((R)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-methylphenoxy)propan-2-ol
56d		(S)-1-chloro-3-(4-(2-(4-((S)-3-(4-((fluoro- ¹⁸ F)methyl)-1 <i>H</i> -1,2,3-triazol-1-yl)-2-hydroxypropoxy)phenyl)propan-2-yl)-2-methylphenoxy)propan-2-ol

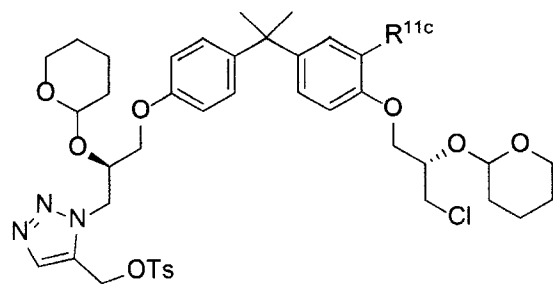
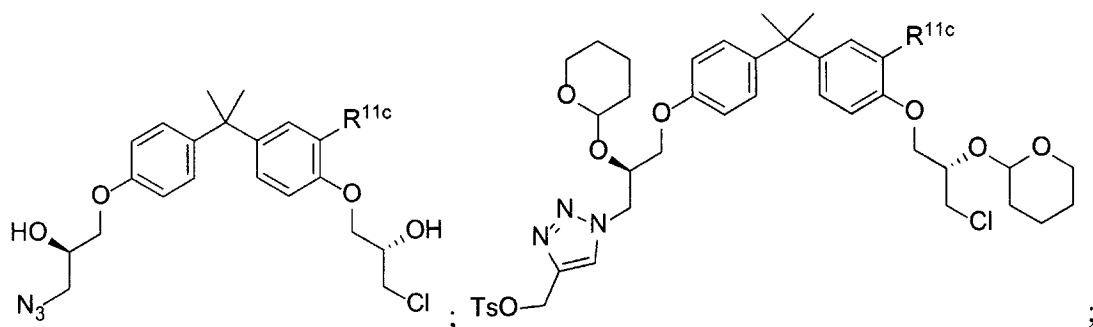
In some embodiments of the compound of Formula 1, the compound has one of the following structures, or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof including all 4 distinct stereoisomers e.g., (*S,R*), (*R,S*), (*R,R*), and (*S,S*):



; wherein R^{11c} is H, Cl, Br, I, F, or C_1-C_6

alkyl.

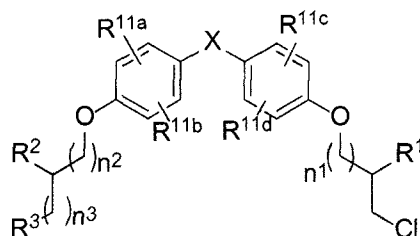
In some embodiments of the compound of Formula I, the compound has one of the following structures, or a pharmaceutically acceptable salt thereof:



; wherein R^{11c} is H, Cl, Br, I, F, or C_1-C_6

alkyl.

In one embodiment, the present invention is directed to a pharmaceutical composition, comprising a compound having a structure of Formula I:



(I)

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-,
 5 -C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-

R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally substituted with one or more R⁶;

R⁵ is each independently H, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

10 R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I, hydroxyl, oxo, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₆-C₁₂ aryl, wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -OS(O)₂-aryl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁸ and R⁹ are each independently H, halogen, -S(O)₀₋₂R⁵, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, aryl, aralkyl, C₁-C₁₀ acyl, or -NR⁵R⁵, or R⁸ and R⁹ can join to form
 15 a unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

R^{11a}, R^{11b}, R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ¹²³I, hydroxyl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl; -OR⁵, -OC(=O)R¹³, C₁-C₁₀ acyl, -S(O)₀₋₂R⁵, -NO₂,
 20 CN, -NH₂, -NHR⁵, or -N(R⁵)₂;

R¹³ is each independently C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

n¹ and n² are each independently 0, 1, or 2; and

n³ is 0, 1, 2, 3, 4, or 5.

In some embodiment, the pharmaceutical composition comprising a compound having a structure of Formula I further comprises a pharmaceutically acceptable carrier. In another embodiment, the pharmaceutical composition comprising a compound having a structure of Formula I further comprises an additional therapeutic agent. In one
5 embodiment, the pharmaceutical composition comprising a compound having a structure of Formula I further comprises a pharmaceutically acceptable carrier and an additional therapeutic agent.

In another embodiment, the pharmaceutical composition comprising a compound having a structure of Formula I further comprises an additional therapeutic agent which is
10 for treating prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, or age-related macular degeneration.

Accordingly, one embodiment comprises the use of the disclosed compounds in
15 combination therapy with one or more currently-used or experimental pharmacological therapies which are utilized for treating the above disease states irrespective of the biological mechanism of action of such pharmacological therapies, including without limitation pharmacological therapies which directly or indirectly inhibit the androgen receptor, pharmacological therapies which are cyto-toxic in nature, and pharmacological
20 therapies which interfere with the biological production or function of androgen (hereinafter, an "additional therapeutic agent"). By "combination therapy" is meant the administration of any one or more of a compound of Formula I with one or more of another therapeutic agent to the same patient such that their pharmacological effects are contemporaneous with one another, or if not contemporaneous, that their effects are
25 synergistic with one another even though dosed sequentially rather than contemporaneously.

Such administration includes without limitation dosing of one or more of a compound of Formula I and one or more of the additional therapeutic agent(s) as separate agents without any comingling prior to dosing, as well as formulations which include one

or more other androgen-blocking therapeutic agents mixed with one or more compound of Formula I as a pre-mixed formulation. Administration of the compound(s) of Formula I in combination with the additional therapeutic agents for treatment of the above disease states also includes dosing by any dosing method including without limitation, intravenous
5 delivery, oral delivery, intra-peritoneal delivery, intra-muscular delivery, or intra-tumoral delivery.

In another aspect of the present disclosure, the one or more of the additional therapeutic agents can be administered to the patient before administration of the compound(s) of Formula I. In another embodiment, the compound(s) of Formula I can be
10 co-administered with one or more of the additional therapeutic agents. In yet another aspect, the one or more additional therapeutic agents can be administered to the patient after administration of the compound(s) of Formula I.

The ratio of the doses of compound(s) of Formula I to that of the one or more additional therapeutic agents can be about 1:1 or can vary, e.g., about 2:1, about 3:1, about
15 4:1, about 5:1, about 6:1, about 7:1, about 8:1, about 9:1, about 10:1, about 1:2, about 1:3, about 1:4, about 1:5, about 1:6, about 1:7, about 1:8, about 1:9, about 1:10, and can be varied accordingly to achieve the optimal therapeutic benefit.

The compound(s) of Formula I that are combined with the one or more additional therapeutic agents for improved treatment of the above disease states can comprise, but are
20 not limited to any compound having a structure of Formula I, including those compounds shown in Tables 1, 2, or 3.

The additional therapeutic agents include without limitation any pharmacological agent which is currently approved by the FDA in the U.S. (or elsewhere by any other regulatory body) for use as pharmacological treatment of any of the above disease states, or
25 which is currently being used experimentally as part of a clinical trial program that relates to the above disease states. Non-limiting examples of the Other Pharmacological Agents comprise, without limitation: the chemical entity known as **ODM-201** (also known as BAY1841788) and related compounds, which appears to bind to the AR and blocks its cellular function, and is currently in clinical development as a treatment for prostate

cancer); the chemical entity known as enzalutamide (4-(3-(4-cyano-3-(trifluoromethyl)phenyl)-5,5-dimethyl-4-oxo-2-thioxoimidazolidin-1-yl)-2-fluoro-N-methylbenzamide) and related compounds, which appears to be a blocker of the androgen receptor (AR) LBD and a FDA-approved treatment for prostate cancer; the chemical entity

5 known as Galeterone and related compounds which appears to be a blocker of the androgen receptor (AR) LBD, and a CYP17 lyase inhibitor, and also appears to decrease overall androgen receptor levels in prostate cancer cells. Galeterone is currently in development as a treatment for prostate cancer; the chemical entity known as ARN-509 (4-[7-[6-cyano-5-(trifluoromethyl)pyridin-3-yl]-8-oxo-6-sulfanylidene-5,7-diazaspiro[3.4]octan-5-yl]-2-

10 fluoro-N-methylbenzamide) and related compounds which appears to be a blocker of the androgen receptor (AR) LBD and is currently in development as a treatment for prostate cancer; the chemical entity known as abiraterone (or CB-7630; (3S,8R,9S,10R,13S,14S)-10,13-dimethyl-17-(pyridin-3-yl) 2,3,4,7,8,9,10,11,12,13,14,15-dodecahydro-1H-cyclopenta[a]phenanthren-3-ol), and related molecules, which appears to block the

15 production of androgen and FDA-approved treatment for prostate cancer; the chemical entity known as bicalutamide (N-[4-cyano-3-(trifluoromethyl)phenyl]-3-[(4-fluorophenyl)sulfonyl]-2-hydroxy-2-methylpropanamide) and related compounds, which appears to be a blocker of the androgen receptor (AR) LBD and which is currently used to treat prostate cancer, the chemical entity known as nilutamide (5,5-dimethyl-3-[4-nitro-3-

20 (trifluoromethyl)phenyl] imidazolidine-2,4-dione) and related compounds, which appears to be a blocker of the AR LBD and which is currently used to treat prostate cancer, the chemical entity known as flutamide (2-methyl-N-[4-nitro-3-(trifluoromethyl)phenyl]-propanamide) and related compounds, which appears to be a blocker of the androgen receptor (AR) LBD and which is currently used to treat prostate cancer, the chemical

25 entities known as cyproterone acetate (6-chloro-1 β ,2 β -dihydro-17-hydroxy-3'*H*-cyclopropa[1,2]pregna-4,6-diene-3,20-dione) and related compounds, which appears to be a blocker of the androgen receptor (AR) LBD and which is currently used to treat prostate cancer, the chemical entity known as docetaxel (Taxotere; 1,7 β ,10 β -trihydroxy-9-oxo-5 β ,20-epoxytax-11-ene-2 α ,4,13 α -triyl 4-acetate 2-benzoate 13-[(2R,3S)-3-[(tert-butoxycarbonyl)amino]-2-hydroxy-3-phenylpropanoate})) and related compounds, which

30

appears to be a cytotoxic antimicrotubule agent and is currently used in combination with prednisone to treat prostate cancer, the chemical entity known as Bevacizumab (Avastin), a monoclonal antibody that recognizes and blocks vascular endothelial growth factor A (VEGF-A) and can be used to treat prostate cancer, the chemical entity known as OSU-
5 HDAC42 ((S)-(+)-N-hydroxy-4-(3-methyl-2-phenylbutyrylamino)-benzamide), and related compounds, which appears to act as a histone deacetylase inhibitor, and is currently being developed as a treatment for prostate cancer, the chemical entity known as VITAXIN which appears to be a monoclonal antibody against the vascular integrin $\alpha\beta3$ to prevent angiogenesis, and which can be used to treat prostate cancer, the chemical entity known as
10 sunitumib (N-(2-diethylaminoethyl)-5-[(Z)-(5-fluoro-2-oxo-1*H*-indol-3-ylidene)methyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxamide) and related compounds, which appears to inhibit multiple receptor tyrosine kinases (RTKs) and can be used for treatment of prostate cancer, the chemical entity known as ZD-4054 (N-(3-Methoxy-5-methylpyrazin-2-yl)-2-[4-(1,3,4-oxadiazol-2-yl)phenyl]pyridin-3-sulfonamid) and related compounds, which appears to
15 block the edta receptor and which can be used for treatment of prostate cancer; the chemical entity known as Cabazitaxel (XRP-6258), and related compounds, which appears to be a cytotoxic microtubule inhibitor, and which is currently used to treat prostate cancer; the chemical entity known as MDX-010 (Ipilimumab), a fully human monoclonal antibody that binds to and blocks the activity of CTLA-4 which is currently in development as an
20 immunotherapeutic agent for treatment of prostate cancer; the chemical entity known as OGX 427 which appears to target HSP27 as an antisense agent, and which is currently in development for treatment of prostate cancer; the chemical entity known as OGX 011 which appears to target clusterin as an antisense agent, and which is currently in development as a treatment for prostate cancer; the chemical entity known as finasteride
25 (Proscar, Propecia; *N*-(1,1-dimethylethyl)-3-oxo-(5 α ,17 β)-4-azaandrost-1-ene-17-carboxamide), and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone, and can be used to treat prostate cancer; the chemical entity known as dutasteride (Avodart; 5 α , 17 β)-*N*-{2,5 bis(trifluoromethyl)phenyl}-3-oxo-4-azaandrost-1-ene-17-carboxamide) and related molecules, which appears
30 to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone, and can be

used in the treatment of prostate cancer; the chemical entity known as turosteride ((4aR,4bS,6aS,7S,9aS,9bS,11aR)-1,4a,6a-trimethyl-2-oxo-N-(propan-2-yl)-N-(propan-2-ylcarbamoyl)hexadecahydro-1H-indeno[5,4-f]quinoline-7-carboxamide), and related molecules, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and can be used in the treatment of prostate cancer; the chemical entity known as bexlosteride (LY-191,704; (4aS,10bR)-8-chloro-4-methyl-1,2,4a,5,6,10b-hexahydrobenzo[f]quinolin-3-one), and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and can be used in the treatment of prostate cancer; the chemical entity known as izonsteride (LY-320,236; (4aR,10bR)-8-[(4-ethyl-1,3-benzothiazol-2-yl)sulfanyl]-4,10b-dimethyl-1,4,4a,5,6,10b-hexahydrobenzo[f]quinolin-3(2H)-one) and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and can be used for the treatment of prostate cancer; the chemical entity known as FCE 28260 and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and can be used for the treatment of prostate cancer; the chemical entity known as SKF105,111, and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and can be used for treatment of prostate cancer.

Accordingly, in some embodiments, the pharmaceutical composition comprising a compound having a structure of Formula I further comprises an additional therapeutic agent selected from the group consisting of enzalutamide, Galeterone, ARN-509; abiraterone, bicalutamide, nilutamide, flutamide, cyproterone acetate, docetaxel, Bevacizumab (Avastin), OSU-HDAC42, VITAXIN, sunitumib, ZD-4054, Cabazitaxel (XRP-6258), MDX-010 (Ipilimumab), OGX 427, OGX 011, finasteride, dutasteride, turosteride, bexlosteride, izonsteride, FCE 28260, SKF105,111, ODM-201, radium 223, or related compounds thereof.

In some embodiments, compounds of Formula I which result in unstable structures and/or unsatisfied valences are not included within the scope of the invention.

In another embodiment, the present disclosure provides a pharmaceutical composition comprising any of the foregoing compounds of Formula I and a pharmaceutically acceptable carrier.

Compounds as described herein can be in the free form or in the form of a salt thereof. In some embodiments, compounds as described herein can be in the form of a pharmaceutically acceptable salt, which are known in the art (Berge et al., *J. Pharm. Sci.* 1977, 66, 1). Pharmaceutically acceptable salt as used herein includes, for example, salts that have the desired pharmacological activity of the parent compound (salts which retain the biological effectiveness and/or properties of the parent compound and which are not biologically and/or otherwise undesirable). Compounds as described herein having one or more functional groups capable of forming a salt can be, for example, formed as a pharmaceutically acceptable salt. Compounds containing one or more basic functional groups can be capable of forming a pharmaceutically acceptable salt with, for example, a pharmaceutically acceptable organic or inorganic acid. Pharmaceutically acceptable salts can be derived from, for example, and without limitation, acetic acid, adipic acid, alginic acid, aspartic acid, ascorbic acid, benzoic acid, benzenesulfonic acid, butyric acid, cinnamic acid, citric acid, camphoric acid, camphorsulfonic acid, cyclopentanepropionic acid, diethylacetic acid, digluconic acid, dodecylsulfonic acid, ethanesulfonic acid, formic acid, fumaric acid, glucoheptanoic acid, gluconic acid, glycerophosphoric acid, glycolic acid, hemisulfonic acid, heptanoic acid, hexanoic acid, hydrochloric acid, hydrobromic acid, hydriodic acid, 2-hydroxyethanesulfonic acid, isonicotinic acid, lactic acid, malic acid, maleic acid, malonic acid, mandelic acid, methanesulfonic acid, 2-naphthalenesulfonic acid, naphthalenedisulphonic acid, p-toluenesulfonic acid, nicotinic acid, nitric acid, oxalic acid, pamoic acid, pectinic acid, 3-phenylpropionic acid, phosphoric acid, picric acid, pimelic acid, pivalic acid, propionic acid, pyruvic acid, salicylic acid, succinic acid, sulfuric acid, sulfamic acid, tartaric acid, thiocyanic acid or undecanoic acid. Compounds containing one or more acidic functional groups can be capable of forming pharmaceutically acceptable salts with a pharmaceutically acceptable base, for example, and without limitation, inorganic bases based on alkaline metals or alkaline earth metals or organic bases such as primary amine compounds, secondary amine compounds, tertiary

amine compounds, quaternary amine compounds, substituted amines, naturally occurring substituted amines, cyclic amines or basic ion-exchange resins. Pharmaceutically acceptable salts can be derived from, for example, and without limitation, a hydroxide, carbonate, or bicarbonate of a pharmaceutically acceptable metal cation such as ammonium, sodium, potassium, lithium, calcium, magnesium, iron, zinc, copper, manganese or aluminum, ammonia, benzathine, meglumine, methylamine, dimethylamine, trimethylamine, ethylamine, diethylamine, triethylamine, isopropylamine, tripropylamine, tributylamine, ethanolamine, diethanolamine, 2-dimethylaminoethanol, 2-diethylaminoethanol, dicyclohexylamine, lysine, arginine, histidine, caffeine, hydrabamine, choline, betaine, ethylenediamine, glucosamine, glucamine, methylglucamine, theobromine, purines, piperazine, piperidine, procaine, *N*-ethylpiperidine, theobromine, tetramethylammonium compounds, tetraethylammonium compounds, pyridine, *N,N*-dimethylaniline, *N*-methylpiperidine, morpholine, *N*-methylmorpholine, *N*-ethylmorpholine, dicyclohexylamine, dibenzylamine, *N,N*-dibenzylphenethylamine, 1-ephedrine, *N,N*-dibenzylethylenediamine or polyamine resins. In some embodiments, compounds as described herein can contain both acidic and basic groups and can be in the form of inner salts or zwitterions, for example, and without limitation, betaines. Salts as described herein can be prepared by conventional processes known to a person skilled in the art, for example, and without limitation, by reacting the free form with an organic acid or inorganic acid or base, or by anion exchange or cation exchange from other salts. Those skilled in the art will appreciate that preparation of salts can occur *in situ* during isolation and purification of the compounds or preparation of salts can occur by separately reacting an isolated and purified compound.

In some embodiments, compounds and all different forms thereof (e.g. free forms, salts, polymorphs, isomeric forms) as described herein can be in the solvent addition form, for example, solvates. Solvates contain either stoichiometric or non-stoichiometric amounts of a solvent in physical association the compound or salt thereof. The solvent can be, for example, and without limitation, a pharmaceutically acceptable solvent. For example, hydrates are formed when the solvent is water or alcoholates are formed when the solvent is an alcohol.

In some embodiments, compounds and all different forms thereof (e.g. free forms, salts, solvates, isomeric forms) as described herein can include crystalline and amorphous forms, for example, polymorphs, pseudopolymorphs, conformational polymorphs, amorphous forms, or a combination thereof. Polymorphs include different crystal packing
5 arrangements of the same elemental composition of a compound. Polymorphs usually have different X-ray diffraction patterns, infrared spectra, melting points, density, hardness, crystal shape, optical and electrical properties, stability and/or solubility. Those skilled in the art will appreciate that various factors including recrystallization solvent, rate of crystallization and storage temperature can cause a single crystal form to dominate.

10 In some embodiments, compounds and all different forms thereof (e.g. free forms, salts, solvates, polymorphs) as described herein include isomers such as geometrical isomers, optical isomers based on asymmetric carbon, stereoisomers, tautomers, individual enantiomers, individual diastereomers, racemates, diastereomeric mixtures and combinations thereof, and are not limited by the description of the formula illustrated for
15 the sake of convenience.

III. Methods

The present compounds find use in any number of methods. For example, in some embodiments the compounds are useful in methods for modulating androgen receptor
20 (AR). Accordingly, in one embodiment, the present disclosure provides the use of any one of the foregoing compounds of Formula I for modulating androgen receptor (AR) activity. For example in some embodiments, modulating androgen receptor (AR) activity is in a mammalian cell. Modulating androgen receptor (AR) can be in a subject in need thereof (e.g., a mammalian subject) and for treatment of any of the described conditions or
25 diseases.

In other embodiments, modulating androgen receptor (AR) activity is for treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar
30 muscular atrophy, age related macular degeneration, and combinations thereof. For

example in some embodiments, the indication is prostate cancer. In other embodiments, the prostate cancer is castration resistant prostate cancer (also referred to as hormone refractory, androgen-independent, androgen deprivation resistant, androgen ablation resistant, androgen depletion-independent, castration-recurrent, anti-androgen-recurrent).

- 5 While in other embodiments, the prostate cancer is androgen dependent prostate cancer. In other embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease.

In some embodiments, compounds as described herein can be administered to a subject. In one embodiment, the present invention is directed to a method of treating castration resistant prostate cancer comprising administering a pharmaceutical composition
10 comprising a compound having a structure of Formula I. In some embodiments, the present invention is directed to a method of treating androgen-dependent prostate cancer comprising administering a pharmaceutical composition comprising a compound having a structure of Formula I. In other embodiments, the present invention is directed to a method of treating androgen-independent prostate cancer comprising administering a
15 pharmaceutical composition comprising a compound having a structure of Formula I.

In other embodiments, the present disclosure provides a method of modulating androgen receptor (AR) activity, the method comprising administering any one of the foregoing compounds of Formula I, pharmaceutically acceptable salt thereof, or pharmaceutical composition of Formula I as described herein (including compositions
20 comprising a compound of Formula I and an additional therapeutic agent), to a subject (e.g., mammal) in need thereof. In some embodiments, modulating androgen receptor (AR) activity is in a mammalian cell. In other embodiments, modulating androgen receptor (AR) activity is in a mammal. In one embodiment, modulating androgen receptor (AR) activity is in a human.

25 The modulating androgen receptor (AR) activity can be for inhibiting AR N-terminal domain activity. The modulating androgen receptor (AR) activity can be for inhibiting androgen receptor (AR) activity. The modulating can be *in vivo*. The modulating androgen receptor (AR) activity can be for treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial
30 cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary

disease, precocious puberty, spinal and bulbar muscular atrophy (e.g., Kennedy's disease), and age related macular degeneration. The indication can be prostate cancer. The prostate cancer can be castration-resistant prostate cancer. The prostate cancer can be androgen dependent prostate cancer.

5 In accordance with another embodiment, there is provided a use of the compounds of Formula I, pharmaceutically acceptable salt thereof, or pharmaceutical composition of Formula I as described herein for preparation of a medicament for modulating androgen receptor (AR).

10 Alternatively, in one embodiment, a method of modulating androgen receptor activity, comprising administering Formula I, pharmaceutically acceptable salt thereof, or pharmaceutical composition of Formula I as described herein, is provided. In some embodiments, the administration can be to a mammal. In other embodiments, the administering can be to a mammal in need thereof and in an effective amount for the treatment of at least one indication selected from the group consisting of: prostate cancer,
15 breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy (e.g., Kennedy's disease), age related macular degeneration, and combinations thereof.

20 As noted above, the presently disclosed compounds can find utility in a number of medical imaging applications, including imaging of the prostate. Many currently available imaging agents tend to accumulate in the bladder, which decreases their effectiveness as imaging tools specifically for the prostate. While not wishing to be bound by theory, the present applicants believe the disclosed compounds are unexpectedly effective for imaging of the prostate due to their ability to accumulate in the prostate, rather than the bladder,
25 allowing the prostate gland to be seen. Accordingly, the compounds can be used in methods for imaging the prostate, for example to image benign prostate diseases. In other embodiments, the compounds can be used in methods to image cancerous prostate diseases, such as tumors of the prostate.

30 Androgen ablation therapy causes a temporary reduction in prostate cancer tumor burden, but the malignancy will begin to grow again in the absence of testicular androgens

to form castrate resistant prostate cancer (CRPC). A rising titer of serum prostate-specific antigen (PSA) after androgen ablation therapy indicates biochemical failure, the emergence of CRPC, and re-initiation of an androgen receptor (AR) transcription program. Most patients succumb to CRPC within two years of biochemical failure.

5 AR is a transcription factor and a validated target for prostate cancer therapy. Current therapies include androgen ablation and administration of antiandrogens. Most CRPC is suspected to be AR-dependent. AR has distinct functional domains that include the C-terminus ligand-binding domain (LBD), a DNA-binding domain (DBD), and an amino-terminal domain (NTD). AR NTD contains the activation function- 1 (AF-1) that
10 contributes most of the activity to the AR. Recently, splice variants of the AR that lack the LBD have been reported in prostate cancer cell lines (VCaP and 22Rv1), and in CRPC tissues. To date more than 20 splice variants of AR have been detected. Splice variants V7 and V567es are clinically relevant with levels of expression correlated to poor survival and CRPC. AR V567es is solely expressed in 20% of metastases. Abiraterone resistance is
15 associated with expression of AR splice variants. Enzalutamide also increases levels of expression of these constitutively active AR splice variants. These splice variants lack LBD and thereby would not be inhibited by current therapies that target the AR LBD such as antiandrogens or androgen ablation therapy. A single patient with advanced prostate cancer can have many lesions throughout the body and skeleton and each tumor can have differing
20 levels of expression of AR.

 Biopsy of metastatic tumors in a patient to determine AR species is neither widely accessible nor feasible to sample tumors in a patient that can have multiple metastases. Thus it is essential to develop approaches to detect the expression of all AR species for the molecular classification of tumors based on the level and extent of expression of AR splice
25 variants, or other AR species that cannot be detected using an imaging agent that interacts with the LBD, to identify patients with potentially aggressive disease and poor prognosis, or to identify patients that will not respond to hormone therapies that target the AR LBD. Accordingly, certain embodiments of the present invention provide a AR NTD-targeted molecular imaging probe (*e.g.*, compound of formula I) which can be used to monitor
30 response to therapy and provide insight into the role of AR in resistance mechanisms.

One current approach to image AR in prostate cancer uses positron emission tomography (PET) with 16β -[^{18}F]-fluoro-5 α dihydrotestosterone (^{18}F -FDHT) that binds to AR LBD. Unfortunately this imaging agent cannot detect splice variants lacking LBD. In some embodiments, the invention employs sequential imaging with ^{18}F -FDHT to detect
5 full-length AR and gamma radiation emitting probes to specifically detect the AR NTD which would be the sum of both full-length AR and variant AR. In other embodiments, the invention employs sequential imaging with two different PET imaging agents to detect only full-length AR and another to specifically detect the AR NTD which would be the sum of both full-length AR and variant AR. Together these data reveal patients with tumors
10 that express variant AR (NTD of variant plus full-length AR detected with NTD isotope minus full-length AR detected with ^{18}F -FDHT). By using sequential imaging, a discordant distribution or discordant level of uptake between ^{18}F -FDHT and a radiolabeled compound of this invention (*i.e.*, compound of Formula I) indicates the presence of overexpression of splice variants lacking the LBD.

15 As described above, radioactive ^{18}F labeled compounds have found use as imaging agents not only to image AR in prostate cancer but for imaging various organs and various tumors. Similarly, radioactive ^{123}I labeled compounds have been known for the use as imaging agents. In one embodiment, the compounds of the present disclosure comprise at least one ^{18}F and/or ^{123}I .

20 In one embodiment, the present invention is directed to a method of imaging cancer by administering a compound having a structure of Formula I to a subject. In another embodiment, the present invention is directed to a method of imaging cancer by administering a pharmaceutical composition comprising a compound having a structure of Formula I to a subject.

25 In some embodiment, the method of imaging cancer by administering a compound having a structure of Formula I or a pharmaceutical composition comprising Formula I to a subject and detecting the presence or absence of cancer by use of SPECT or PET. In other embodiments, the method of imaging cancer by administering a compound having a structure of Formula I to a subject and the method identifies the presence or absence of a
30 tumor. In one embodiment, the method of imaging cancer by administering a compound

having a structure of Formula I to a subject and the method identifies the location of a tumor. In one embodiment, the method of imaging cancer by administering a compound having a structure of Formula I to a subject and the method identifies a presence of a prostate cancer. In other embodiments, the prostate cancer is androgen-dependent prostate cancer. In some embodiments, the subject is a mammal such as a human.

In some other embodiments, the method is useful for detecting the presence of AR splice variants or other AR species that cannot be detected by imaging agents that interact with the AR LBD (i.e., mutations, truncations). Without wishing to be bound by any particular theory, since the present compounds bind to the AR N-terminal domain (NTD), even mutants or variants which lack the AR LBD can be imaged employing the present compounds. Thus, the present methods can be useful for detecting AR species, including mutants and variants, which lack the LBD or have LBD mutations, but do comprise the AR NTD. In other embodiments the method detects the presence or overexpression of AR splice variants lacking the ligand binding domain. For example, the method can include sequential imaging with ^{18}F -FDHT and a compound of the invention and a discordant distribution or discordant level of uptake between ^{18}F -FDHT and the compound of the invention indicates the presence or overexpression of splice variants lacking the ligand binding domain.

In other embodiments, the compounds of the invention are used in single photon emission computed tomography methods to monitor a patient's response to therapy. In other embodiments, the methods comprise use of a compound of the invention to detect the AR NTD.

In accordance with a further embodiment, the method of imaging a cancer is by administering a compound as described anywhere herein. In one embodiment, the method of imaging a cancer is by administering a compound represented in Table 2, Table 3 or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof. In some embodiments, the imaging is in a mammalian cell. In one embodiment, the imaging is in a mammal. In other embodiments, the imaging is in a human patient.

The administering and imaging can be to a mammal in need of diagnosis of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian

cancer, endometrial cancer, salivary gland carcinoma, benign prostatic hyperplasia, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy (*e.g.*, Kennedy's disease), and age-related macular degeneration. The imaging can be for imaging AR splice variants, mutants or other AR species which comprise the AR NTD.

In some embodiments, the compounds as described herein or pharmaceutically acceptable salts thereof can be used for imaging and diagnosis of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, benign prostatic hyperplasia, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, and age-related macular degeneration. In some embodiments, the compounds as described herein or acceptable salts thereof above can be used in the preparation of a medicament or a composition for imaging the prostate, for example for imaging benign prostate conditions or for imaging prostate cancer in a subject in need of such imaging (for example for diagnosis and/or location of prostate tumors).

In some embodiments, pharmaceutical compositions useful in modulating androgen receptor (AR) activity or useful for imaging, in accordance with this invention can comprise a salt of such a compound, preferably a pharmaceutically or physiologically acceptable salt. Pharmaceutical preparations will typically comprise one or more carriers, excipients or diluents acceptable for the mode of administration of the preparation, be it by injection, inhalation, topical administration, lavage, or other modes suitable for the selected treatment. Suitable carriers, excipients or diluents are those known in the art for use in such modes of administration.

Suitable pharmaceutical compositions can be formulated by means known in the art and their mode of administration and dose determined by the skilled practitioner. For parenteral administration, a compound can be dissolved in sterile water or saline or a pharmaceutically acceptable vehicle used for administration of non-water soluble compounds such as those used for vitamin K. For enteral administration, the compound can be administered in a tablet, capsule or dissolved in liquid form. The tablet or capsule can be enteric coated, or in a formulation for sustained release. Many suitable formulations are

known, including, polymeric or protein microparticles encapsulating a compound to be released, ointments, pastes, gels, hydrogels, or solutions which can be used topically or locally to administer a compound. A sustained release patch or implant can be employed to provide release over a prolonged period of time. Many techniques known to one of skill in the art are described in *Remington: the Science & Practice of Pharmacy* by Alfonso Gennaro, 20th ed., Lippencott Williams & Wilkins, (2000). Formulations for parenteral administration can, for example, contain excipients, polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, or hydrogenated naphthalenes. Biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxyethylene-polyoxypropylene copolymers can be used to control the release of the compounds. Other potentially useful parenteral delivery systems for modulatory compounds include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation can contain excipients, for example, lactose, or can be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or can be oily solutions for administration in the form of nasal drops, or as a gel.

In another aspect, the present disclosure provides a pharmaceutical composition comprising a compound as described herein, and an additional therapeutic agent and/or a pharmaceutically acceptable carrier. In some embodiments, the additional therapeutic agent is for treating prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy or age related macular degeneration. In other embodiments, the additional therapeutic agent is enzalutamide, galeterone, ARN-509, ODN-201 abiraterone, bicalutamide, nilutamide, flutamide, cyproterone acetate, docetaxel, Bevacizumab (Avastin), OSU-HDAC42, VITAXIN, sunitumib, ZD-4054, Cabazitaxel (XRP-6258), MDX-010 (Ipilimumab), OGX 427, OGX 011, finasteride, dutasteride, turosteride, bexlosteride, izonsteride, FCE 28260, SKF105,111 ODM-201, or related compounds thereof.

Compounds described herein can also be used in assays and for research purposes. Definitions used include ligand dependent activation of the androgen receptor (AR) by

androgens such as dihydrotestosterone (DHT) or the synthetic androgen (R1881) used for research purposes. Ligand-independent activation of the androgen receptor (AR) refers to transactivation of the full length androgen receptor (AR) in the absence of androgen (ligand) by, for example, stimulation of the cAMP dependent protein kinase (PKA) pathway with forskolin (FSK). Some compounds and compositions of this invention can inhibit both FSK and androgen (e.g. R1881, a synthetic androgen) induction of ARE luciferase (ARE-luc). Constitutive activity of the androgen receptor (AR) refers to splice variants lacking the androgen receptor (AR) ligand-binding domain. Such compounds can block a mechanism that is common to both ligand dependent and ligand independent activation of the androgen receptor (AR), as well as constitutively active splice variants of the androgen receptor (AR) that lack ligand-binding domain. This could involve any step in activation of the androgen receptor (AR) including dissociation of heatshock proteins, essential posttranslational modifications (e.g., acetylation, phosphorylation), nuclear translocation, protein-protein interactions, formation of the transcriptional complex, release of co repressors, and/or increased degradation. Some compounds and compositions of this invention can inhibit ligand-only activity and can interfere with a mechanism specific to ligand dependent activation (e.g., accessibility of the ligand binding domain (LBD) to androgen). Numerous disorders in addition to prostate cancer involve the androgen axis (e.g., acne, hirsutism, alopecia, benign prostatic hyperplasia) and compounds interfering with this mechanism can be used to treat such conditions. Some compounds and compositions of this invention can only inhibit FSK induction and can be specific inhibitors to ligand independent activation of the androgen receptor (AR). These compounds and compositions can interfere with the cascade of events that normally occur with FSK and/or PKA activity or any downstream effects that can play a role on the androgen receptor (AR) (e.g. FSK increases MAPK activity which has a potent effect on androgen receptor (AR) activity). Examples can include an inhibitor of cAMP and or PKA or other kinases. Some compounds and compositions of this invention can induce basal levels of activity of the AR (no androgen or stimulation of the PKA pathway). Some compounds and compositions of this invention can increase induction by R1881 or FSK. Such compounds and compositions can stimulate transcription or transactivation of the AR.

Some compounds and compositions of this invention can inhibit activity of the androgen receptor. Interleukin 6 (IL 6) also causes ligand independent activation of the androgen receptor (AR) in LNCaP cells and can be used in addition to FSK.

Compounds or pharmaceutical compositions in accordance with this invention or
5 for use in this invention can be administered by means of a medical device or appliance such as an implant, graft, prosthesis, stent, etc. Also, implants can be devised which are intended to contain and release such compounds or compositions. An example would be an implant made of a polymeric material adapted to release the compound over a period of time.

10 It is to be noted that dosage values can vary with the exact imaging protocol. For any particular subject, specific dosage regimens can be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions. Dosage ranges set forth herein are exemplary only and do not limit the dosage ranges that can be selected by medical practitioners. The
15 amount of active compound(s) in the composition can vary according to factors such as the disease state, age, sex, and weight of the subject. Dosage regimens can be adjusted to provide the optimum imaging result. For example, a single 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 imaging results. It can be advantageous to
20 formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage.

In general, compounds of the invention should be used without causing substantial toxicity. Toxicity of the compounds of the invention can be determined using standard techniques, for example, by testing in cell cultures or experimental animals and
25 determining the therapeutic index, *i.e.*, the ratio between the LD50 (the dose lethal to 50% of the population) and the LD100 (the dose lethal to 100% of the population). In some circumstances, such as in severe disease conditions, substantial excesses of the compositions can be administered for therapeutic effects. Some compounds of this invention can be toxic at some concentrations. Titration studies can be used to determine
30 toxic and non-toxic concentrations. Toxicity can be evaluated by examining a particular

compound's or composition's specificity across cell lines using PC3 or DU145 cells as possible negative controls since these cells do not express functional AR. Animal studies can be used to provide an indication if the compound has any effects on other tissues. Systemic therapy that targets the AR will not likely cause major problems to other tissues
5 since antiandrogens and androgen insensitivity syndrome are not fatal.

Compounds for use in the present invention can be obtained from medical sources or modified using known methodologies from naturally occurring compounds. In addition, methods of preparing or synthesizing compounds of the present invention will be understood by a person of skill in the art having reference to known chemical synthesis
10 principles. For example, Auzou *et al* 1974 *European Journal of Medicinal Chemistry* 9(5), 548-554 describes suitable synthetic procedures that can be considered and suitably adapted for preparing compounds of any one of the compounds of structure (I) as set out above. Other references that can be helpful include: Debasish Das, Jyh-Fu Lee and Soofin Cheng "Sulfonic acid functionalized mesoporous MCM-41 silica as a convenient catalyst
15 for Bisphenol-A synthesis" *Chemical Communications*, (2001) 2178-2179; US Patent 2571217 Davis, Orris L.; Knight, Horace S.; Skinner, John R. (Shell Development Co.) "Halohydrin ethers of phenols." (1951); and Rokicki, G.; Pawlicki, J.; Kuran, W. "Reactions of 4-chloromethyl-1,3-dioxolan-2-one with phenols as a new route to polyols and cyclic carbonates." *Journal fuer Praktische Chemie (Leipzig)* (1985) 327, 718-722.

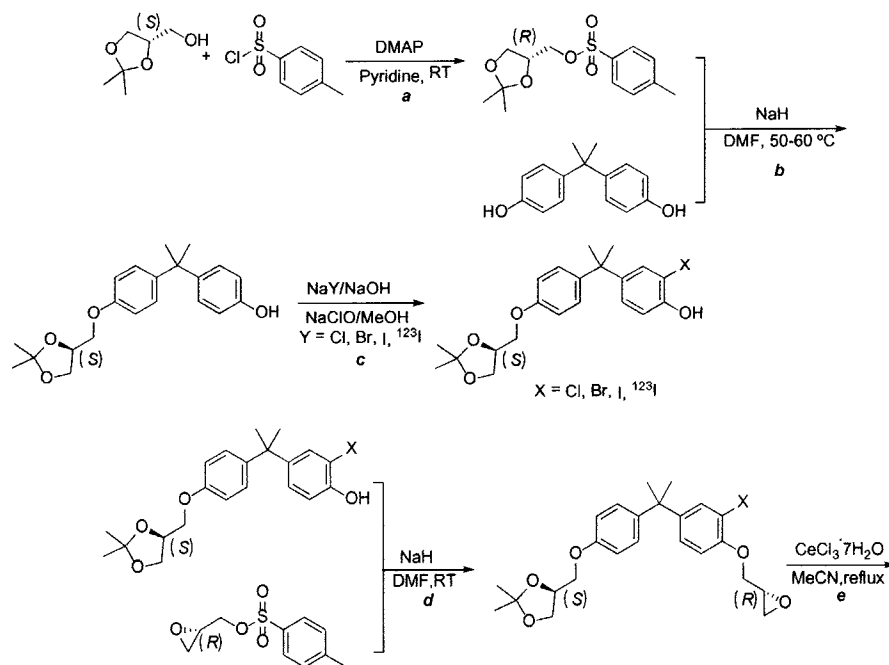
20 In some embodiments, compounds and all different forms thereof as described herein can be used, for example, and without limitation, in combination with other treatment methods for at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal
25 and bulbar muscular atrophy, and age related macular degeneration. For example, compounds and all their different forms as described herein can be used as neoadjuvant (prior), adjunctive (during), and/or adjuvant (after) therapy with surgery, radiation (brachytherapy or external beam), or other therapies (eg. HIFU), and in combination with chemotherapies, androgen ablation, antiandrogens or any other therapeutic approach.

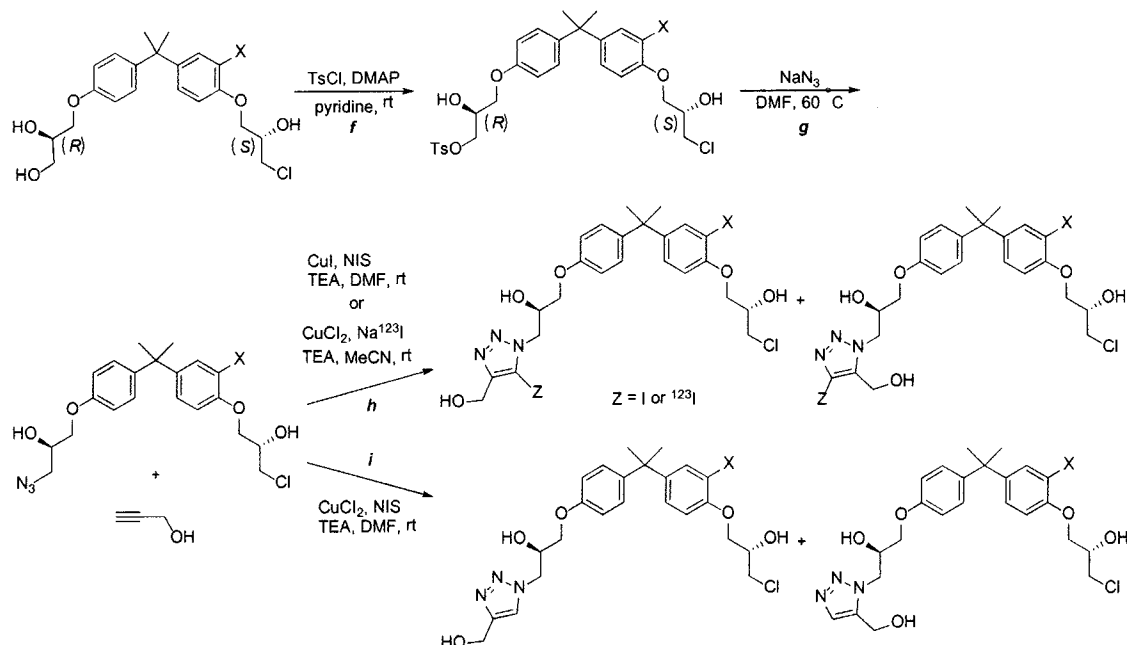
In an exemplary embodiment for imaging the prostate, a dose of the disclosed compounds in solution (typically 5 to 10 millicuries or 200 to 400 MBq) is typically injected rapidly into a saline drip running into a vein, in a patient. Then, the patient is placed in the SPECT for a series of one or more scans which can take from 20 minutes to
 5 as long as an hour (often, only about one quarter of the body length can be imaged at a time). Methods for SPECT scanning are well known in the art.

The compounds described herein can be used for in vivo or in vitro research uses (i.e. non-clinical) to investigate the mechanisms of orphan and nuclear receptors (including steroid receptors such as androgen receptor (AR)). Furthermore, these compounds can be
 10 used individually or as part of a kit for in vivo or in vitro research to investigate signal transduction pathways and/or the activation of orphan and nuclear receptors using recombinant proteins, cells maintained in culture, and/or animal models.

For example, exemplary compounds of the present invention can be prepared with
 15 reference to the following General Reaction Scheme I:

General Reaction Scheme I





It should be noted that, although General Reaction Scheme I depicts a certain stereochemistry in the products, one skilled in the art would recognize that other stereoisomers can be synthesized by using starting materials of different stereochemistry, such as for step *a* and step *d*.

Referring to General Reaction Scheme I, (*S*)-(2,2-dimethyl-1,3-dioxolan-4-yl)methanol can be tosylated under basic conditions as shown in step *a*. In step *b*, tosyl group (*p*-toulenesulfonate) can be displaced with bisphenol derivatives under basic conditions. Optionally, in step *c*, the bisphenol derivative is halogenated on the phenyl ring. In step *d*, the unreacted phenol portion undergoes another elimination reaction to afford a stereospecific bisphenol derivative with an epoxide on one side and a protected diol on the other. In step *e*, epoxide is opened using CeCl₃·7H₂O and the diol is deprotected *in situ*. In step *f*, primary alcohol is selectively tosylated then in step *g*, the tosyl group is displaced with an azide using sodium azide. Lastly, Click chemistry provides triazole containing products in steps *h* and *i*. Tosyl leaving groups have been found to be particularly useful in the synthesis outlined in General Scheme I.

The radioactive iodine moiety (¹²³I) can be installed by step *c* with an appropriate iodinating reagent, for example Na¹²³I and a suitable oxidant (e.g., NaClO). It should be noted that, although General Reaction Scheme I depicts iodination at only one position,

other compounds of structure (I) with ^{123}I at different positions and/or multiple ^{123}I substitutions can be prepared according to analogous methods known to those of ordinary skill in the art. Further, ^{123}I atoms can be introduced via any number of reagents, and iodination is not limited to those methods depicted or described above. Methods for such
5 iodination are well known in the art. Methodologies for preparation of specific exemplary compounds of structure I are described in more detail in the following examples.

One skilled in the art will recognize that variations to the order of the steps and reagents discussed in reference to the above General Synthetic Schemes I are possible. For example, epoxidation can precede dioxalone formation.

10 In addition, protecting group strategies can be employed for preparation of the compounds disclosed herein. Such strategies are well known to those of skill in the art. Exemplary protecting groups and related strategies are disclosed in Greene's Protective Groups in Organic Synthesis, Wiley-InterScience; 4 edition (October 30, 2006), which is hereby incorporated by reference in its entirety. In certain embodiments, a protecting
15 group is used to mask an alcohol moiety while performing other chemical transformations. After removal of the protecting group, the free hydroxyl is obtained. Such protecting groups and strategies are well known in the art.

Various alternative embodiments and examples of the invention are described herein. These embodiments and examples are illustrative and should not be construed as
20 limiting the scope of the invention. The following examples are provided for purposes of illustration, not limitation.

EXAMPLES

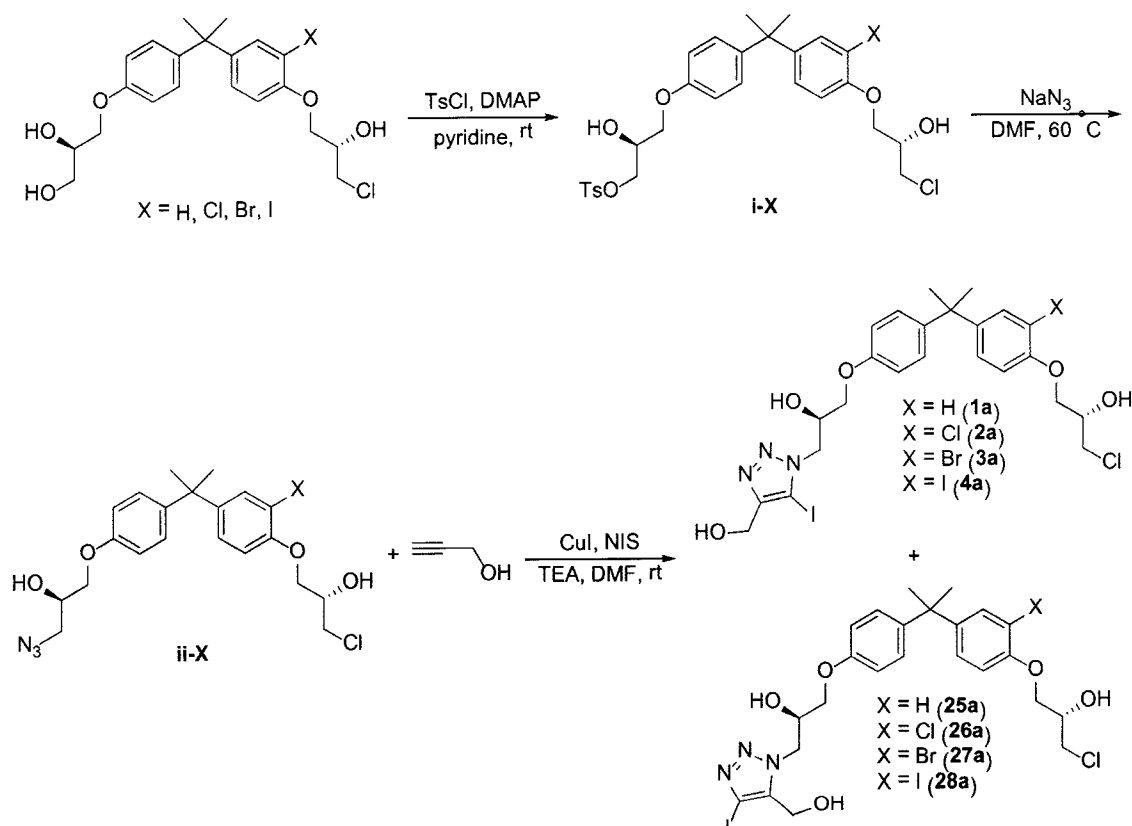
All non-aqueous reactions were performed in flame-dried round bottomed flasks. The flasks were fitted with rubber septa and reactions were conducted under a positive
25 pressure of argon unless otherwise specified. Stainless steel syringes were used to transfer air- and moisture-sensitive liquids. Flash column chromatography was performed as described by Still et al. (Still, W. C.; Kahn, M.; Mitra, A. *J. Org. Chem.* **1978**, *43*, 2923) using 230-400 mesh silica gel. Thin-layer chromatography was performed using aluminum plates pre-coated with 0.25 mm 230-400 mesh silica gel impregnated with a fluorescent

indicator (254 nm). Thin-layer chromatography plates were visualized by exposure to ultraviolet light and a "Seebach" staining solution (700 mL water, 10.5 g Cerium (IV) sulphate tetrahydrate, 15.0 g molybdate phosphoric acid, 17.5 g sulphuric acid) followed by heating (~1 min) with a heating gun (~250 °C). Organic solutions were concentrated on
5 Büchi R-114 rotatory evaporators at reduced pressure (15-30 torr, house vacuum) at 25-40 °C.

Commercial reagents and solvents were used as received. All solvents used for extraction and chromatography were HPLC grade. Normal-phase Si gel Sep paksTM were purchased from waters, Inc. Thin-layer chromatography plates were Kieselgel 60F₂₅₄. All
10 synthetic reagents were purchased from Sigma Aldrich and Fisher Scientific Canada.

EXAMPLE 1

SYNTHESIS OF (*S*)-1-(4-(2-(3-BROMO-4-((*R*)-3-CHLORO-2-HYDROXYPROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)-3-(4-(HYDROXYMETHYL)-5-iodo-1*H*-1,2,3-triazol-1-yl)PROPAN-2-OL (**1a**) AND (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(5-(HYDROXYMETHYL)-4-iodo-1*H*-1,2,3-triazol-1-yl)PROPOXY)PHENYL)PROPAN-2-yl)PHENOXY)PROPAN-2-OL (**25a**)

**Compound i-H (X = H)**

- 10 *p*-Toluenesulfonyl chloride (~1.5 equiv) was added portionwise over a period of approximately 10 min to a solution of (*R*)-3-(4-(2-(4-((*S*)-3-chloro-2-hydroxypropoxy)phenyl)propan-2-yl)phenoxy)propane-1,2-diol (1 equiv) and DMAP (~0.01 equiv) in anhydrous pyridine (~1M) in a water bath. The resulting solution was stirred at room temperature. The pyridine was removed under reduced pressure, and the
- 15 residue was diluted with ethyl acetate, washed subsequently with water (x 2), cold aqueous

1 M HCl, saturated NaHCO₃ and water. The organic layer was dried over Mg₂SO₄, filtered and concentrated the crude product. The crude product was purified by column chromatography (eluent: ethyl acetate/hexane) to afford Compound **i-H**. ¹H NMR and ¹³C NMR spectra are shown in Figures 1 and 2, respectively.

5

Compound ii-H (X = H)

Sodium azide (~1.5 equiv) was added to a solution of Compound **i-H** (1 equiv) in anhydrous dimethyl formamide, at room temperature. The reaction mixture was heated at 60 °C. Once the reaction is complete, any excess sodium azide was quenched using common laboratory procedures. The product was extracted with ethyl acetate (x 3). The organic layer was washed with saturated NaHCO₃ and water, dried over anhydrous magnesium sulfate, filtered, and then concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **ii-H**. ¹H NMR and ¹³C NMR spectra are shown in Figures 3 and 4, respectively.

15

Compounds 1a and 25a

CuI and NIS was added to a solution of Compound **ii-H** (1 equiv) and propargyl alcohol (~1.5equiv) in DMF and triethylamine. The reaction mixture is stirred at room temperature. Once the reaction was complete, any excess reagents were quenched using common laboratory procedures. The product was extracted with ethyl acetate (x 3). The organic layer was washed with saturated NaHCO₃ and water, dried over anhydrous magnesium sulfate, filtered, and then concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **1a** and Compound **25a**.

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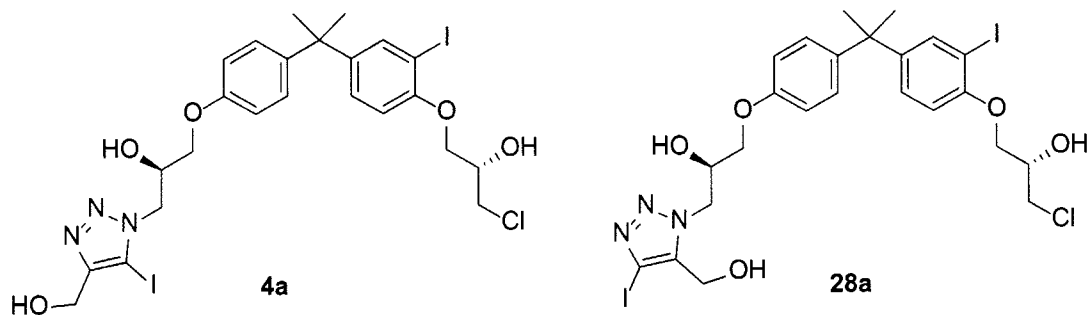
Compound **1a**: ¹H NMR and ¹³C NMR spectra are shown in Figures 5 and 6 respectively. MS (ES⁺): *m/z* 624.3 [M+Na]⁺.

Compound **25a**: ¹H NMR and ¹³C NMR spectra are shown in Figures 7 and 8, respectively. MS (ES⁺): *m/z* 624.3 [M+Na]⁺.

30

EXAMPLE 2

SYNTHESIS OF (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(4-(HYDROXYMETHYL)-5-iodo-1*H*-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol (**4a**) AND (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(5-(HYDROXYMETHYL)-4-iodo-1*H*-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)-2-iodophenoxy)propan-2-ol (**28a**)

**Compounds 4a and 28a**

Compound **4a** and Compound **28a** were synthesized according to Example 1 from (*R*)-3-(4-(2-(4-((*S*)-3-chloro-2-hydroxypropoxy)-3-iodophenyl)propan-2-yl)phenoxy)propane-1,2-diol.

Compound i-I (X = I): ¹H NMR and ¹³C NMR spectra are shown in Figures 9 and 10, respectively.

Compound ii-I (X = I): ¹H NMR and ¹³C NMR spectra are shown in Figures 11 and 12, respectively.

Compound **4a**: ¹H NMR and ¹³C NMR spectra are shown in Figures 13 and 14, respectively.

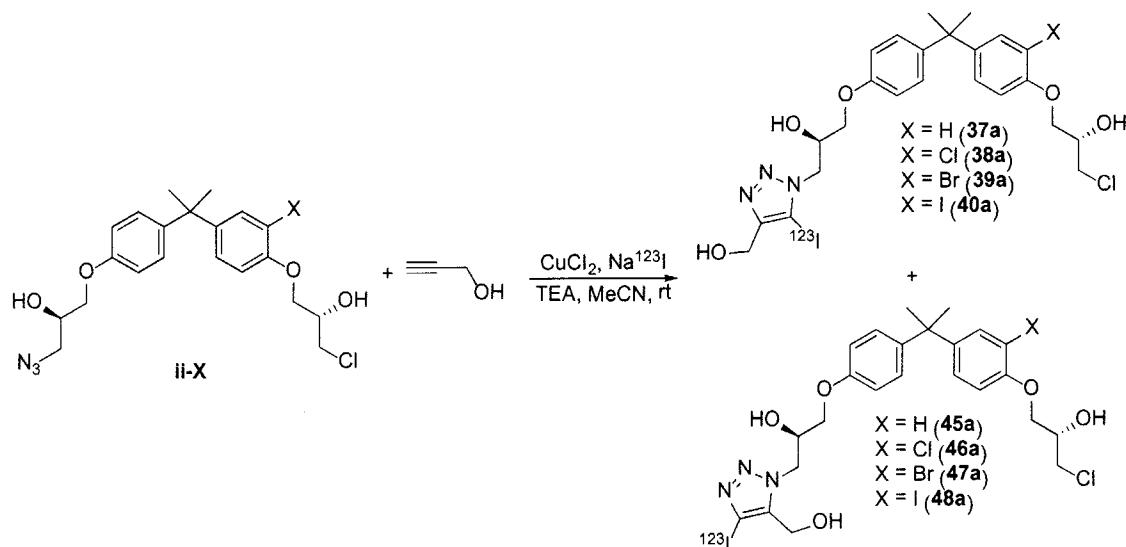
Compound **28a**: ¹H NMR and ¹³C NMR spectra are shown in Figures 15 and 16, respectively.

20 EXAMPLE 3

SYNTHESIS OF (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(4-(HYDROXYMETHYL)-5-(iodo-¹²³I)-1*H*-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol (**33a**) AND (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(5-(HYDROXYMETHYL)-4-(iodo-

¹²³I)-1*H*-1,2,3-triazol-1-yl)propoxy)phenyl)propan-2-yl)phenoxy)propan-2-ol

(41a)

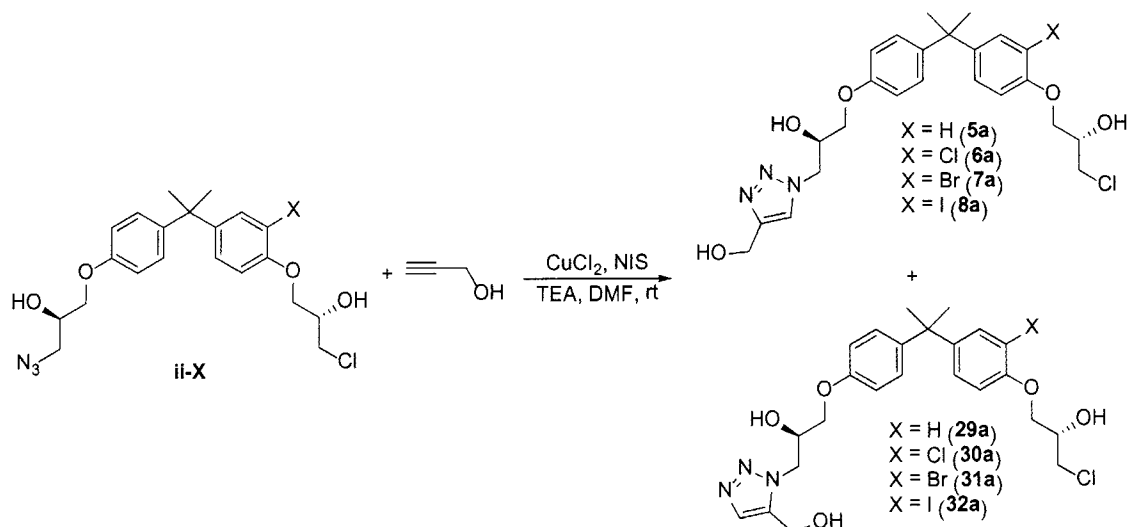


5 Compounds 37a and 45a

CuCl₂ and Na¹²³I are added to a solution of Compound ii-H (1 equiv) and propargyl alcohol (~1.5equiv) in acetonitrile and triethylamine. The reaction mixture is stirred at room temperature. Once the reaction is complete, any excess reagents are quenched using common laboratory procedures. The product is extracted with ethyl acetate (x 3). The organic layer is washed with saturated NaHCO₃ and water, dried over anhydrous magnesium sulfate, filtered, and then concentrated under reduced pressure. The resulting residue is purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound 37a and Compound 45a.

EXAMPLE 4

SYNTHESIS OF (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(4-(HYDROXYMETHYL)-1*H*-1,2,3-TRIAZOL-1-YL)PROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**5a**) AND (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-2-HYDROXY-3-(5-(HYDROXYMETHYL)-1*H*-1,2,3-TRIAZOL-1-YL)PROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**29a**)

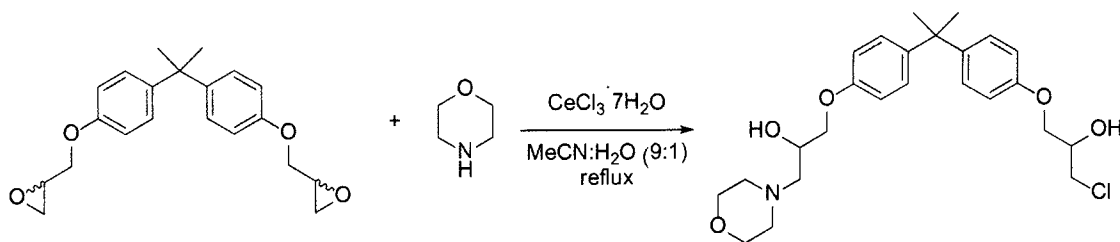


Compounds 5a and 29a

CuCl₂ and NIS is added to a solution of Compound **ii-H** (1 equiv) and propargyl alcohol (~1.5equiv) in DMF and triethylamine. The reaction mixture is stirred at room temperature. Once the reaction is complete, any excess reagents are quenched using common laboratory procedures. The product is extracted with ethyl acetate (x 3). The organic layer is washed with saturated NaHCO₃ and water, dried over anhydrous magnesium sulfate, filtered, and then concentrated under reduced pressure. The resulting residue is purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **5a** and Compound **29a**.

EXAMPLE 5

SYNTHESIS OF 1-CHLORO-3-(4-(2-(4-(2-HYDROXY-3-MORPHOLINOPROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**13**)



5

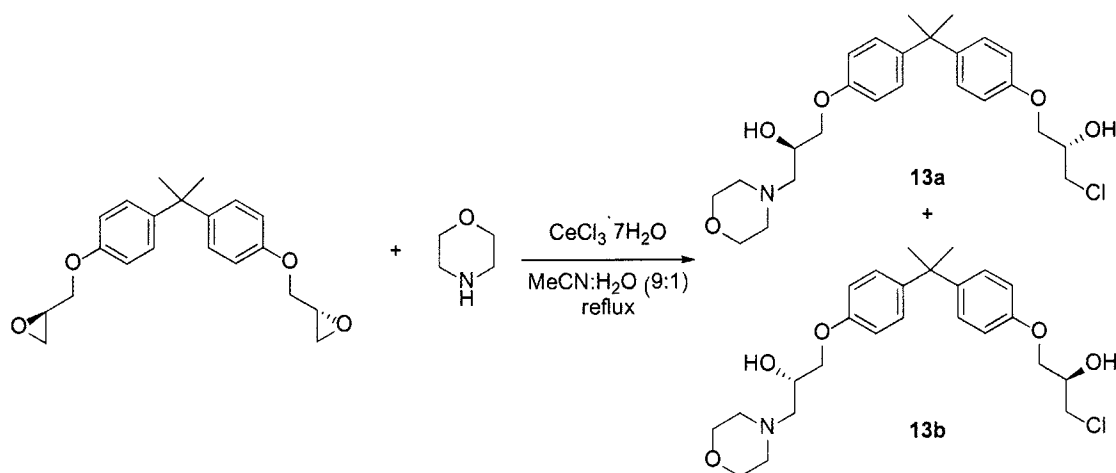
Compound 13

To a solution of 2,2'-(((propane-2,2-diylbis(4,1-phenylene))bis(oxy))bis(methylene))bis(oxirane) (1.0 equiv) and morpholine (~1 equiv) in acetonitrile/water (9:1) was added $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ (~0.5 equiv) and the mixture was refluxed. The resulting crude mixture was filtered and washed with ethyl acetate, and the resulting suspension was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide the title Compound **13**. ^1H NMR spectrum is shown in Figures 17. MS (ESI⁺): m/z 464.3 $[\text{M}]^+$.

10

EXAMPLE 6

SYNTHESIS OF (S)-1-CHLORO-3-(4-(2-(4-((R)-2-HYDROXY-3-MORPHOLINOPROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**13a**)



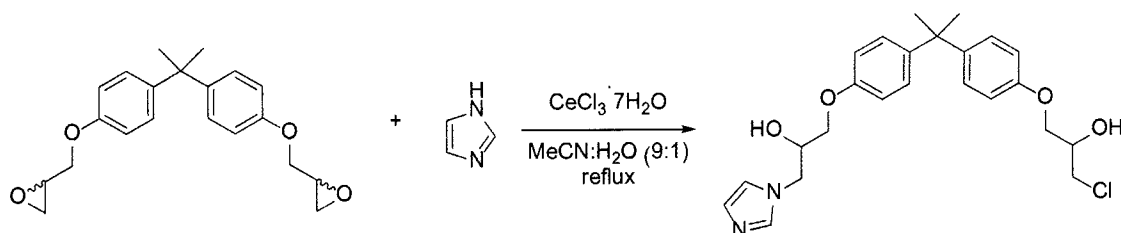
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Compound 13a

To a solution of (2R,2'R)-2,2'-(((propane-2,2-diylbis(4,1-phenylene))bis(oxy))bis(methylene))bis(oxirane) (1.0 equiv) and morpholine (~1 equiv) in acetonitrile/water (9:1) is added $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ (~0.5 equiv) and the mixture is refluxed. The resulting crude mixture is filtered and washed with ethyl acetate, and the resulting suspension is concentrated under reduced pressure. The resulting residue is purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **13a** and Compound **13b**.

EXAMPLE 7

SYNTHESIS OF 1-CHLORO-3-(4-(2-(4-(2-HYDROXY-3-(1*H*-IMIDAZOL-1-
YL)PROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**9**)



5

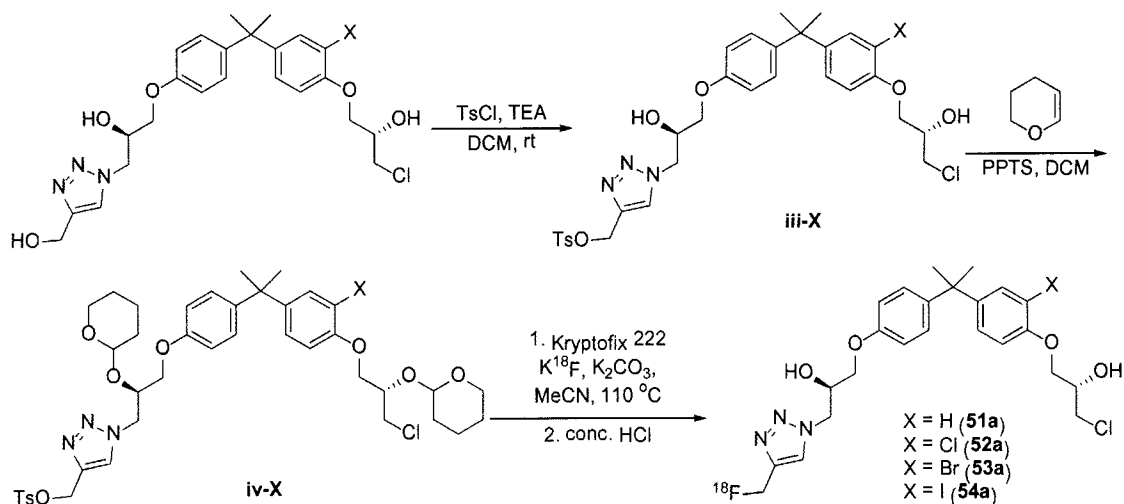
Compound 9

To a solution of 2,2'-(((propane-2,2-diylbis(4,1-phenylene))bis(oxy))bis(methylene))bis(oxirane) (1 equiv) and 1*H*-imidazole (~1 equiv) in acetonitrile/water (9:1) were added $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ (~0.5 equiv) and the mixture was refluxed.

- 10 The resulting crude mixture was filtered and washed with ethyl acetate, and the resulting suspension was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **9**.

EXAMPLE 8

- 15 SYNTHESIS OF (*S*)-1-CHLORO-3-(4-(2-(4-((*R*)-3-(4-((FLUORO- ^{18}F)METHYL)-1*H*-1,2,3-TRIAZOL-1-YL)-2-HYDROXYPROPOXY)PHENYL)PROPAN-2-YL)PHENOXY)PROPAN-2-OL (**51a**)



Compound iii-H (X = H)

p-Toluenesulfonyl chloride (~1.5 equiv) is added portion wise over a period of approximately 10 min to a solution of Compound **5a** (1 equiv) and TEA in DCM (~1M) in a water bath. The resulting solution is stirred at room temperature. The solvent is removed under reduced pressure, and the residue is diluted with ethyl acetate, washed subsequently with water (x 2), cold aqueous 1 M HCl, saturated NaHCO₃ and water. The organic layer was dried over Mg₂SO₄, filtered and concentrated the crude product. The crude product is purified by column chromatography (eluent: ethyl acetate/hexane) to afford Compound **iii-H**.

Compound iv-H (X = H)

3,4-Dihydro-2H-pyran and PPTS is added to a solution of Compound **iii-H** (1 equiv) in DCM. The resulting solution is stirred until reaction is complete. The solvent is removed under reduced pressure, and the residue is diluted with ethyl acetate, washed subsequently with saturated NaHCO₃ and water. The organic layer was dried over Mg₂SO₄, filtered and concentrated the crude product. The crude product is purified by column chromatography (eluent: ethyl acetate/hexane) to afford Compound **iv-H**.

Compound 51a

To a solution of Compound **iv-H** (1 equiv) in MeCN is added K¹⁸F and K₂CO₃, and the reaction mixture is heated at 110 °C. Once the reaction is complete, concentrated HCl is

added and stirred. The product is extracted with ethyl acetate, and the resulting suspension is concentrated under reduced pressure. The resulting residue is purified by flash column chromatography on silica gel (eluent: ethyl acetate/hexane) to provide Compound **51a**.

EXAMPLE 9

5 COMPOUND ACTIVITY

LNCaP cells were transiently transfected with PSA (6.1 kb)-luciferase for 24 h prior to pre-treatment with compounds of the invention (*e.g.*, compounds **1a**, **9**, **13**) ranging in concentration from 62.5 ng/ml to 1.5 ug/ml for 1 hour before the addition of vehicle, or synthetic androgen, R1881 (1 nM) to induce luciferase production. After 24 h of incubation
10 with R1881, the cells were harvested, and relative luciferase activities were determined. To determine the IC₅₀, treatments were normalized to the predicted maximal activity induction (in the absence of test compounds, vehicle only).

Table 4. IC₅₀ values for selected compounds (μM) with standard error

15

Compound	Number of Trials	Average IC ₅₀
1a	4	3.23 +/- 0.37
9	6	8.45 +/- 3.14
13	3	12.84 +/- 1.95

Although various embodiments of the invention are disclosed herein, many adaptations and modifications can be made within the scope of the invention in accordance with the common general knowledge of those skilled in this art. Such modifications
20 include the substitution of known equivalents for any aspect of the invention in order to achieve the same result in substantially the same way. Numeric ranges are inclusive of the numbers defining the range. The word "comprising" is used herein as an open-ended term, substantially equivalent to the phrase "including, but not limited to", and the word

"comprises" has a corresponding meaning. As used herein, the singular forms "a", "an" and "the" include plural referents unless the context clearly dictates otherwise. Thus, for example, reference to "a thing" includes more than one such thing. Citation of references herein is not an admission that such references are prior art to the present invention. Any

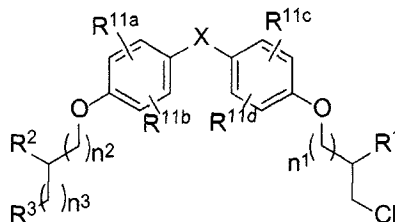
5 priority document(s) and all publications, including but not limited to patents and patent applications, cited in this specification are incorporated herein by reference as if each individual publication were specifically and individually indicated to be incorporated by reference herein and as though fully set forth herein. The invention includes all embodiments and variations substantially as hereinbefore described and with reference to

10 the examples and drawings.

CLAIMS

What is claimed is:

1. A compound having the following structure (I):



(I),

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-, -C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-,

R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally substituted with one or more R⁶;

R⁵ is each independently H, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I, hydroxyl, oxo, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₆-C₁₂ aryl, wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -OS(O)₂-aryl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁸ and R⁹ are each independently H, halogen, -S(O)₀₋₂R⁵, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, aryl, aralkyl, C₁-C₁₀ acyl, or -NR⁵R⁵, or R⁸ and R⁹ can join to form a unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

R^{11a}, R^{11b}, R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ¹²³I, hydroxyl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl; -OR⁵, -OC(=O)R¹³, C₁-C₁₀ acyl, -S(O)₀₋₂R⁵, -NO₂-, CN, -NH₂, -NHR⁵, or -N(R⁵)₂;

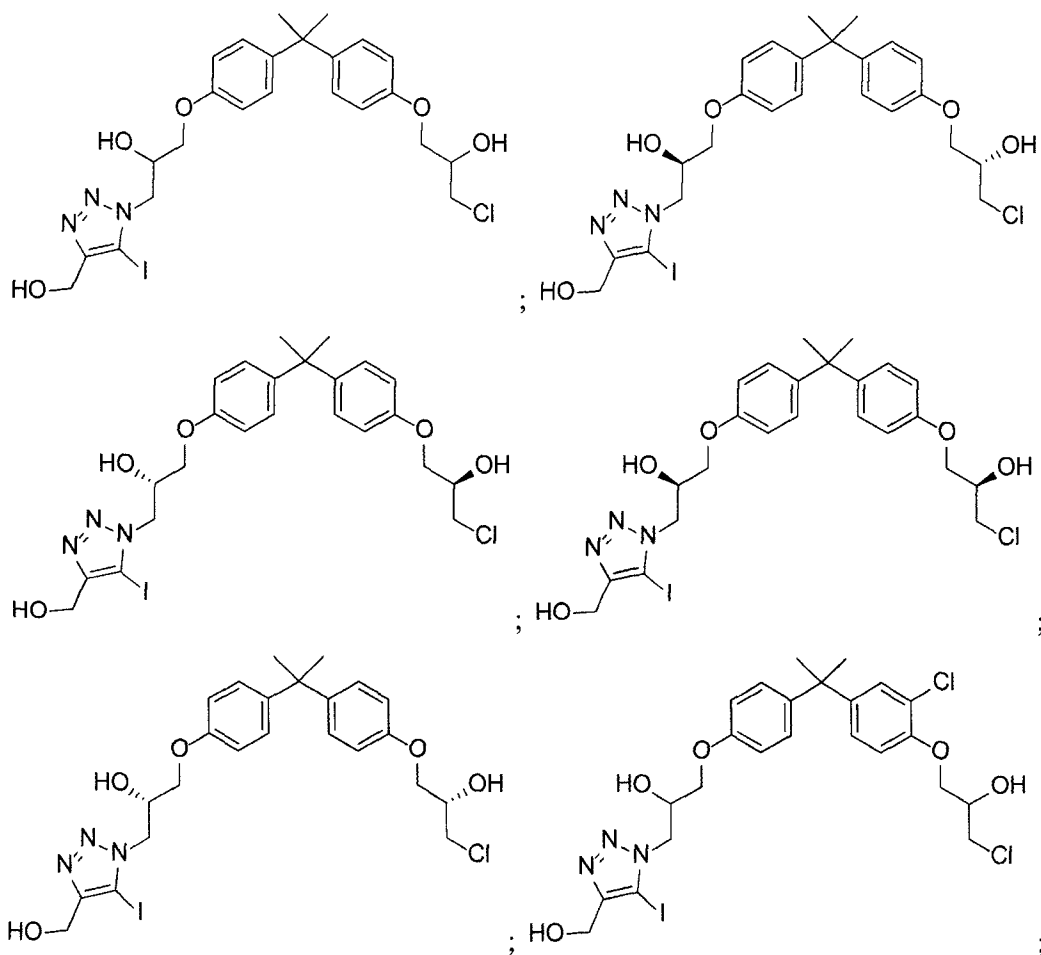
R¹³ is each independently C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

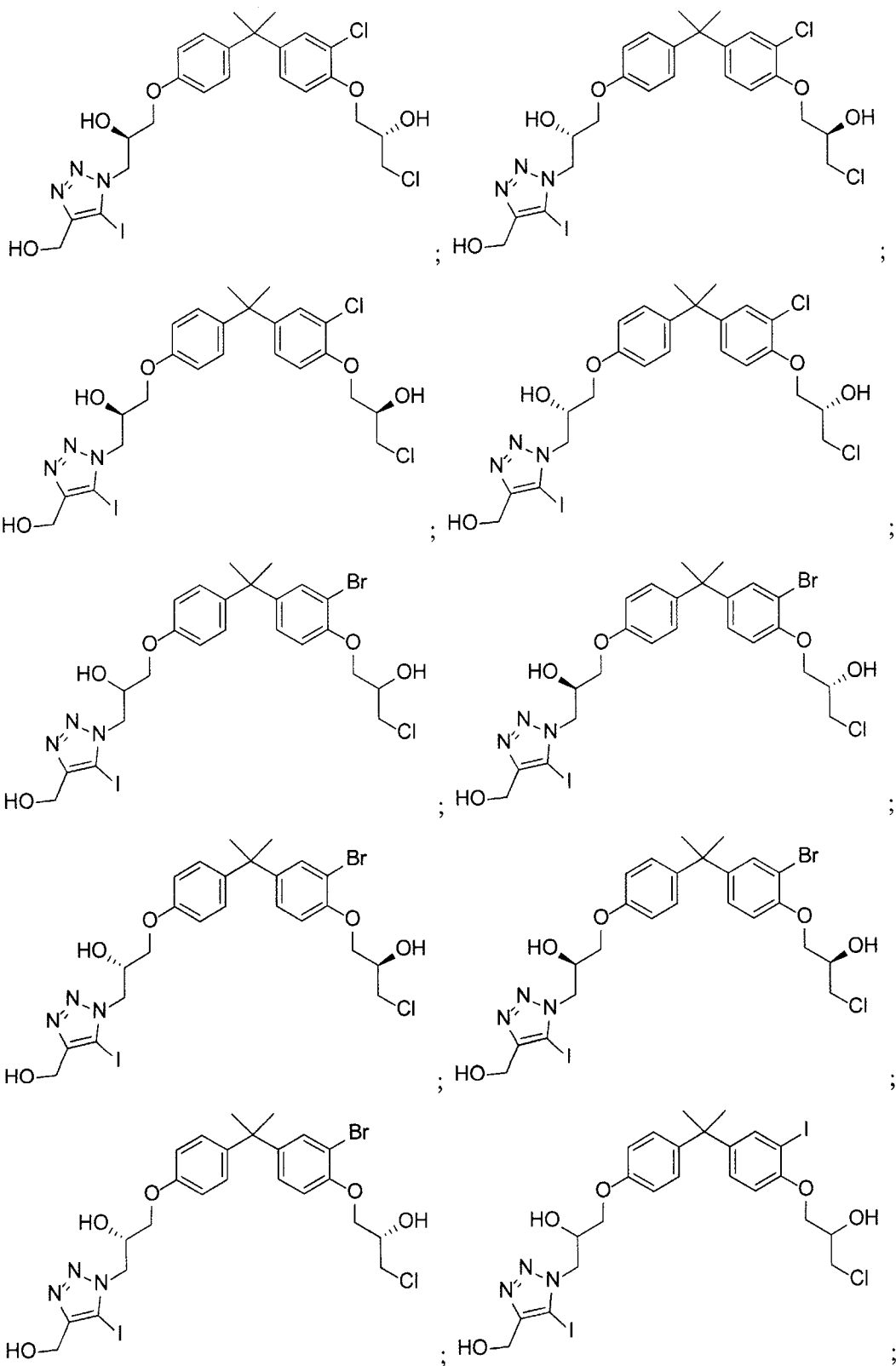
n¹ and n² are each independently 0, 1, or 2; and

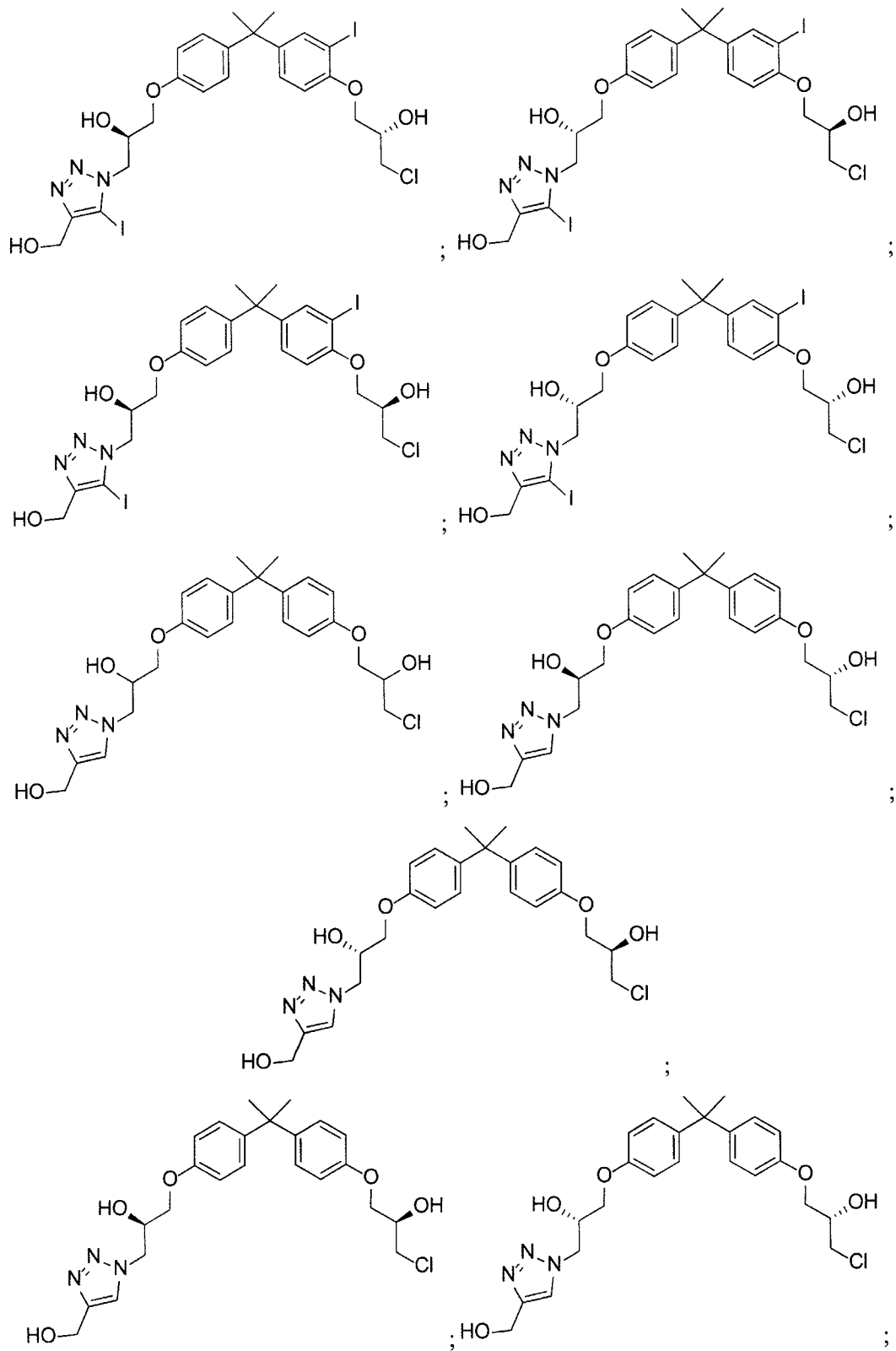
n^3 is 0, 1, 2, 3, 4, or 5.

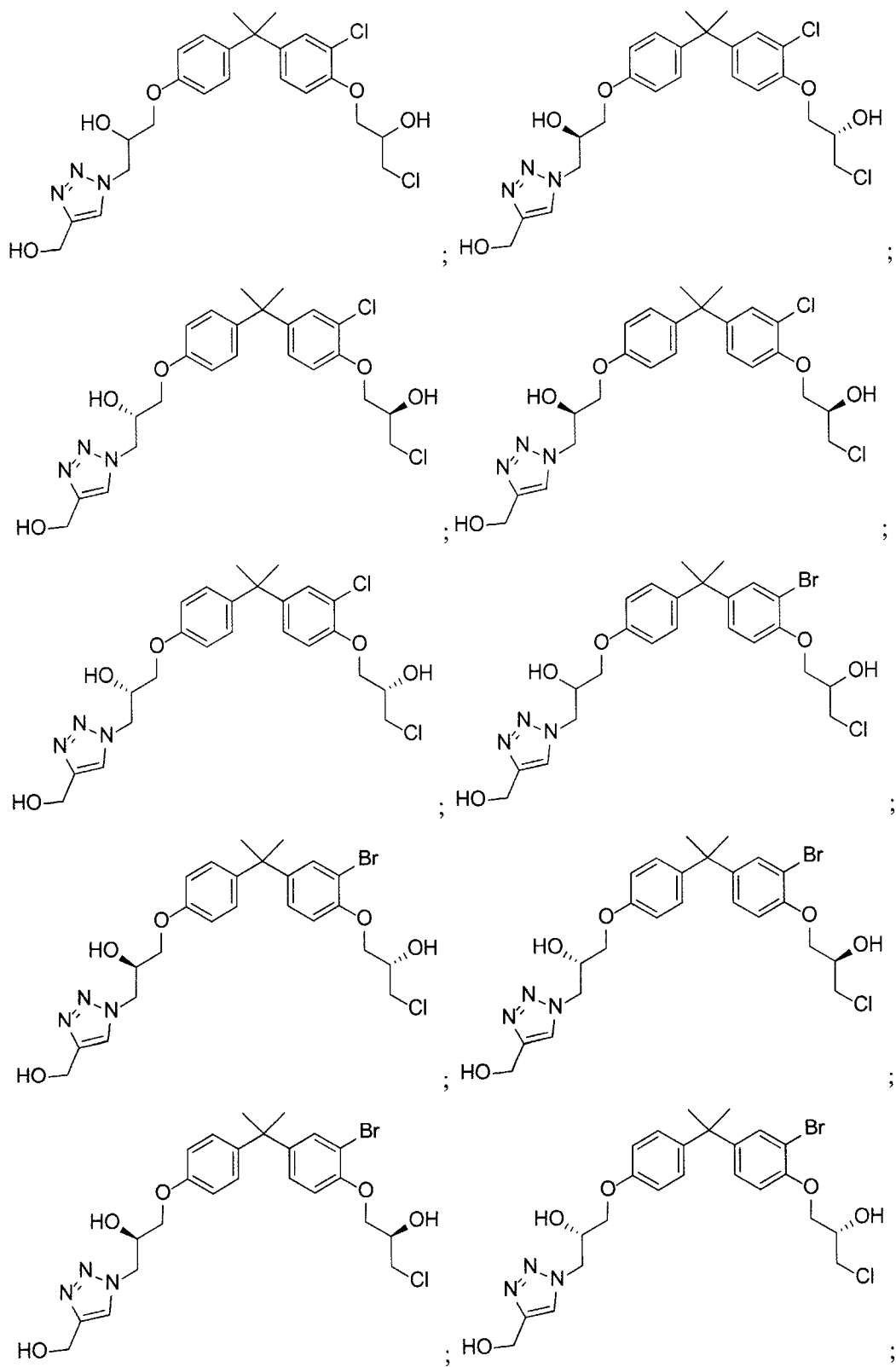
2. The compound of claim 1, wherein X is $-C(R^8R^9)-$.
3. The compound of claim 1 or 2, wherein R^8 and R^9 are C_1 - C_{10} alkyl.
4. The compound of claim 1 or 2, wherein R^8 and R^9 are methyl.
5. The compound of claim 1, wherein R^3 is 5-6 membered heteroaryl or 3-7 membered heterocyclyl, wherein said heteroaryl or said heterocyclyl comprises at least one N atom in the ring.
6. The compound of claim 1, wherein R^3 is selected from a group consisting of pyrrole, furan, thiophene, pyrazole, pyridine, pyridazine, pyrimidine, imidazole, thiazole, isoxazole, oxadiazole, thiadiazole, oxazole, triazole, isothiazole, oxazine, triazine, azepine, pyrrolidine, pyrroline, imidazoline, imidazolidine, pyrazoline, pyrazolidine, piperidine, dioxane, morpholine, dithiane, thiomorpholine, piperazine, and tetrazine.
7. The compound of claim 1, wherein R^3 is substituted with at least one ^{123}I or I.
8. The compound of claim 1, wherein R^3 is substituted with at least one R^6 , wherein at least one R^6 is further substituted with at least one of ^{123}I , I, or ^{18}F .
9. The compound of claim 1, wherein at least one R^6 is C_1 - C_6 alkyl, wherein at least one R^6 is further substituted with at least one of ^{123}I , I, or ^{18}F .
10. The compound of claim 1, wherein each R^{13} is independently methyl, ethyl or propyl.
11. The compound of claim 1, wherein each R^{13} is methyl.

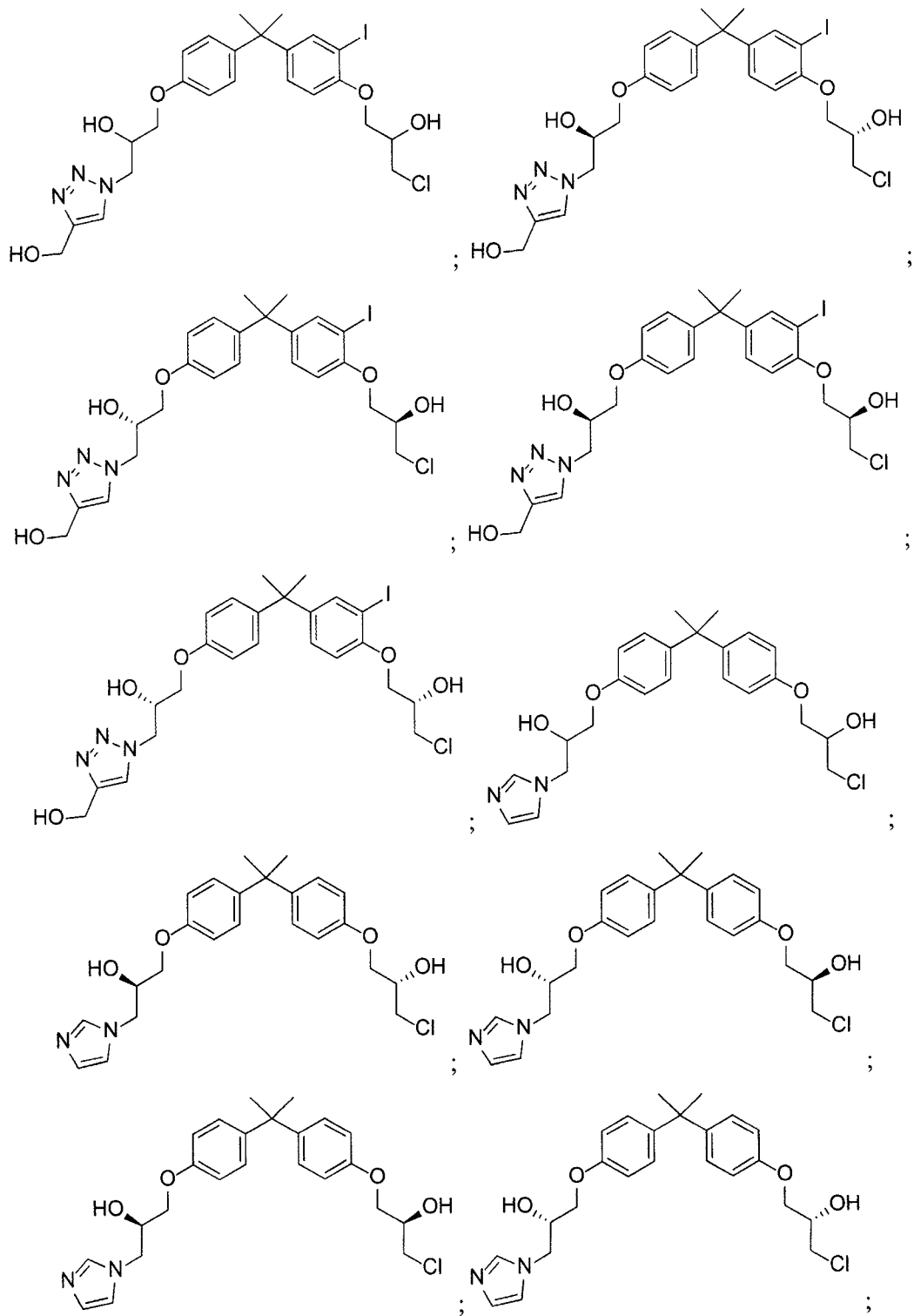
12. The compound of claim 1, wherein at least one of R^{11a} , R^{11b} , R^{11c} and R^{11d} is Cl, Br, I or ^{123}I .
13. The compound of claim 1, wherein at least one of R^5 , R^{11a} , R^{11b} , R^{11c} and R^{11d} is I; and at least one of R^5 , R^{11a} , R^{11b} , R^{11c} and R^{11d} is ^{123}I .
14. The compound of claim 1, wherein n^3 is 0, 1, or 2.
15. The compound of claim 1, wherein the compound has one of the following structures:

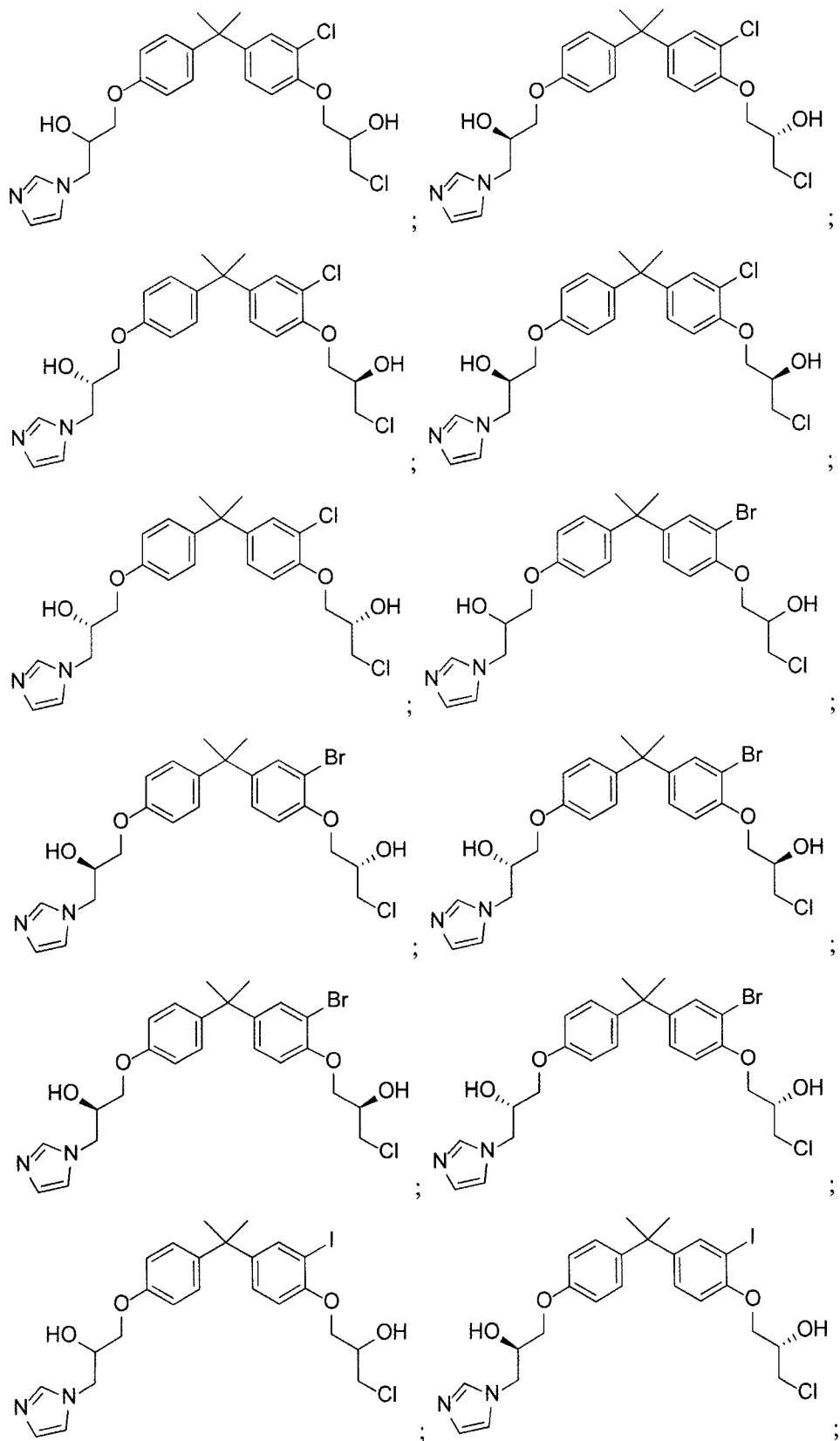


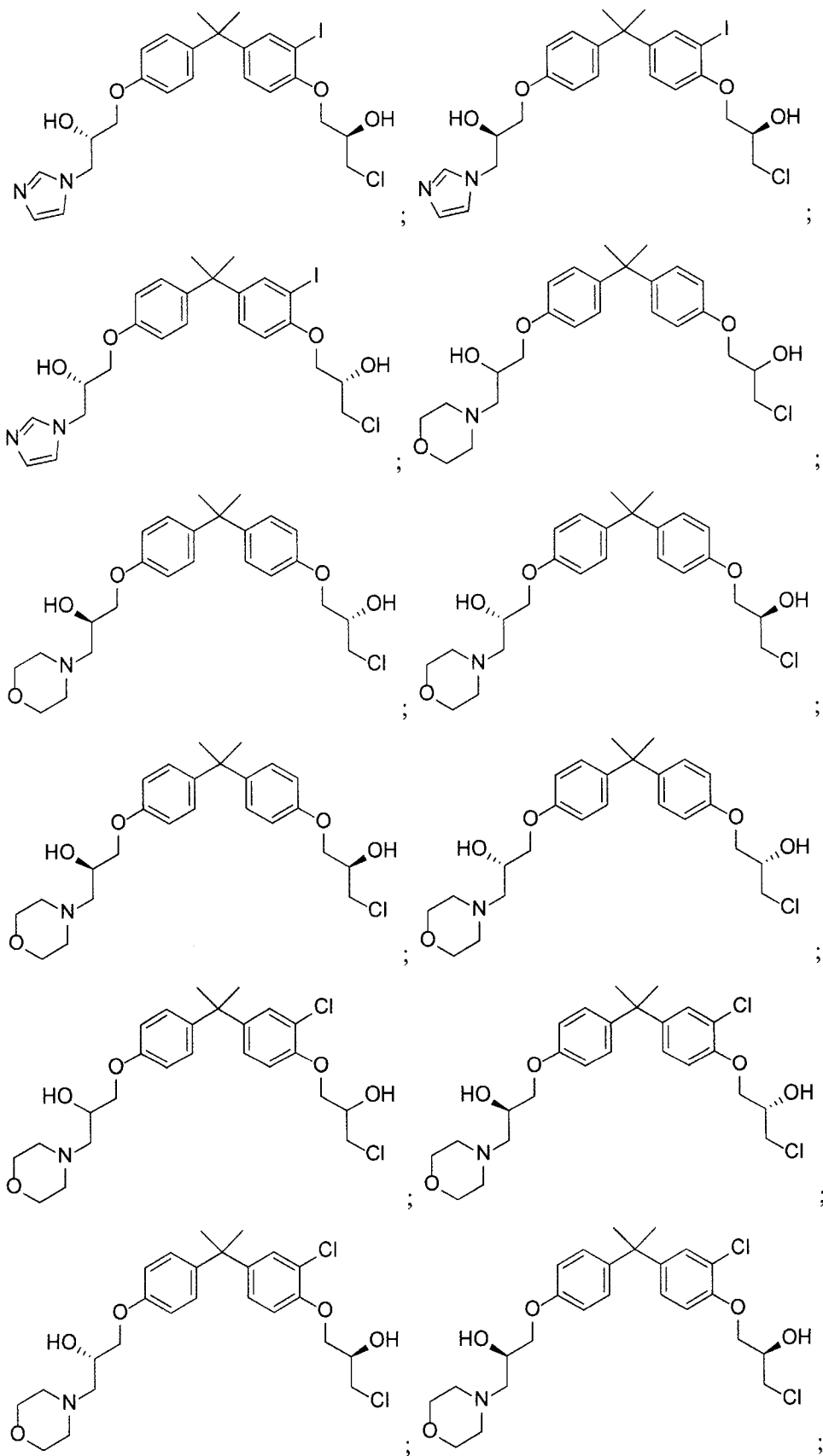


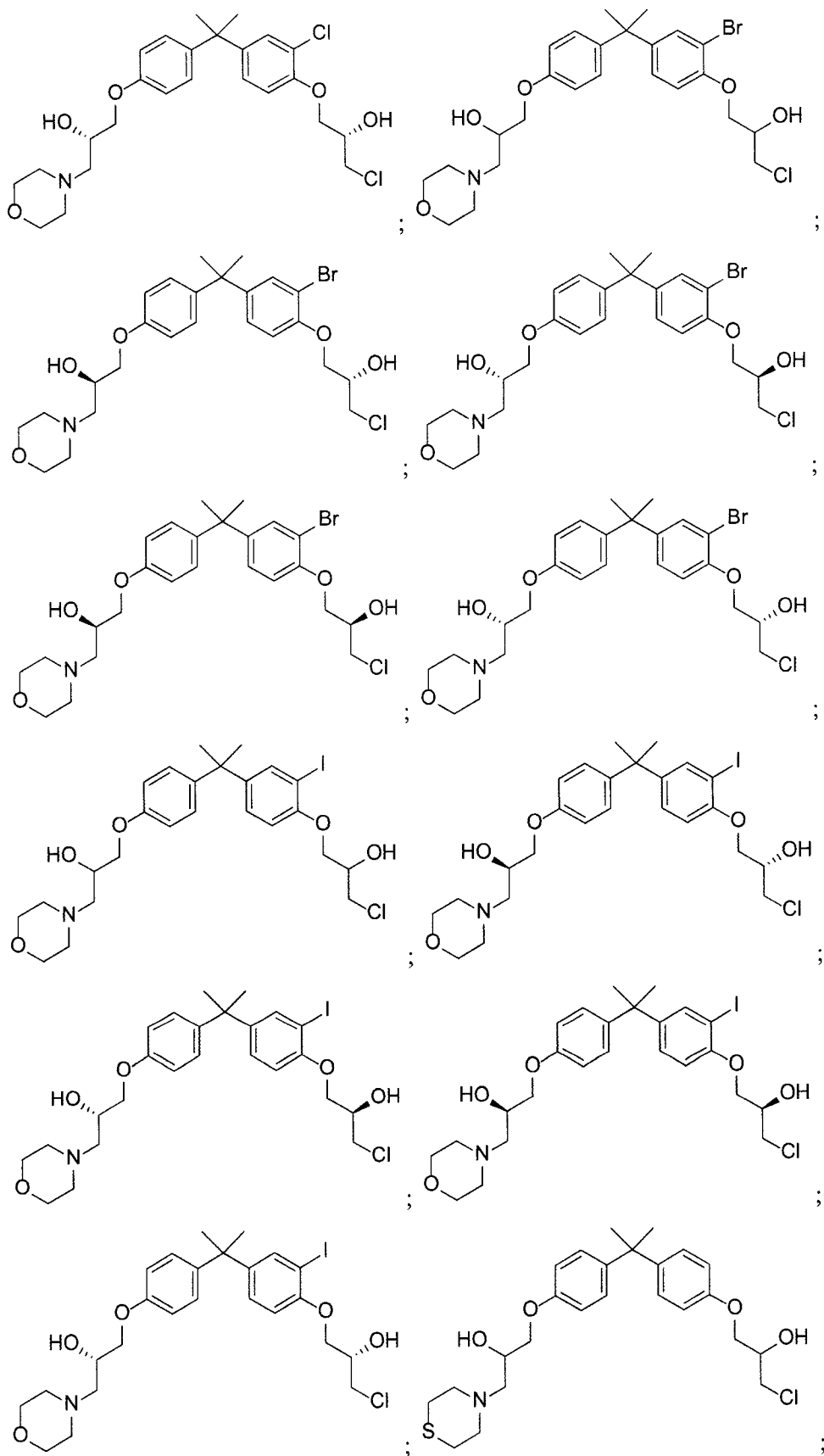


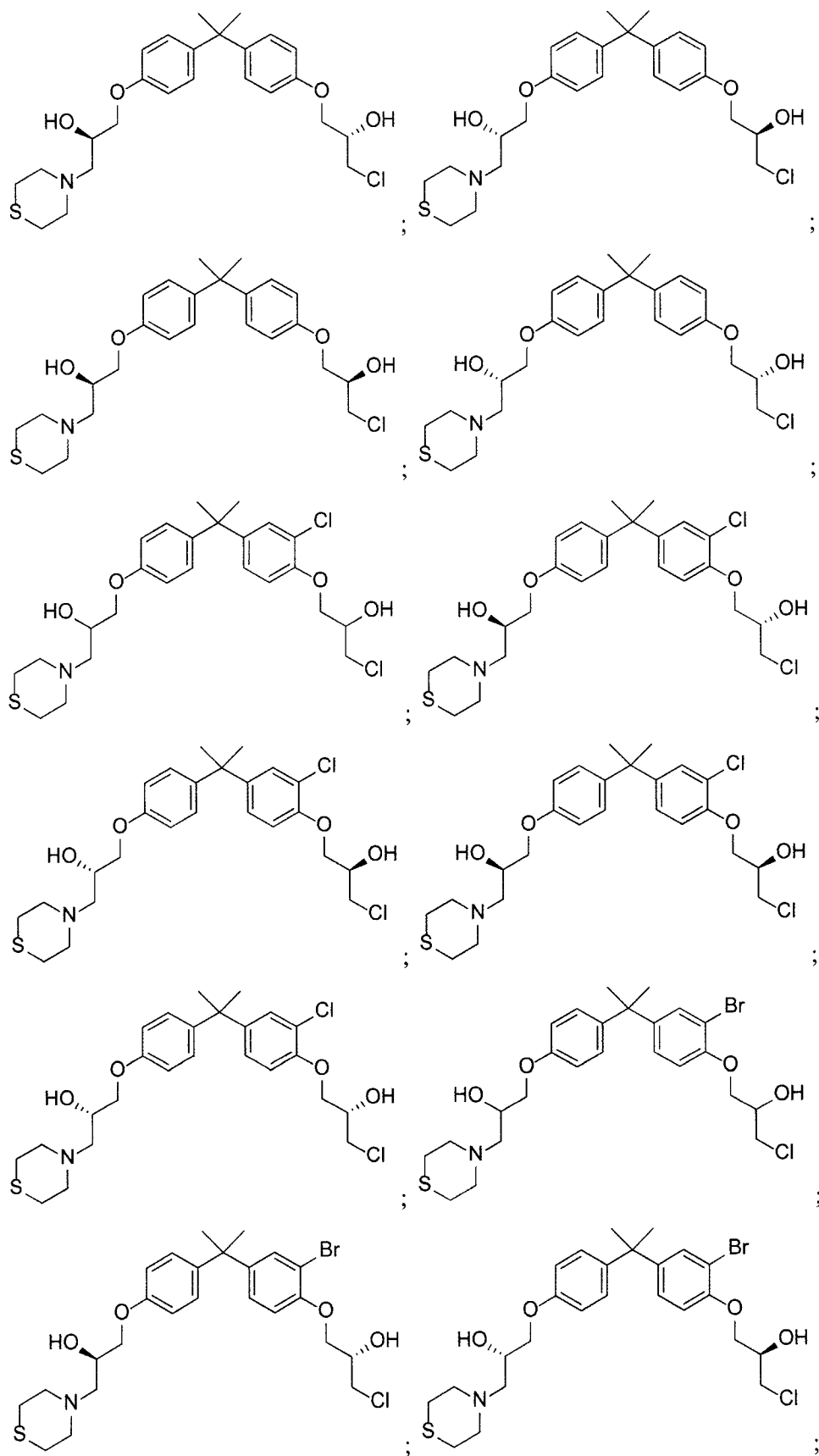


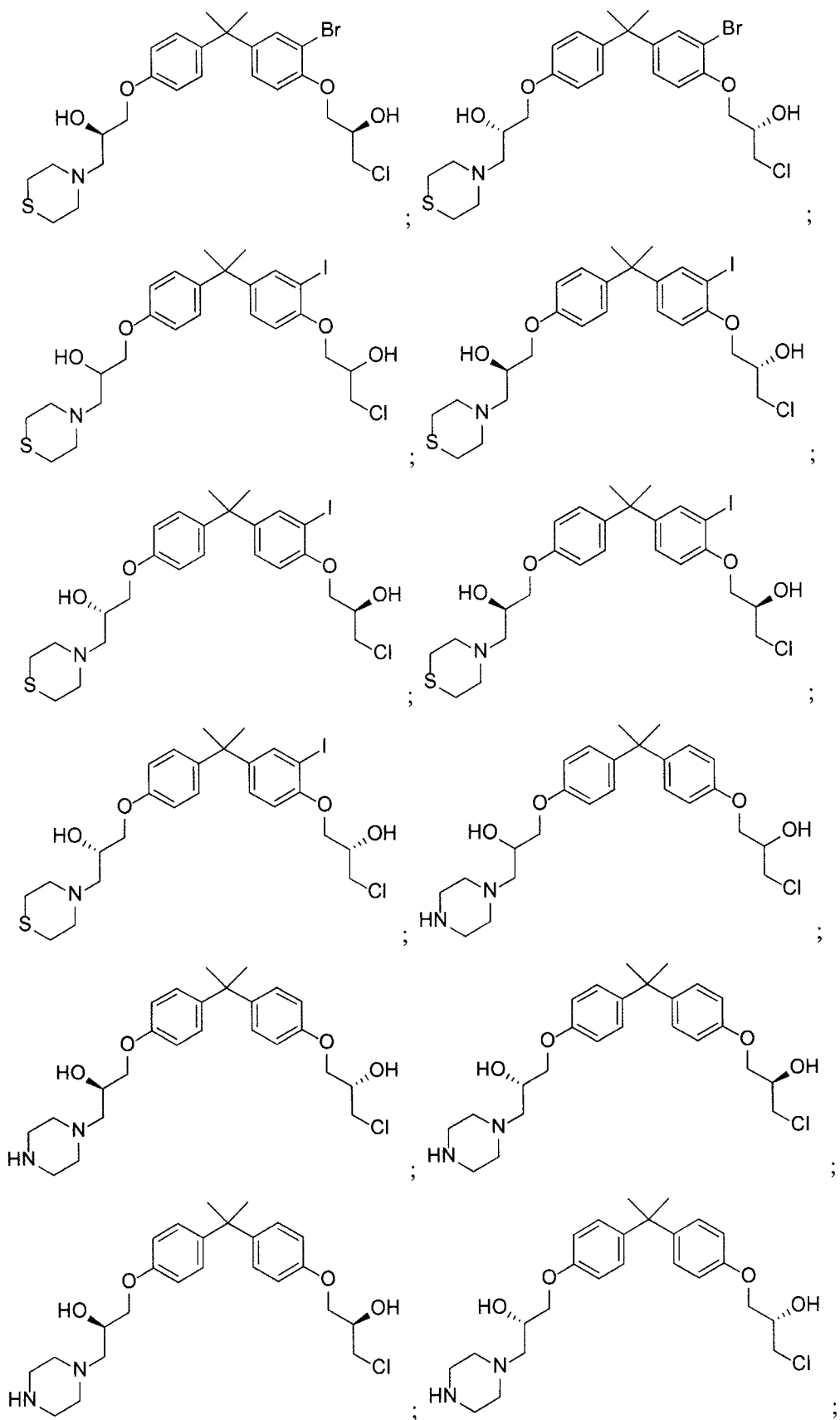


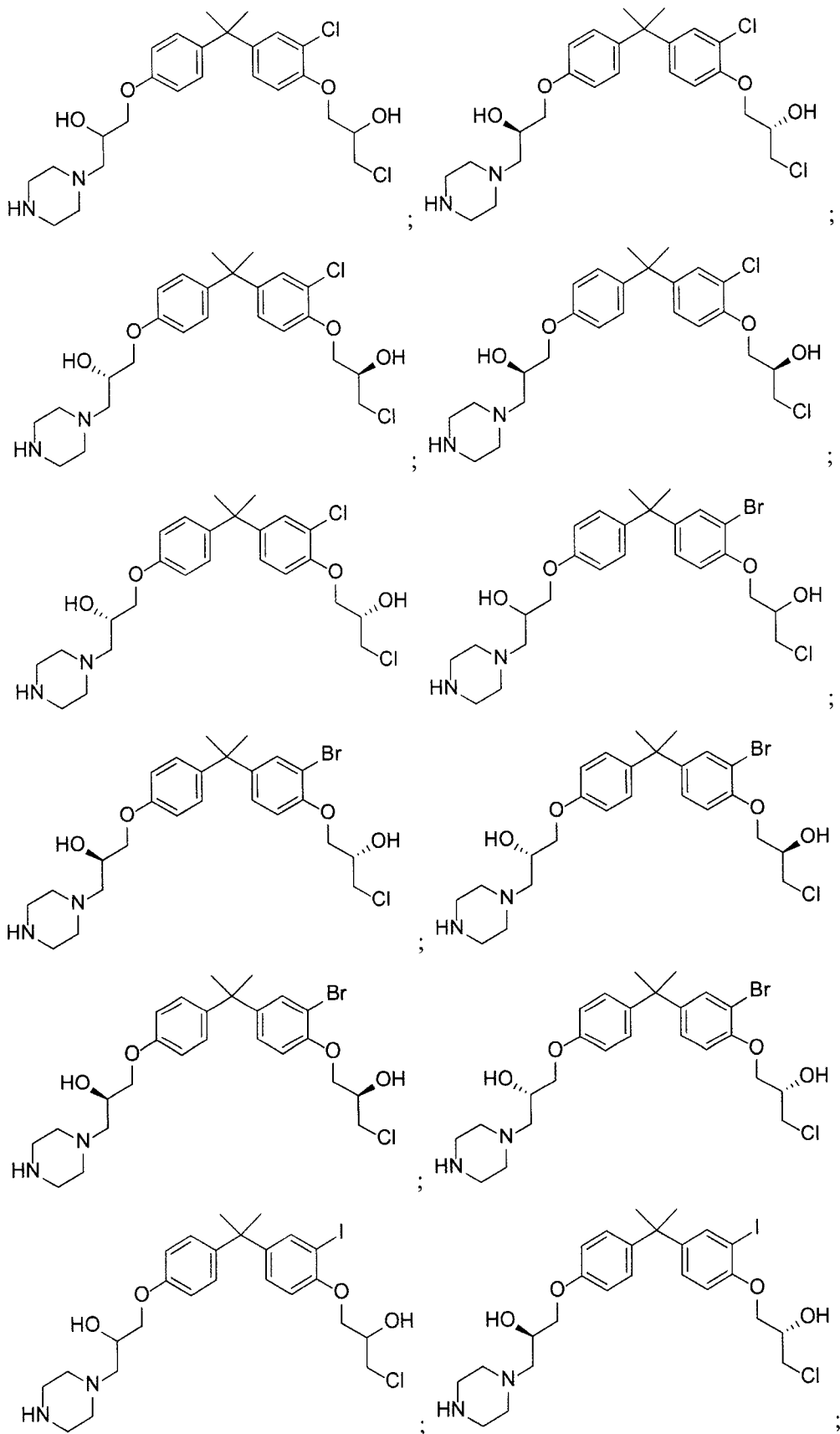


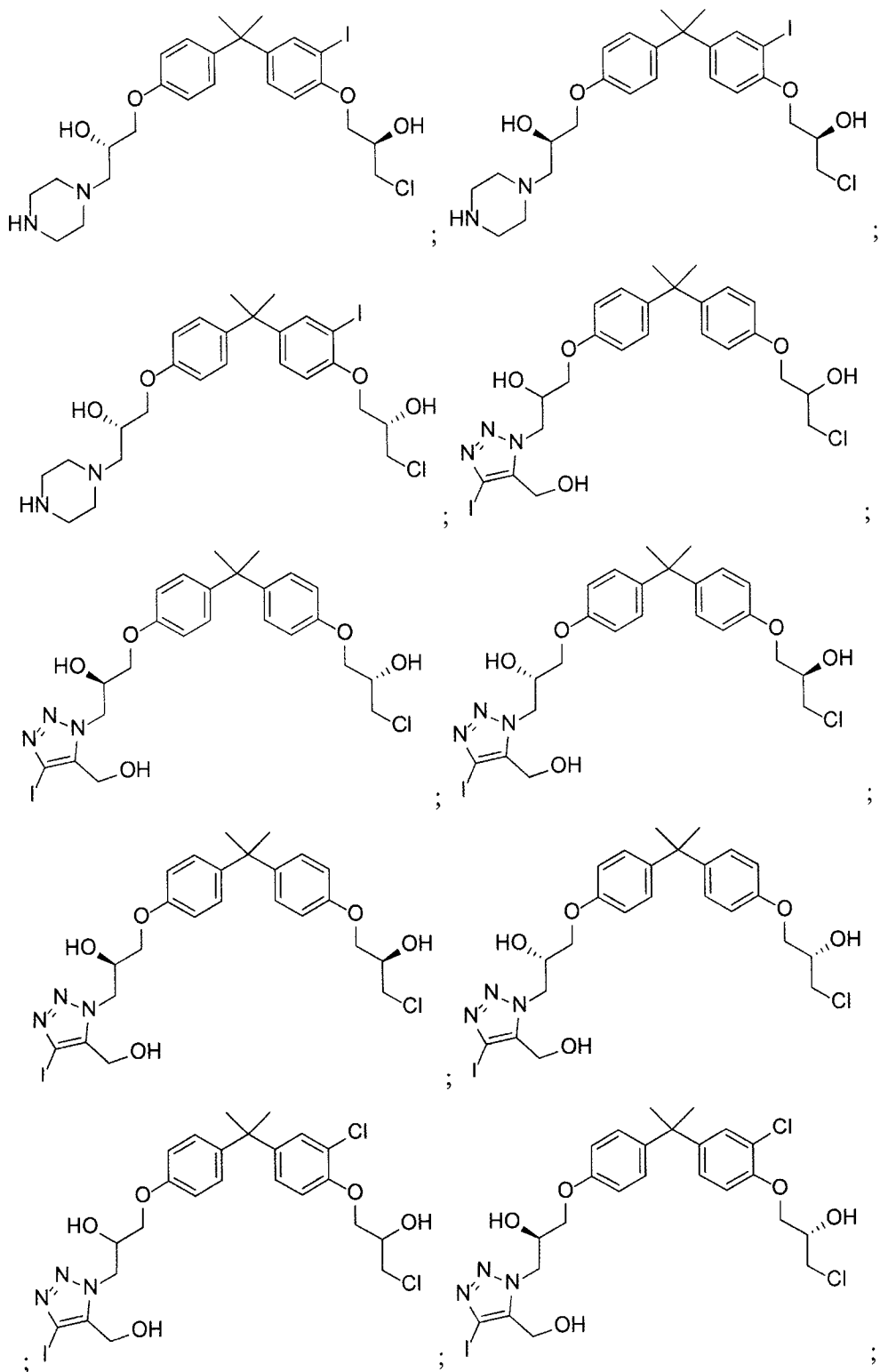


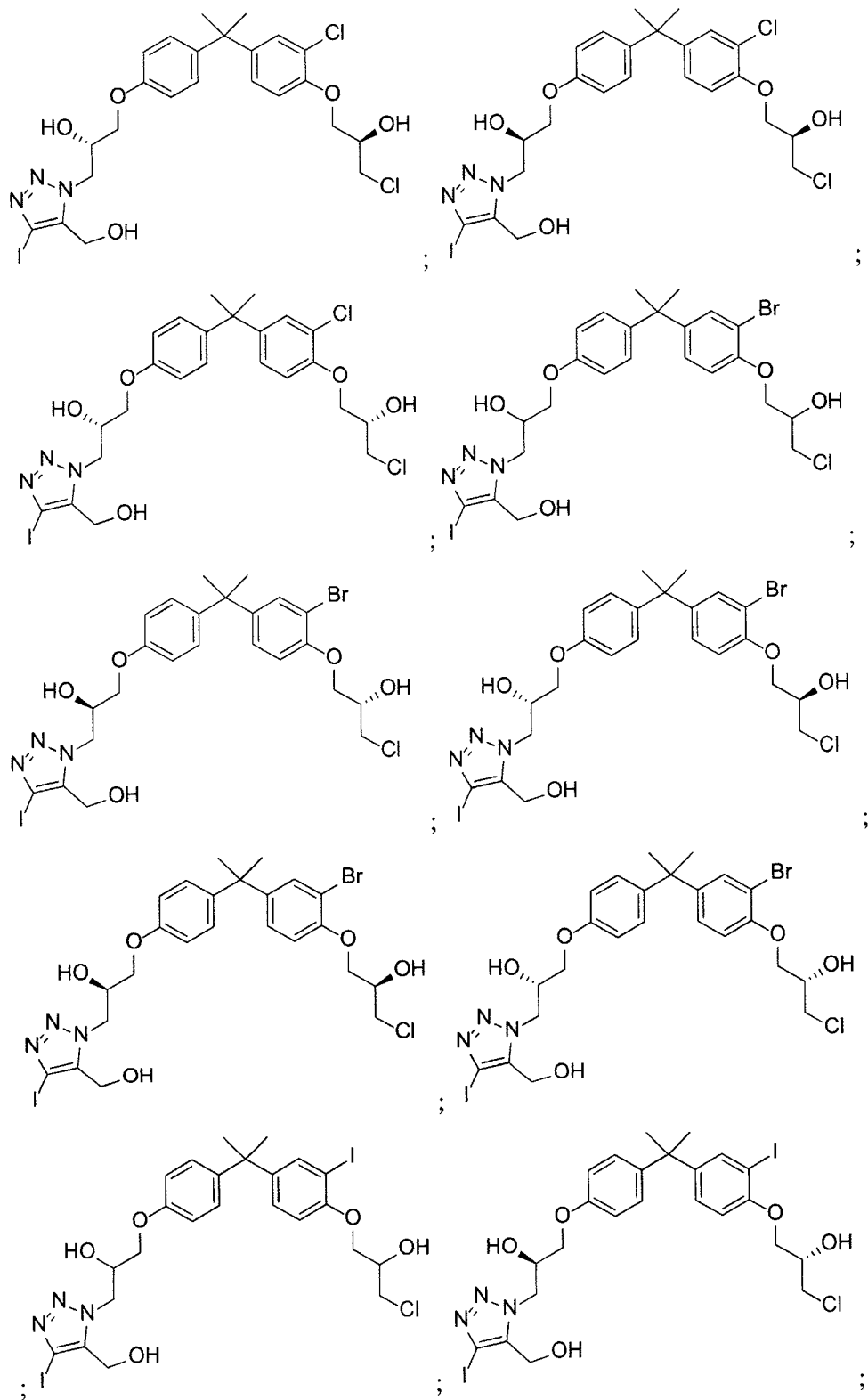


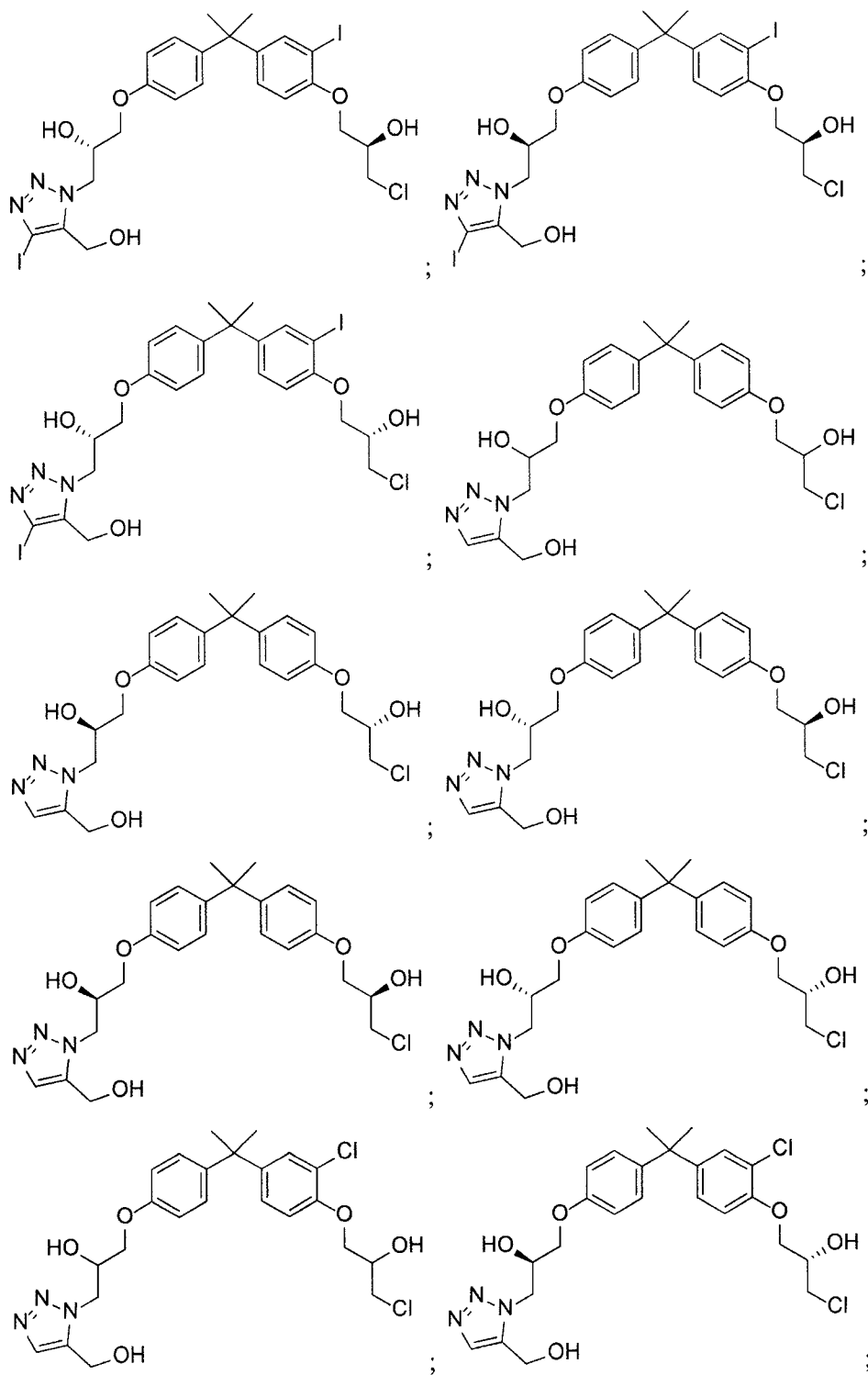


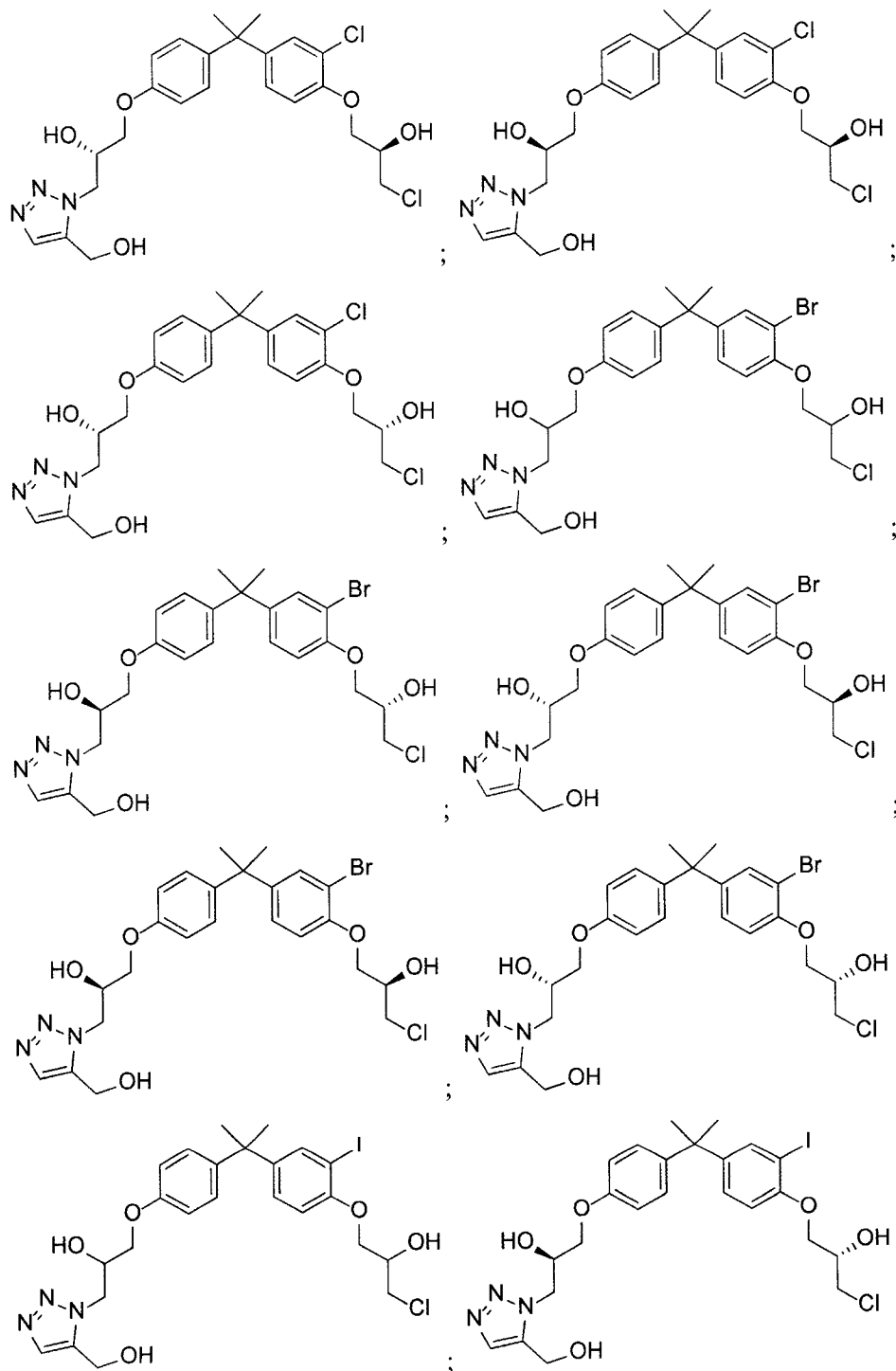


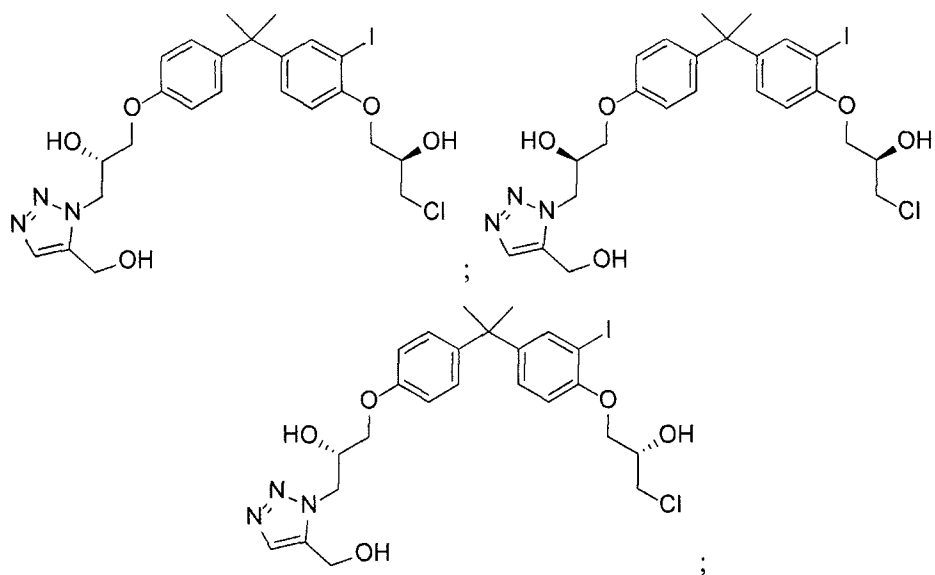






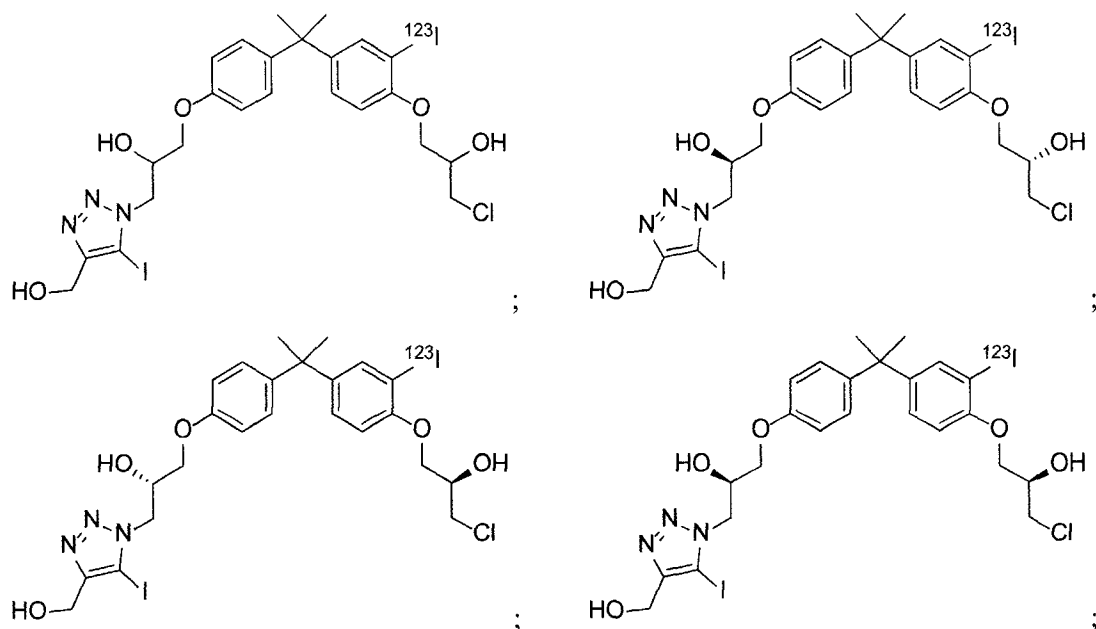


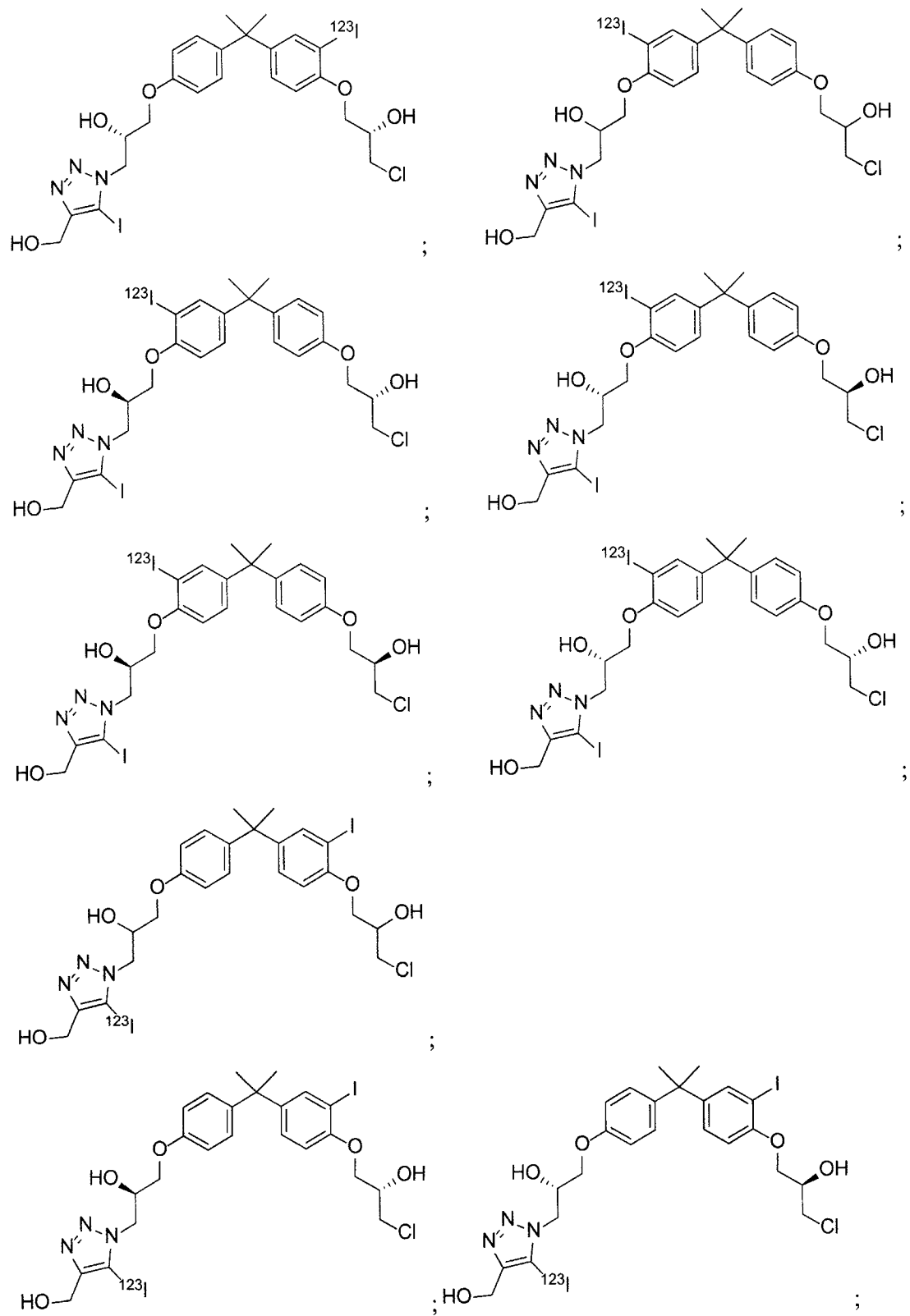


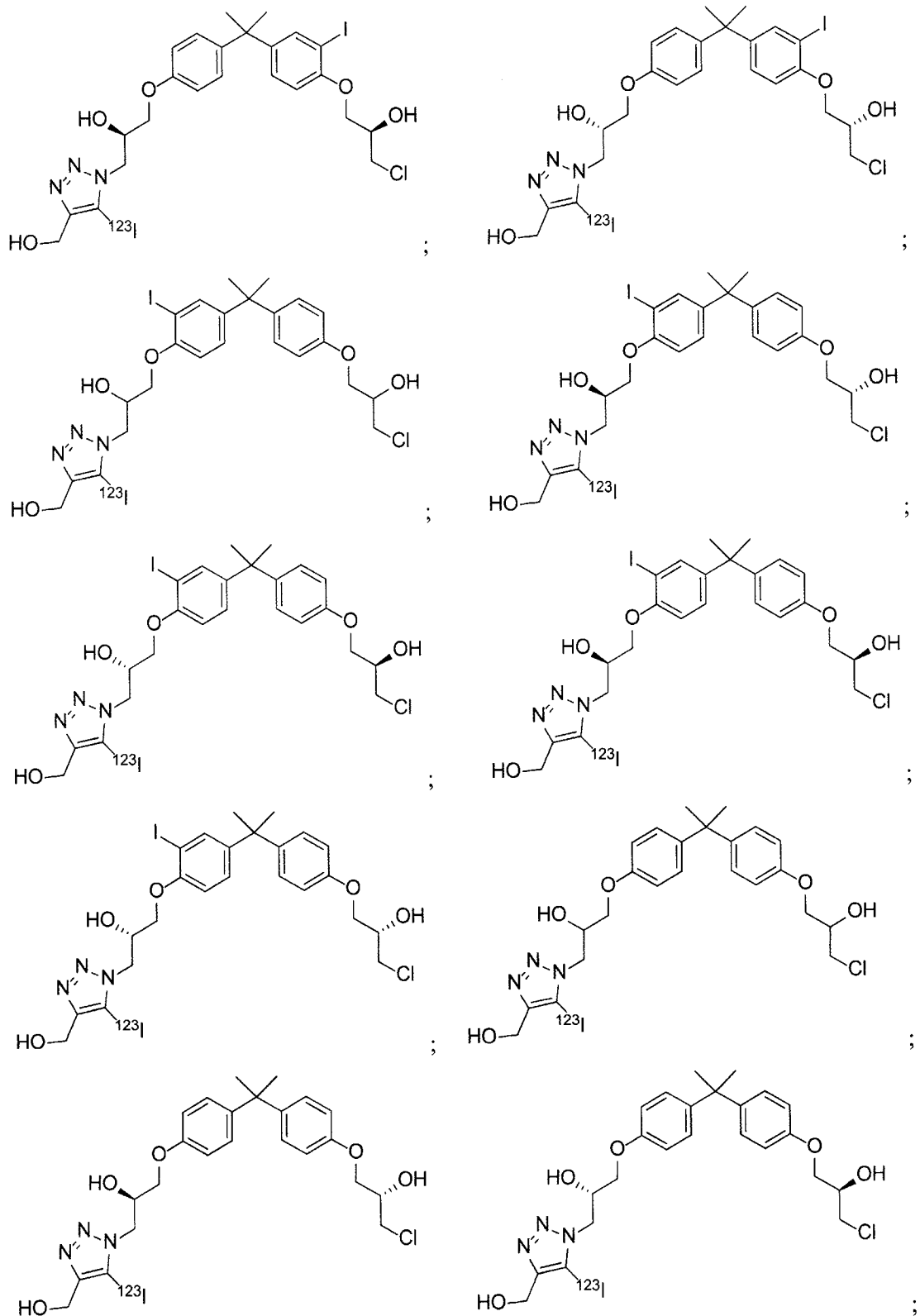


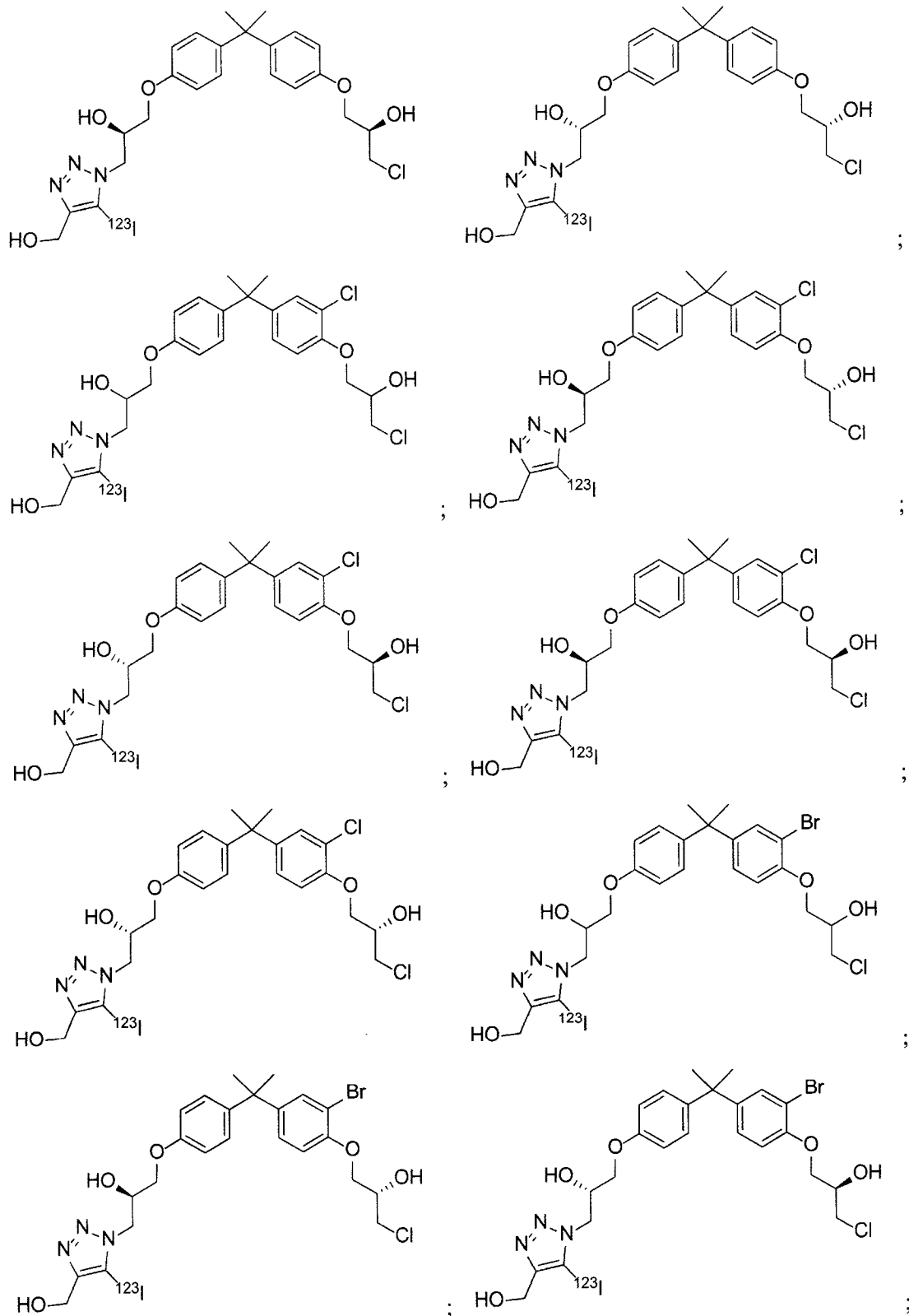
or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof.

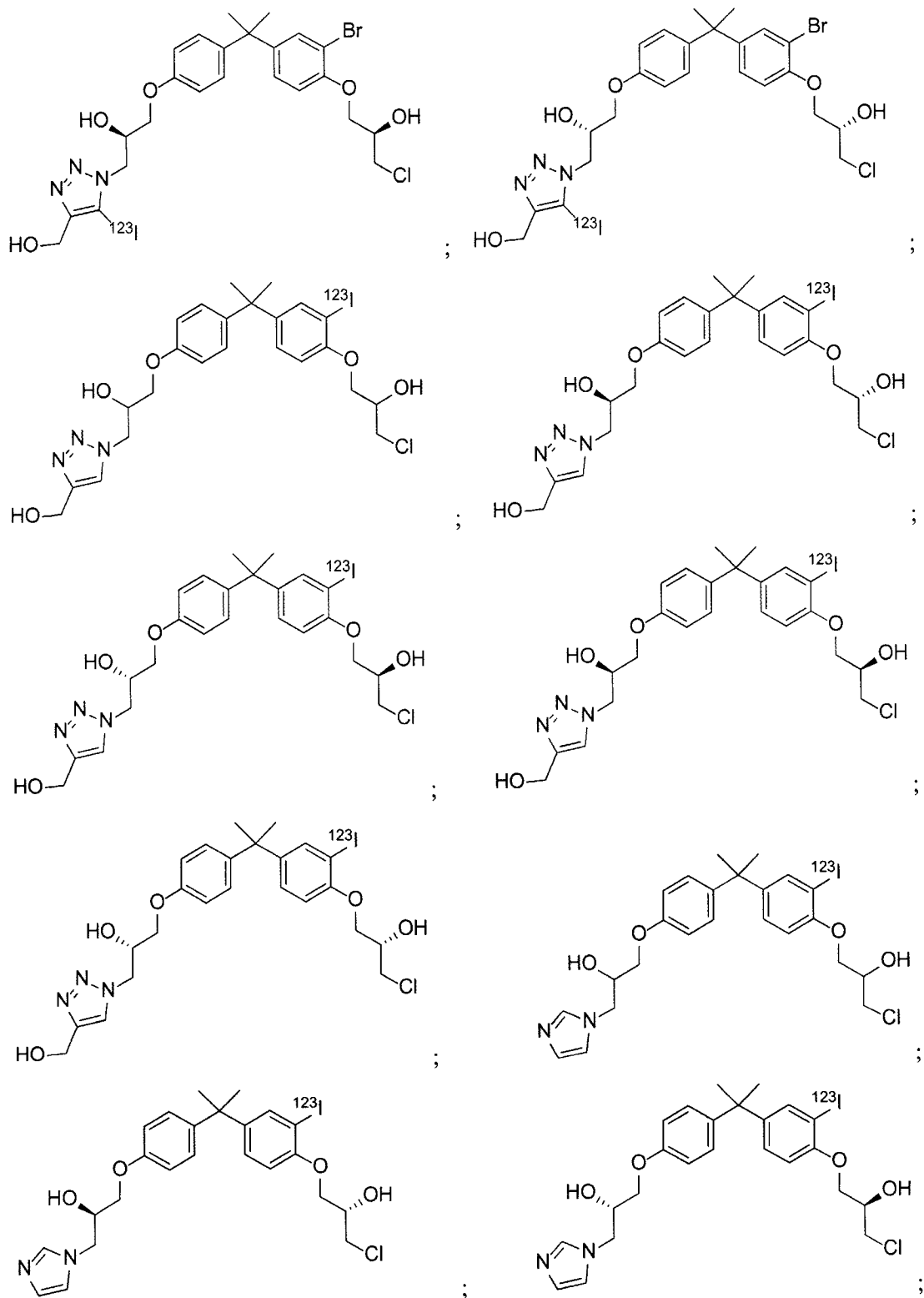
16. The compound of claim 1, wherein the compound has one of the following structures:



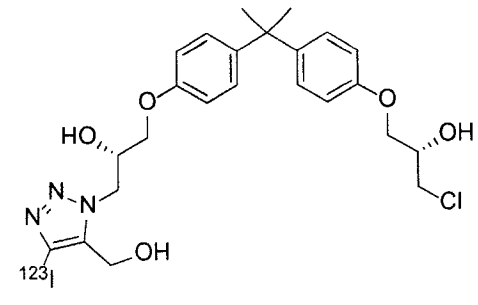
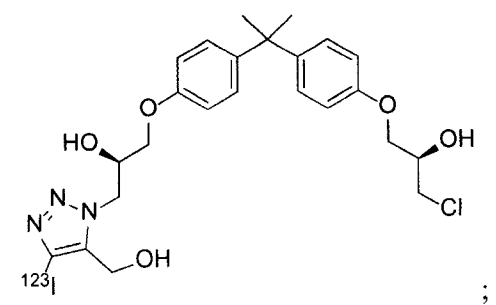
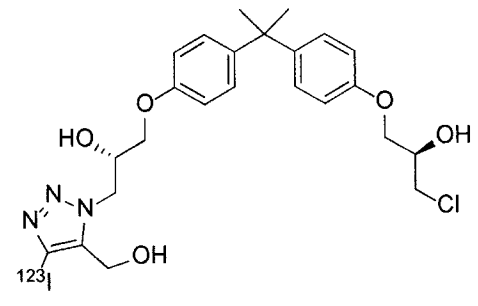
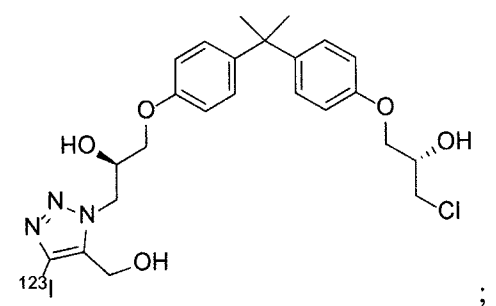
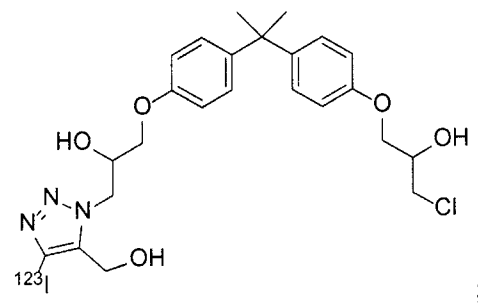
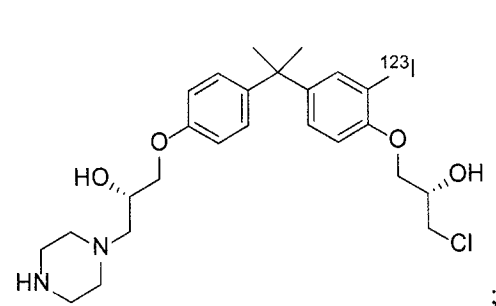
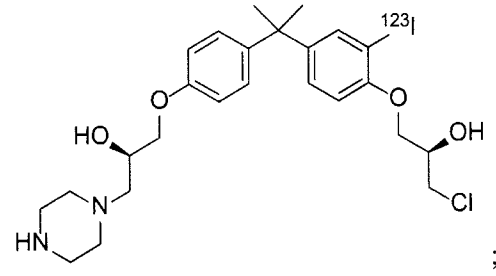
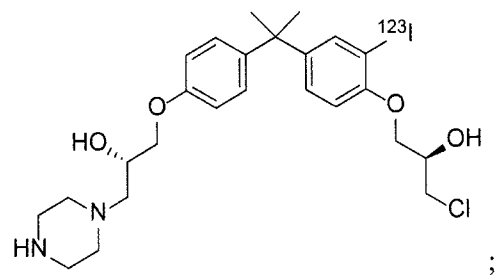
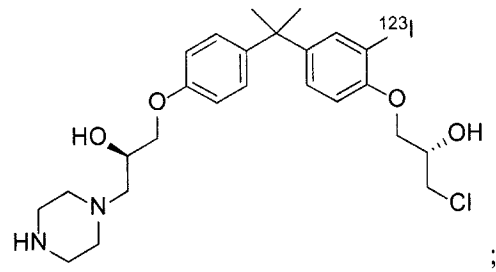
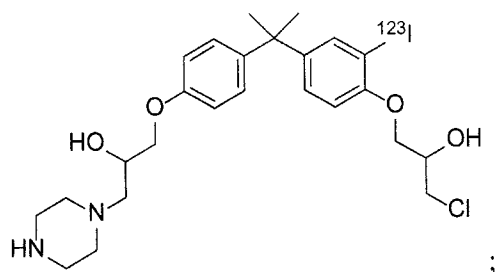


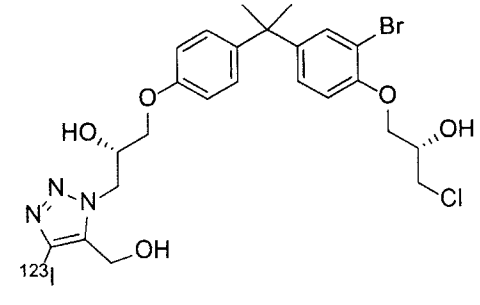
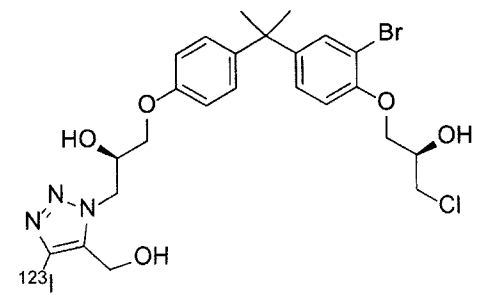
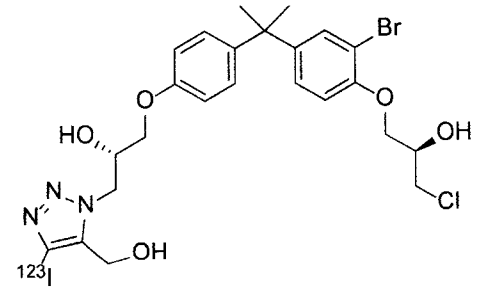
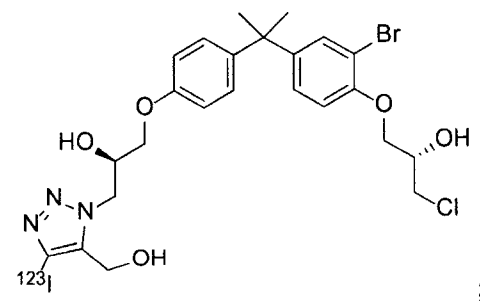
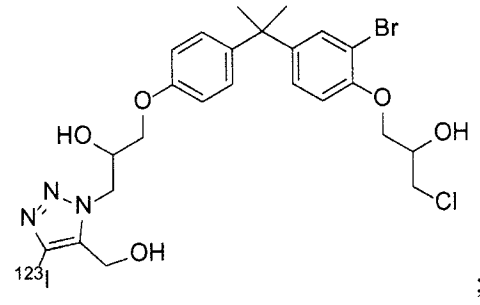
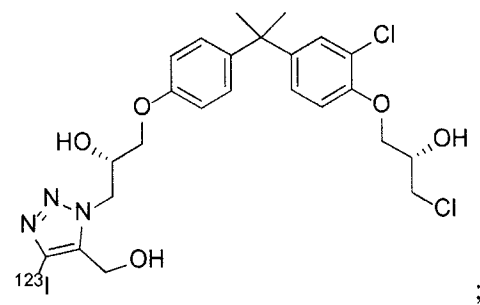
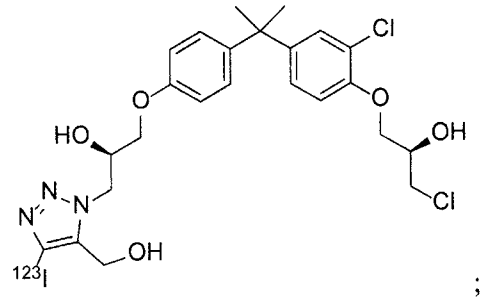
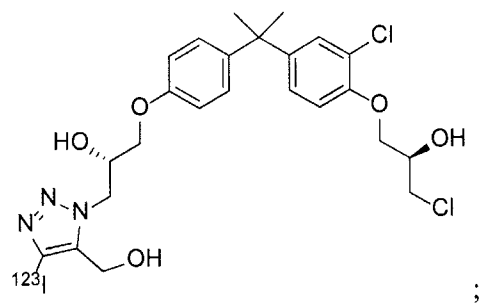
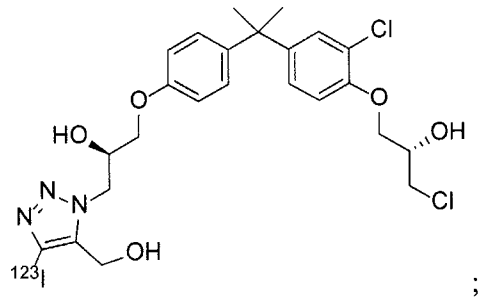
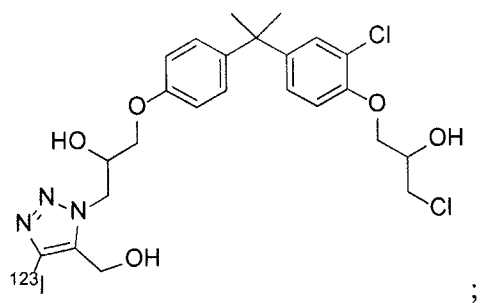


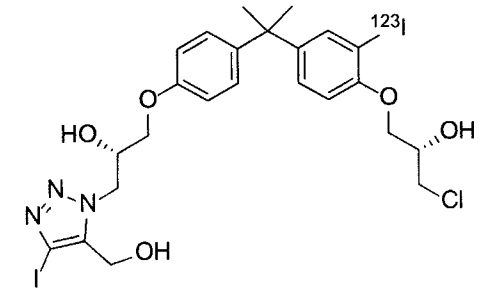
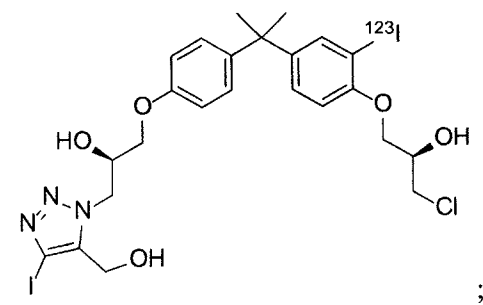
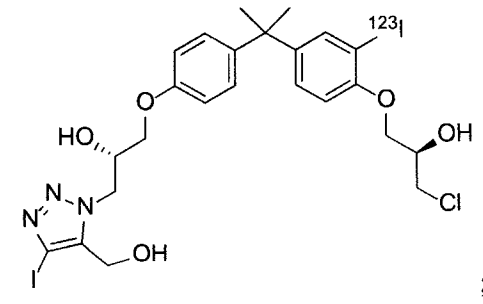
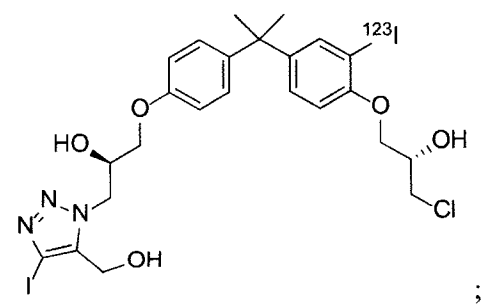
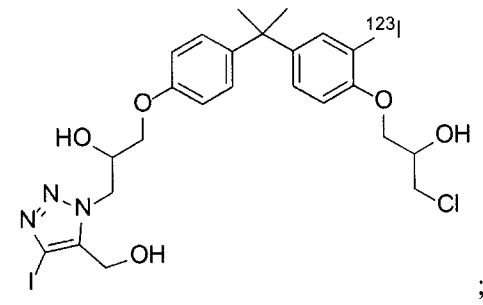
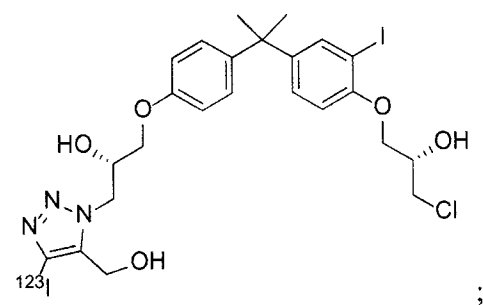
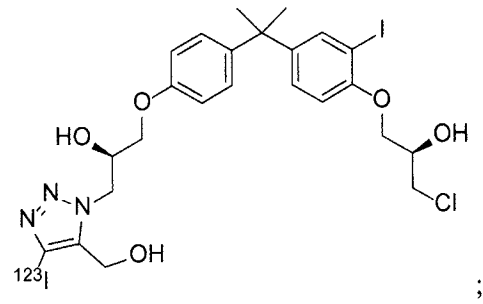
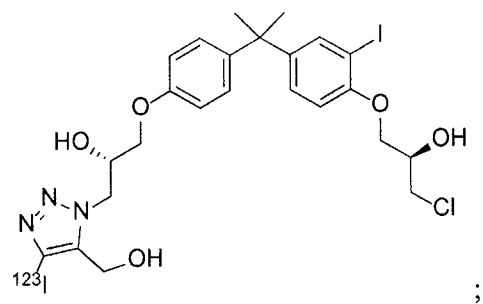
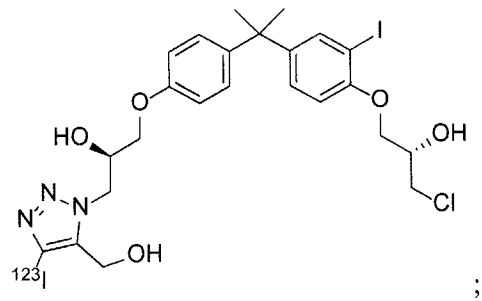
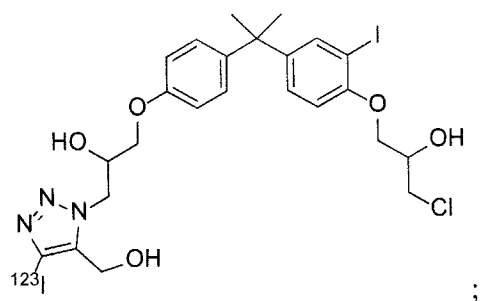


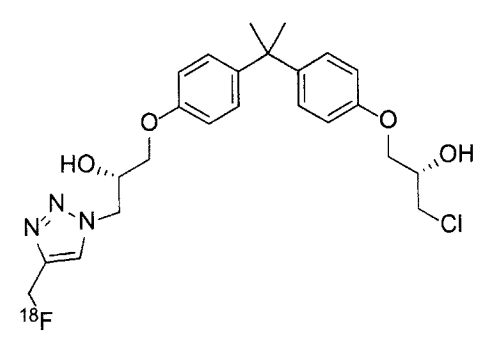
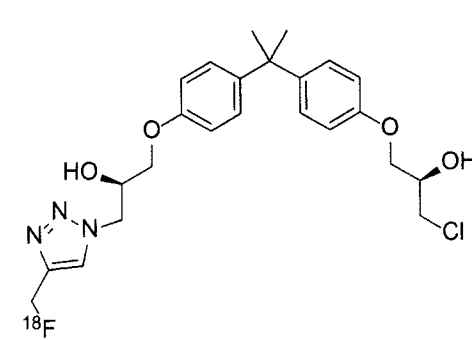
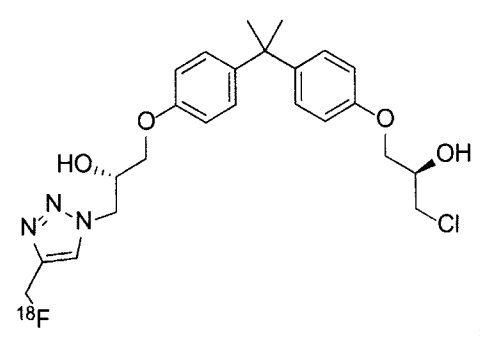
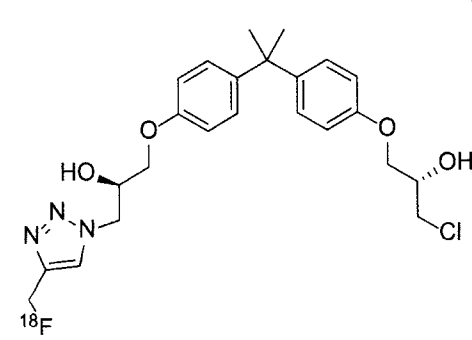
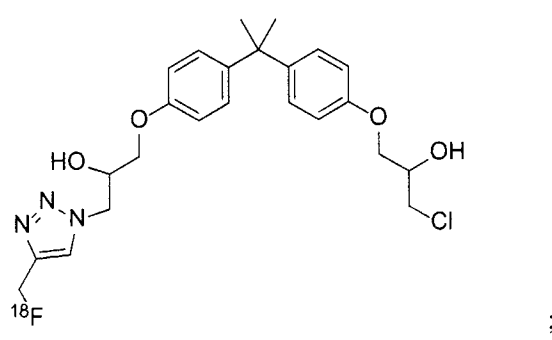
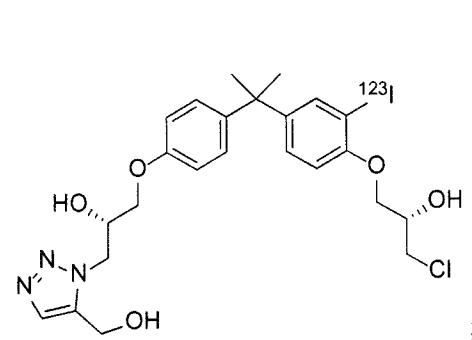
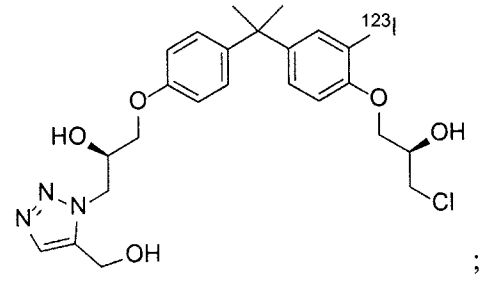
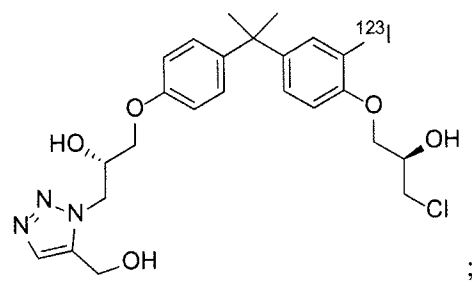
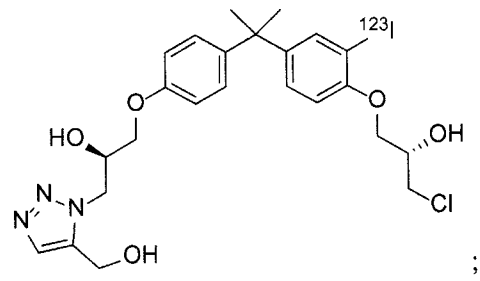
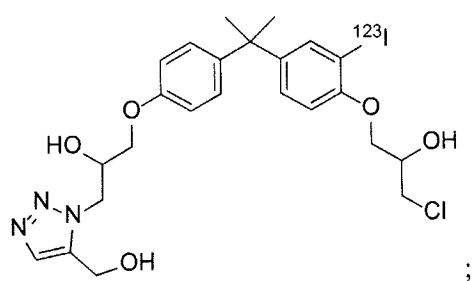


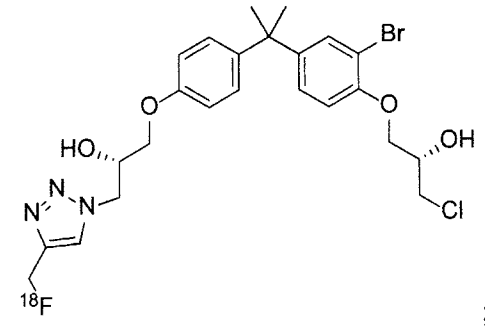
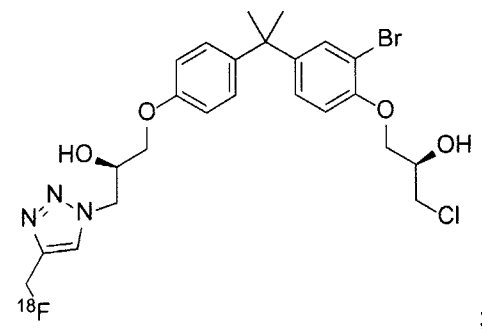
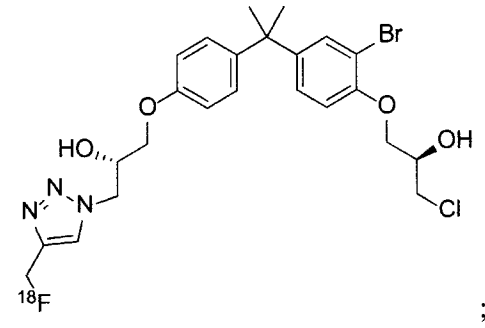
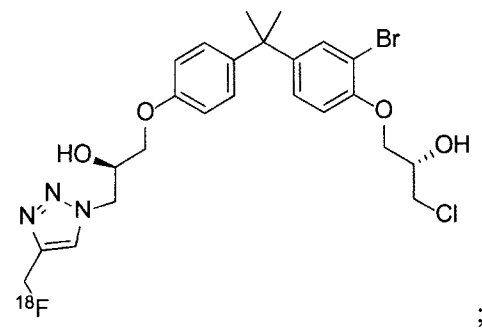
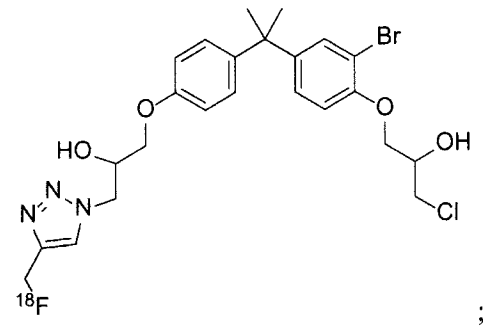
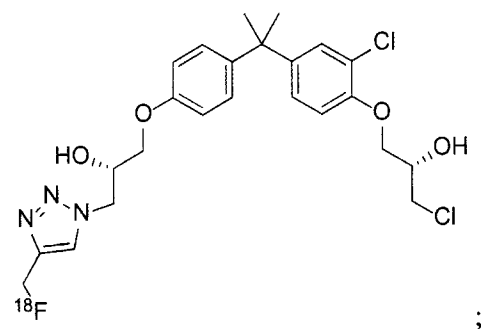
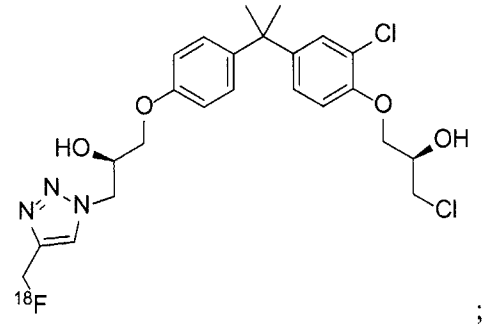
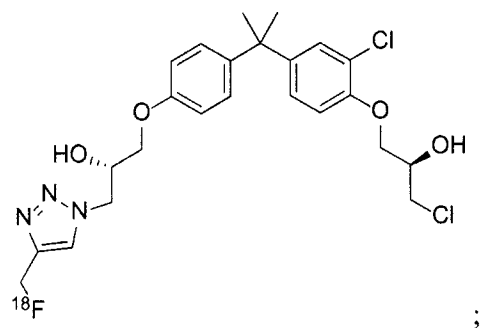
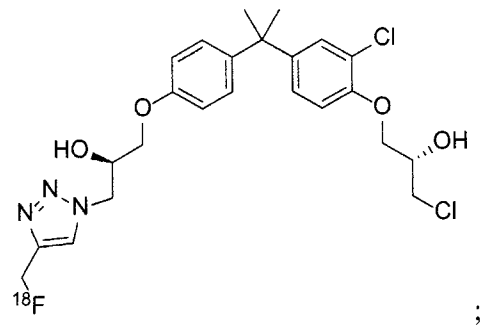
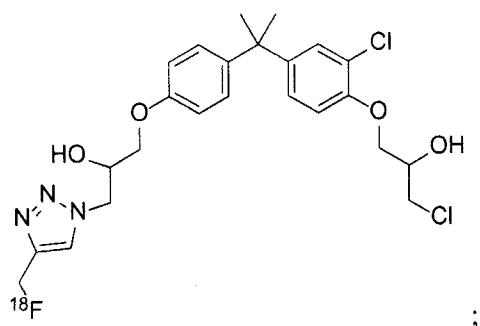


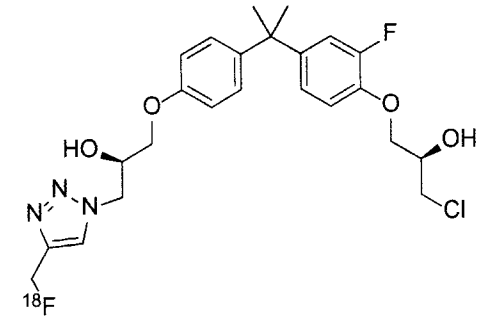
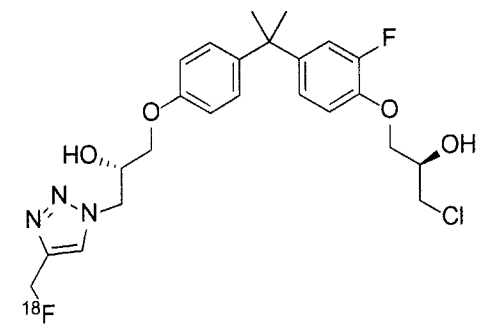
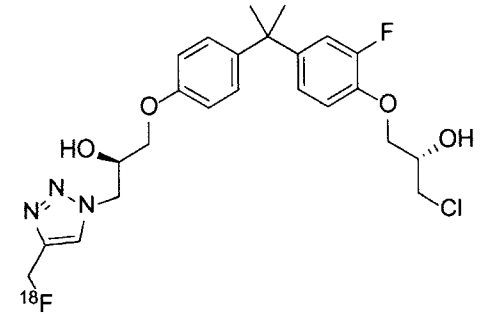
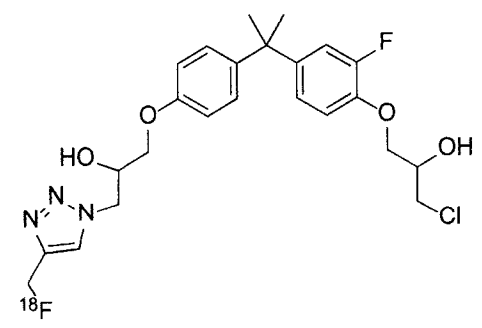
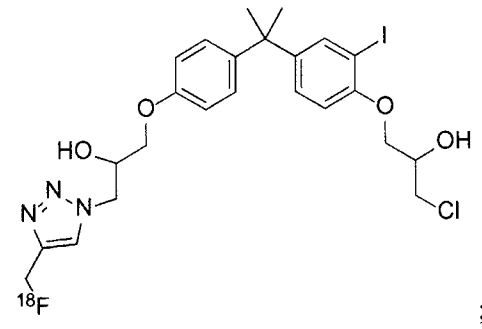
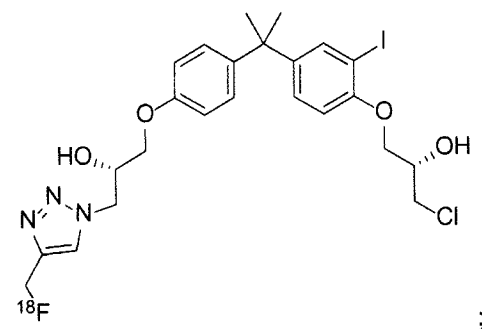
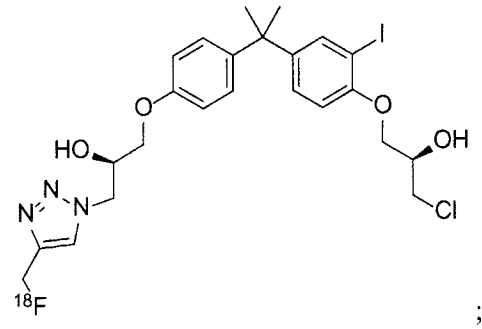
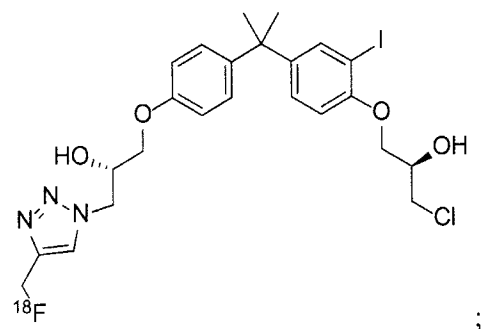
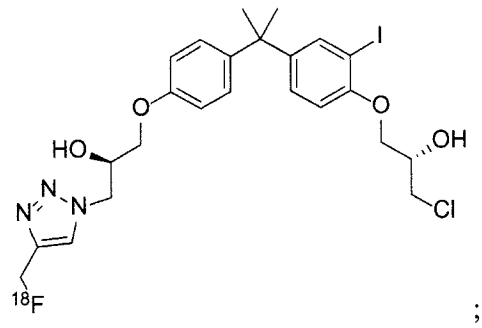
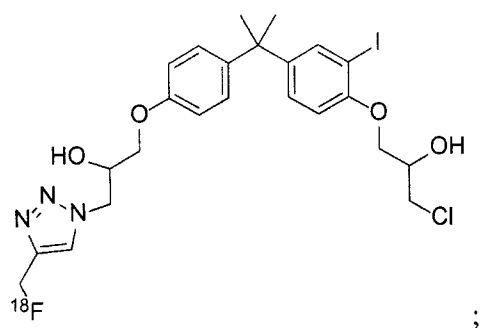


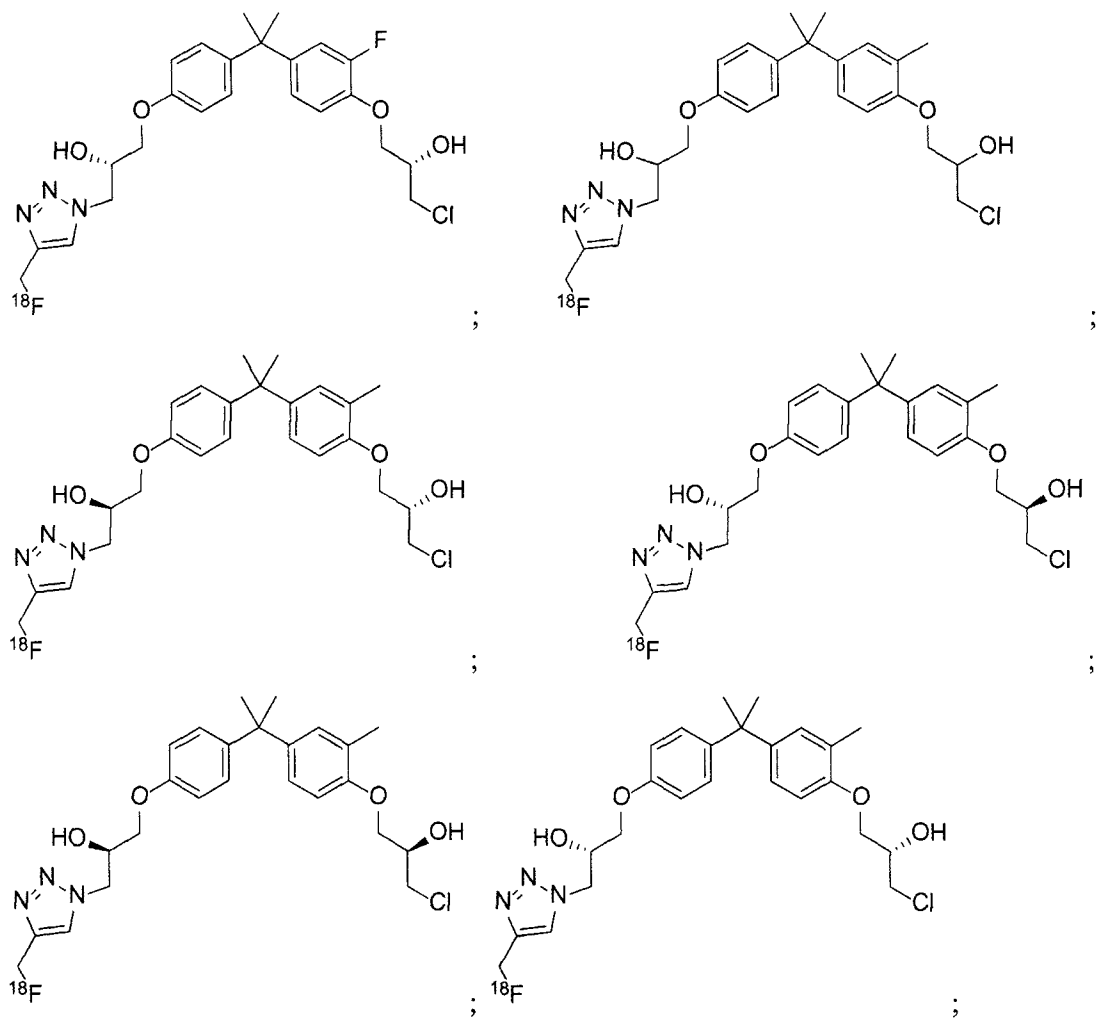






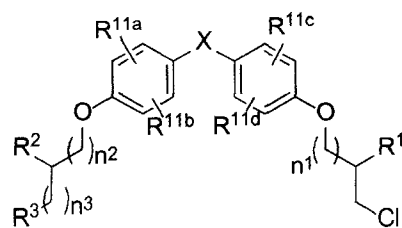






or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof.

17. A pharmaceutical composition, comprising: a compound of formula (I):



(I),

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-,
C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-

R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally substituted with one or more R⁶;

R⁵ is each independently H, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I, hydroxyl, oxo, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₆-C₁₂ aryl, wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -OS(O)₂-aryl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁸ and R⁹ are each independently H, halogen, -S(O)₀₋₂R⁵, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, aryl, aralkyl, C₁-C₁₀ acyl, or -NR⁵R⁵, or R⁸ and R⁹ can join to form a unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

R^{11a}, R^{11b}, R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ¹²³I, hydroxyl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl; -OR⁵, -OC(=O)R¹³, C₁-C₁₀ acyl, -S(O)₀₋₂R⁵, -NO₂-, CN, -NH₂, -NHR⁵, or -N(R⁵)₂;

R¹³ is each independently C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

n¹ and n² are each independently 0, 1, or 2; and

n³ is 0, 1, 2, 3, 4, or 5.

18. The pharmaceutical composition of claim 17, further comprising a pharmaceutically acceptable carrier.

19. The pharmaceutical composition of claim 17, further comprising a pharmaceutically acceptable carrier and an additional therapeutic agent.

20. The pharmaceutical composition of claim 19, wherein the additional therapeutic agent is for treating prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary

gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, or age-related macular degeneration.

21. The pharmaceutical composition of claim 19, wherein the additional therapeutic agent is enzalutamide, Galeterone, ARN-509; abiraterone, bicalutamide, nilutamide, flutamide, cyproterone acetate, docetaxel, Bevacizumab (Avastin), OSU-HDAC42, VITAXIN, sunitumib, ZD-4054, , Cabazitaxel (XRP-6258), MDX-010 (Ipilimumab), OGX 427, OGX 011, finasteride, dutasteride, turosteride, bexlosteride, izonsteride, FCE 28260, SKF105,111, ODM-201, radium 233, or related compounds thereof.

22. A method for modulating androgen receptor activity, comprising: administering a pharmaceutical composition according to claim 17 to a patient in need thereof.

23. A method for treating a condition or disease that is responsive to modulation of androgen receptor activity, comprising: administering a pharmaceutical composition according to claim 17 to a patient in need thereof,

wherein said condition or disease is selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, and age-related macular degeneration.

24. The method of claim 23, wherein the condition or disease is prostate cancer.

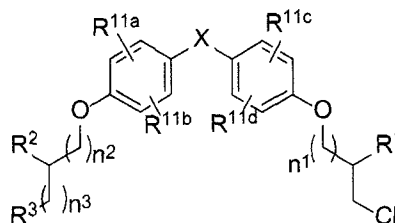
25. The method of claim 23, wherein the condition or disease is castration resistant prostate cancer.

26. The method of claim 23, wherein the condition or disease is androgen-dependent prostate cancer.

27. A method for treating a condition or disease selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, salivary gland carcinoma, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty, spinal and bulbar muscular atrophy, and age-related macular degeneration, comprising:

administering to a patient in need thereof a compound having the following structure

(I):



(I),

or a pharmaceutically acceptable salt, tautomer or stereoisomer thereof, wherein:

X is -O-, -S(O)₀₋₂-, -C(=O)-, -C(OR⁵)₂-, -C(OR⁵)(OC(=O)R¹³)-, -C(R⁸R⁹)-, -C(=CR⁸R⁹)-, -N(R⁹)-, -N(COR⁹)-, -CHNR⁸R⁹-, -C(=NR⁹)-, -C(=NOR⁵)-, -C(=N-NHR⁵)-,

R¹ and R² are each independently H, hydroxyl, -O-heterocyclyl, or -OC(=O)R¹³;

R³ is -N₃, aryl, carbocyclyl, heteroaryl or heterocyclyl which are optionally substituted with one or more R⁶;

R⁵ is each independently H, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁶ is each independently selected from the group consisting of H, F, Cl, Br, I, ¹²³I, hydroxyl, oxo, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₆-C₁₂ aryl, wherein each R⁶ is optionally substituted with one or more of halogen, ¹²³I, ¹⁸F, hydroxyl, -OS(O)₂-aryl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl;

R⁸ and R⁹ are each independently H, halogen, -S(O)₀₋₂R⁵, C₁-C₁₀ alkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, aryl, aralkyl, C₁-C₁₀ acyl, or -NR⁵R⁵, or R⁸ and R⁹ can join to form a unsubstituted or substituted mono-, bi-, or tri-cyclic carbocycle or heterocycle containing from 3 to 20 carbon atoms;

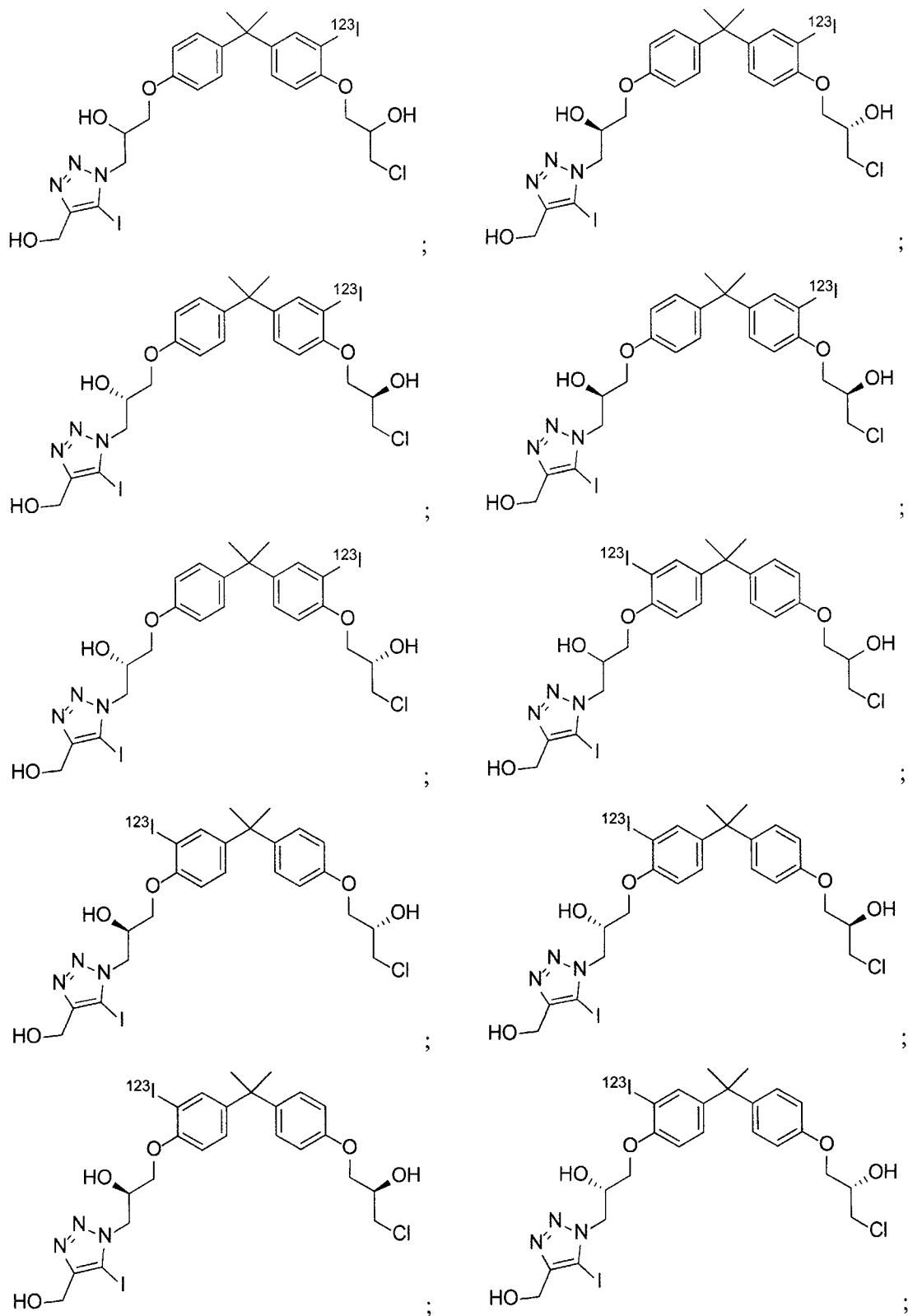
R^{11a} , R^{11b} , R^{11c} and R^{11d} are each independently H, F, Cl, Br, I, ^{123}I , hydroxyl, $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, or $\text{C}_2\text{-C}_6$ alkynyl; $-\text{OR}^5$, $-\text{OC}(=\text{O})\text{R}^{13}$, $\text{C}_1\text{-C}_{10}$ acyl, $-\text{S}(\text{O})_{0-2}\text{R}^5$, $-\text{NO}_2$, $-\text{CN}$, $-\text{NH}_2$, $-\text{NHR}^5$, or $-\text{N}(\text{R}^5)_2$;

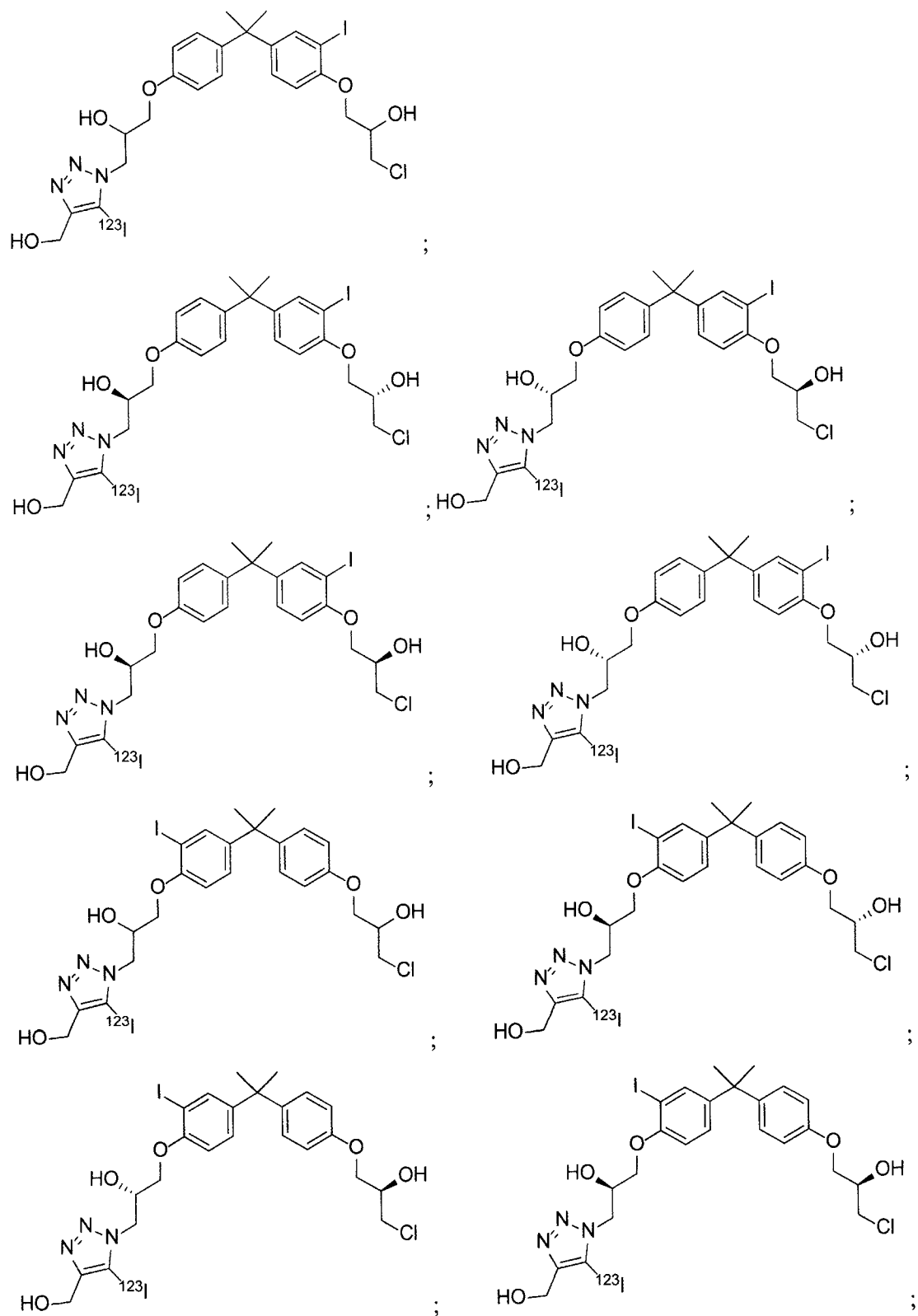
R^{13} is each independently $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, or $\text{C}_2\text{-C}_6$ alkynyl;

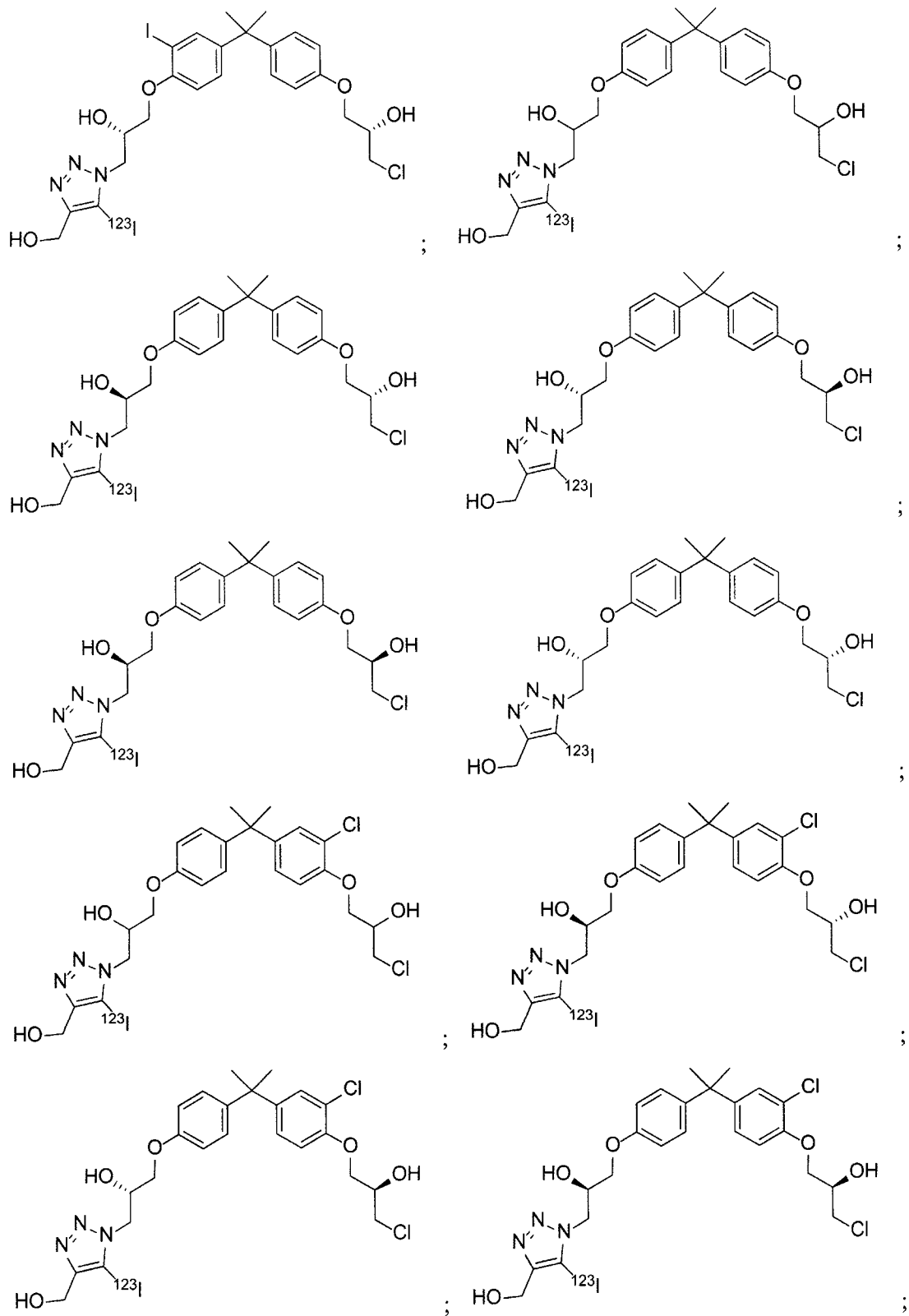
n^1 and n^2 are each independently 0, 1, or 2; and

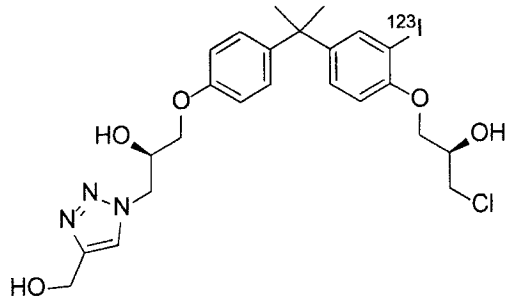
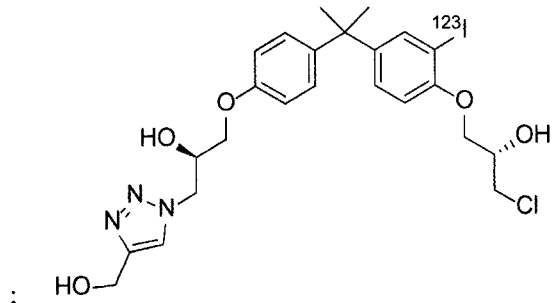
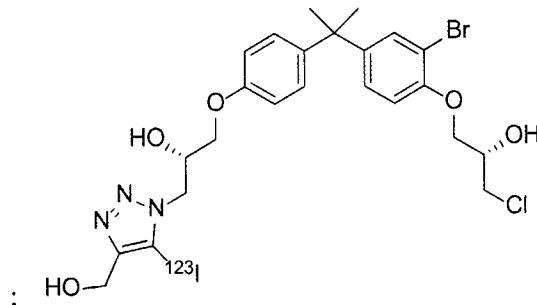
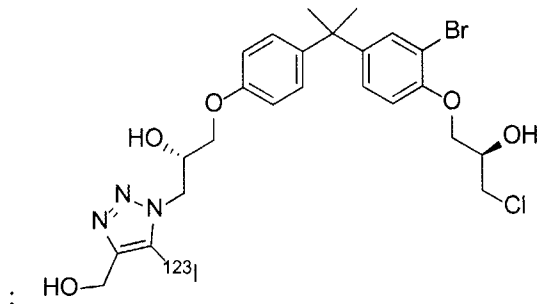
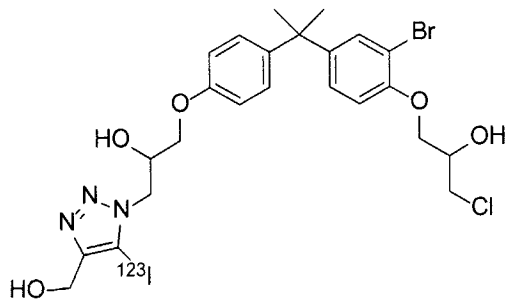
n^3 is 0, 1, 2, 3, 4, or 5.

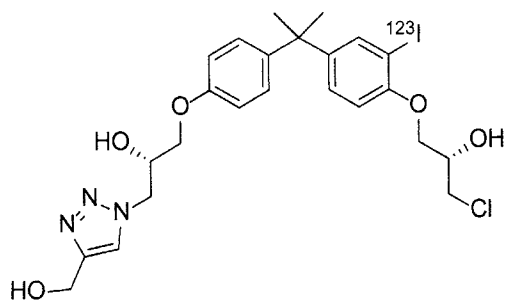
28. The method of claim 27, wherein the condition or disease is prostate cancer.
29. The method of claim 27, wherein the condition or disease is castration resistant prostate cancer or androgen-dependent prostate cancer.
30. A method of imaging cancer, the method comprising administering a compound of claim 1 to a subject and detecting the presence or absence of cancer by use of SPECT or PET.
31. The method of claim 30, wherein the method identifies the presence or absence of a tumor.
32. The method of claim 30, wherein the method identifies the location of a tumor.
33. The method of claim 30, wherein the cancer is prostate cancer.
34. The method of claim 30, wherein the compound of claim 1 is selected from the group consisting of:



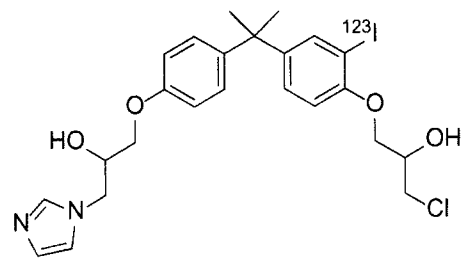




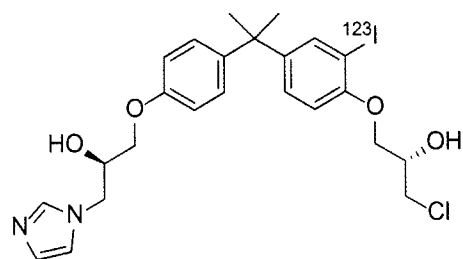




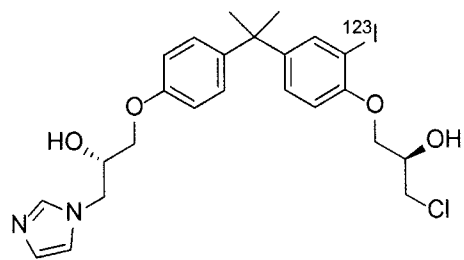
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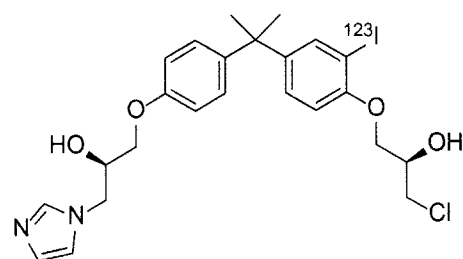
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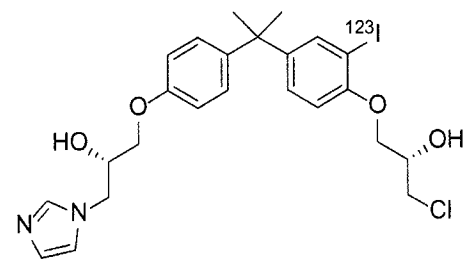
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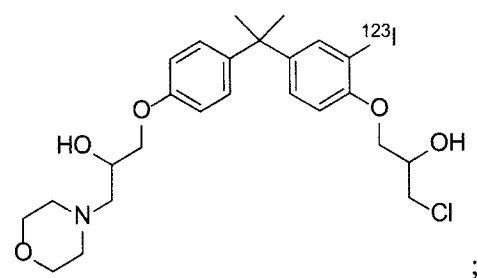
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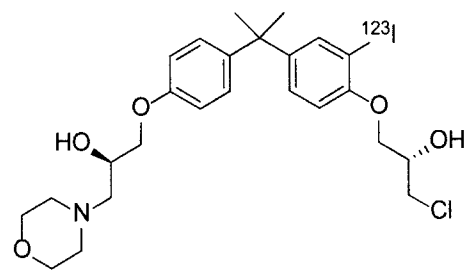
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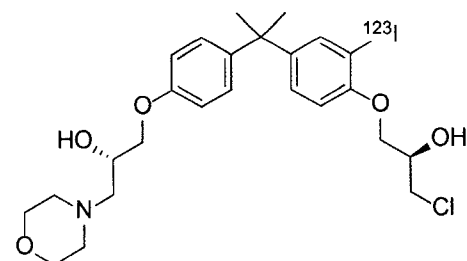
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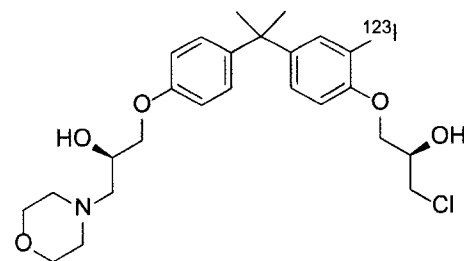
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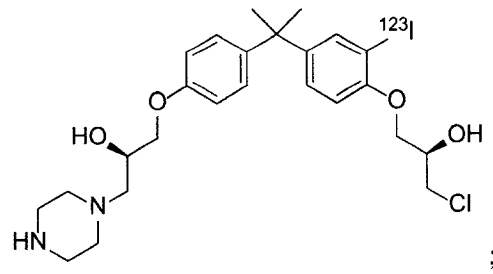
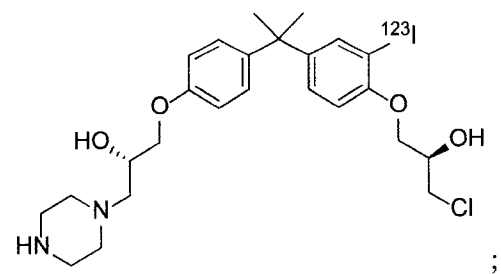
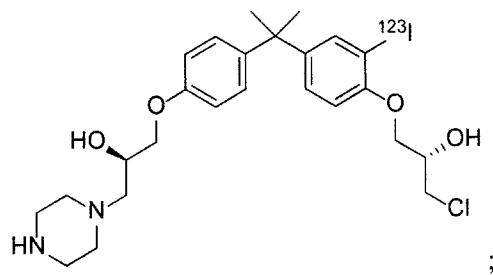
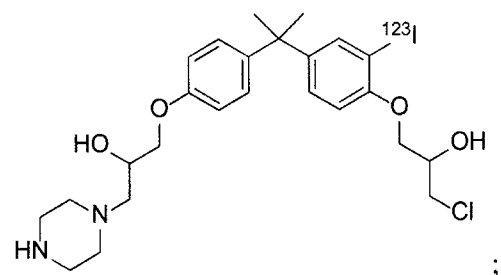
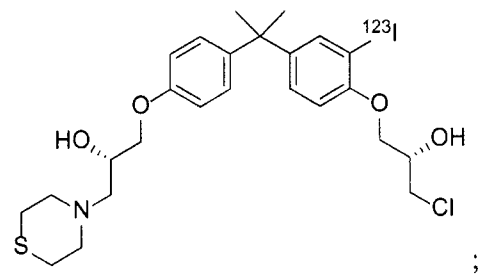
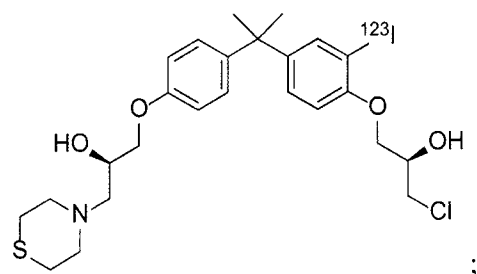
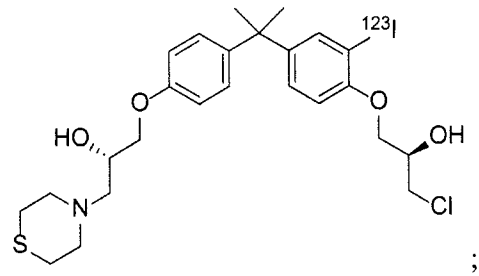
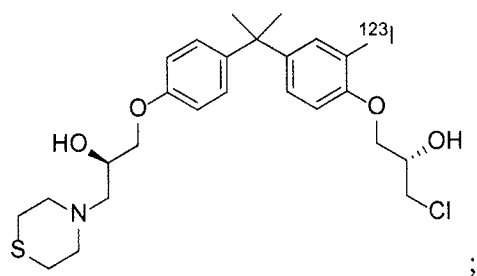
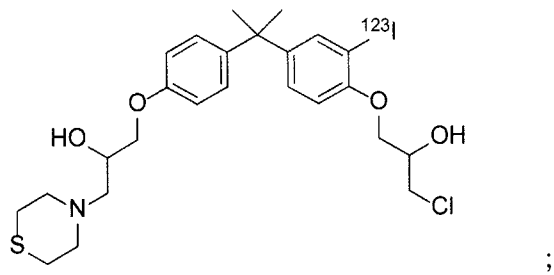
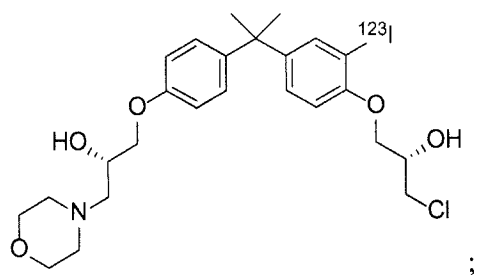
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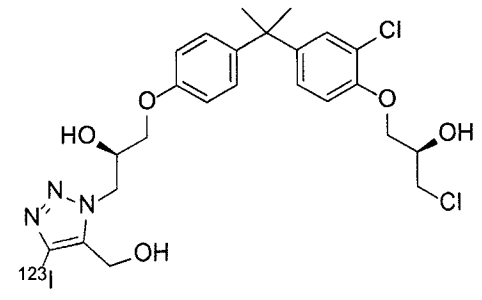
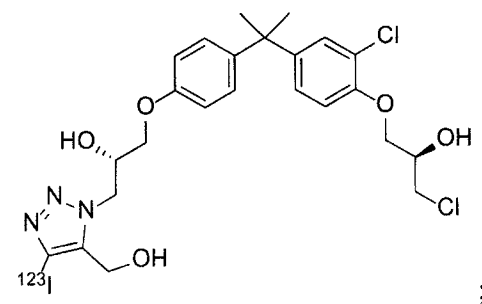
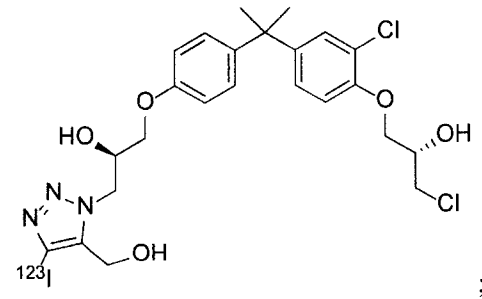
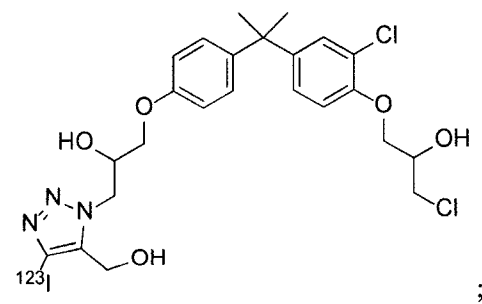
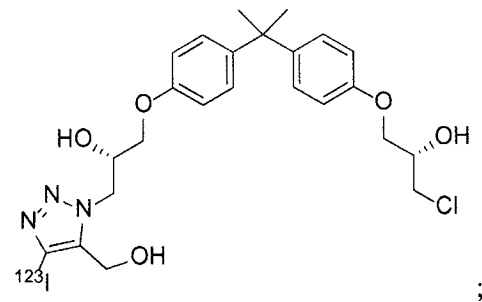
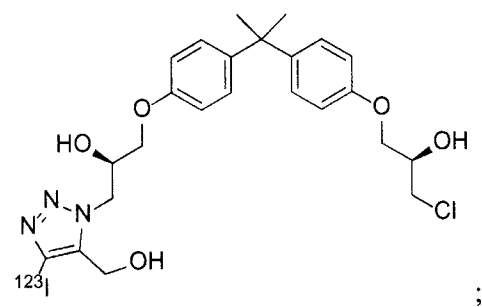
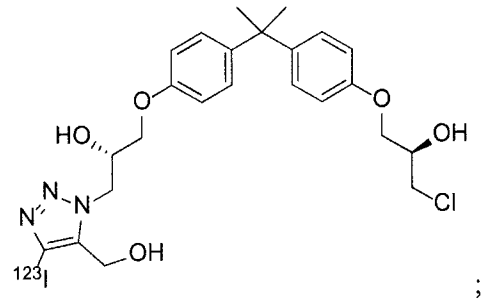
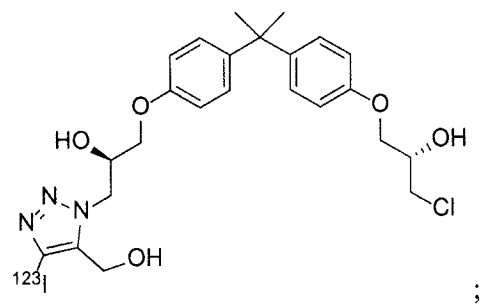
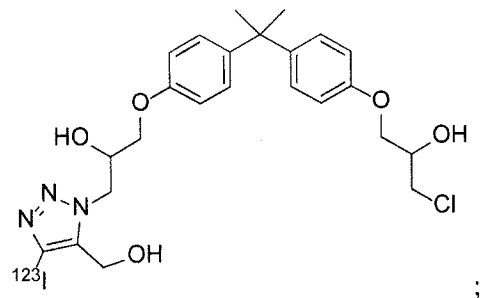
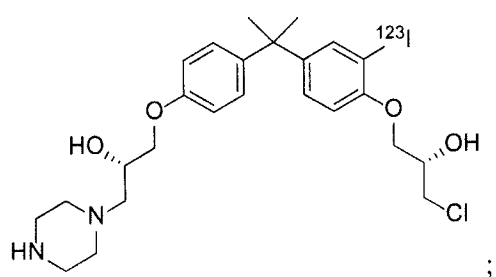


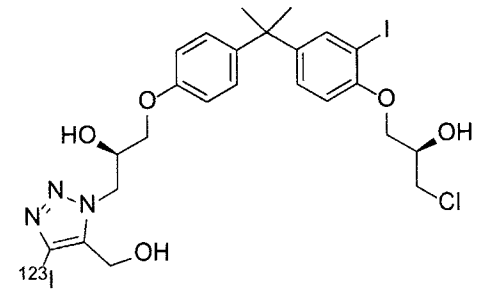
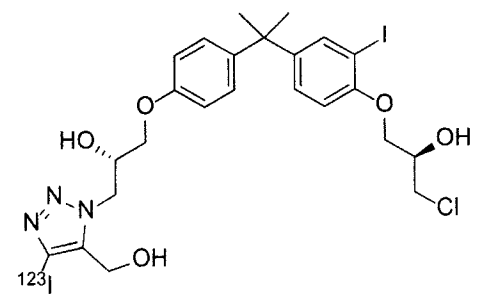
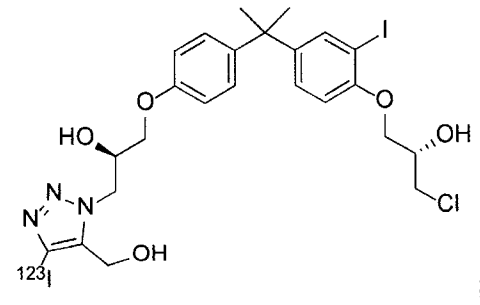
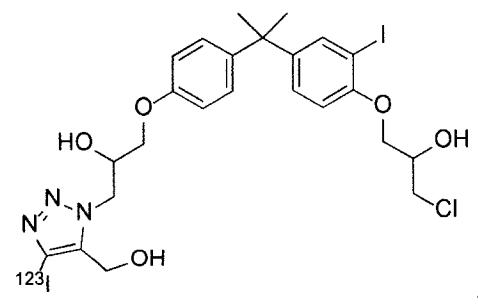
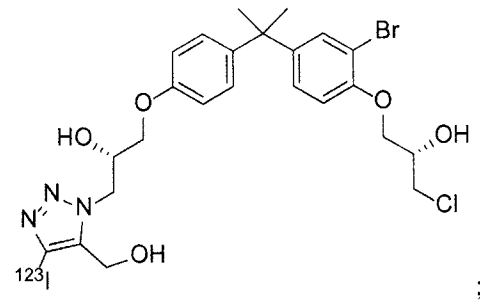
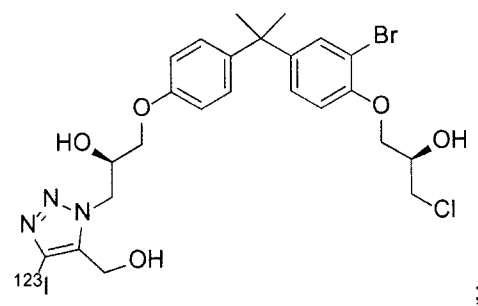
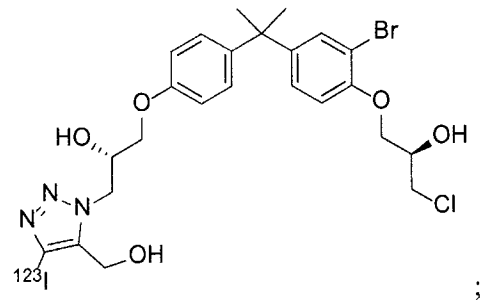
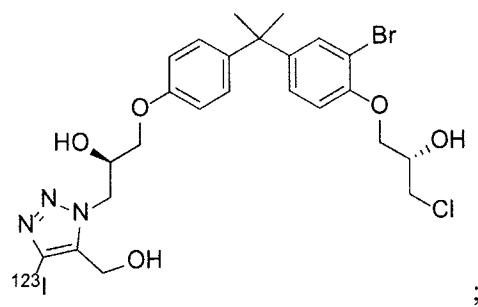
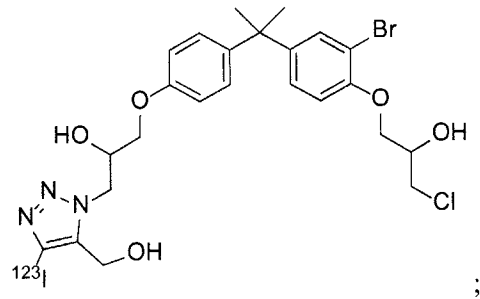
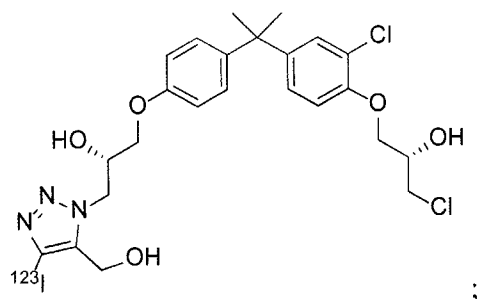
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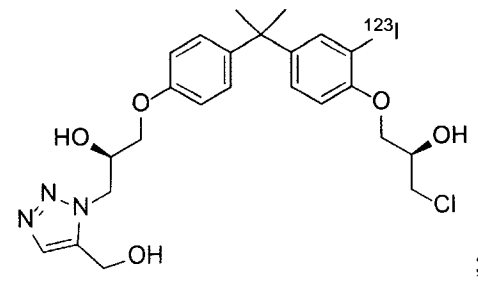
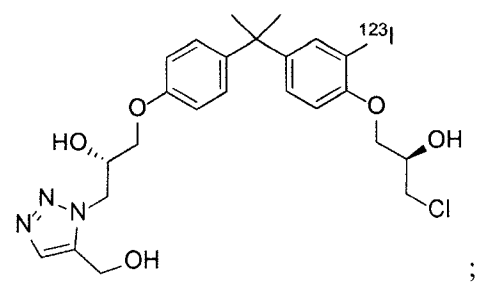
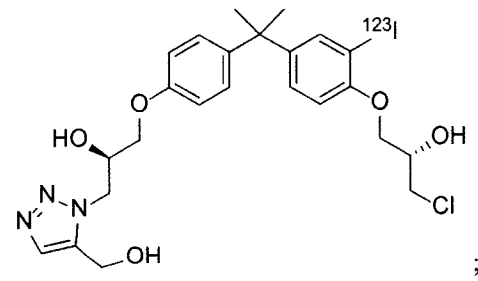
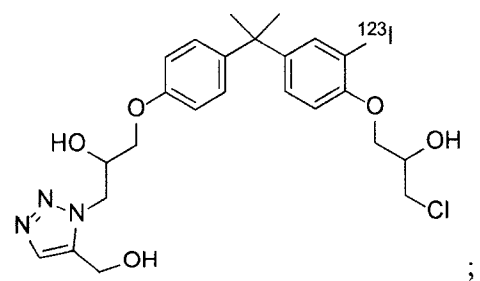
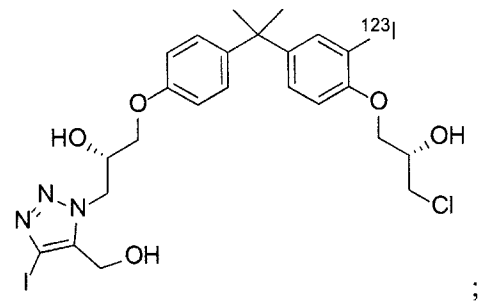
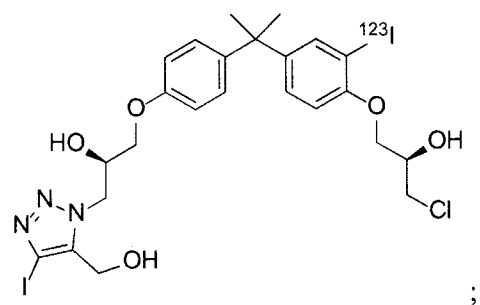
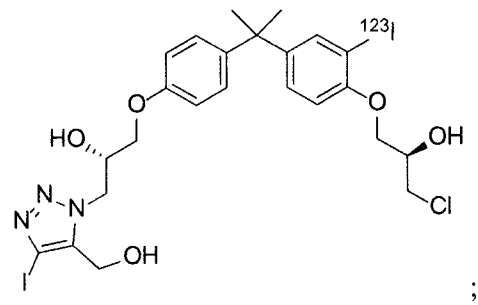
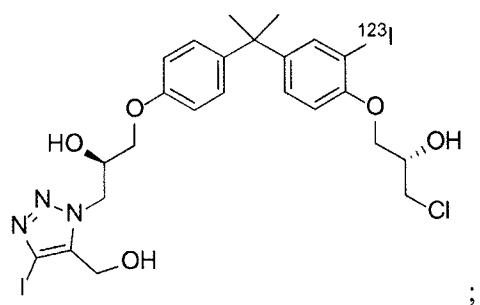
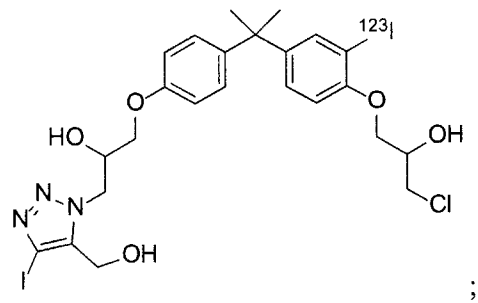
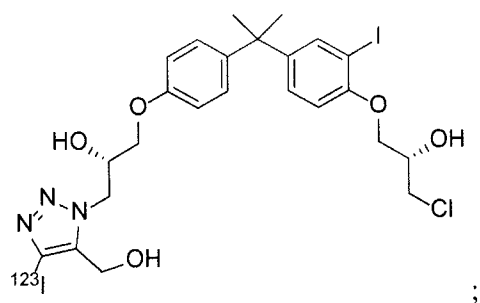


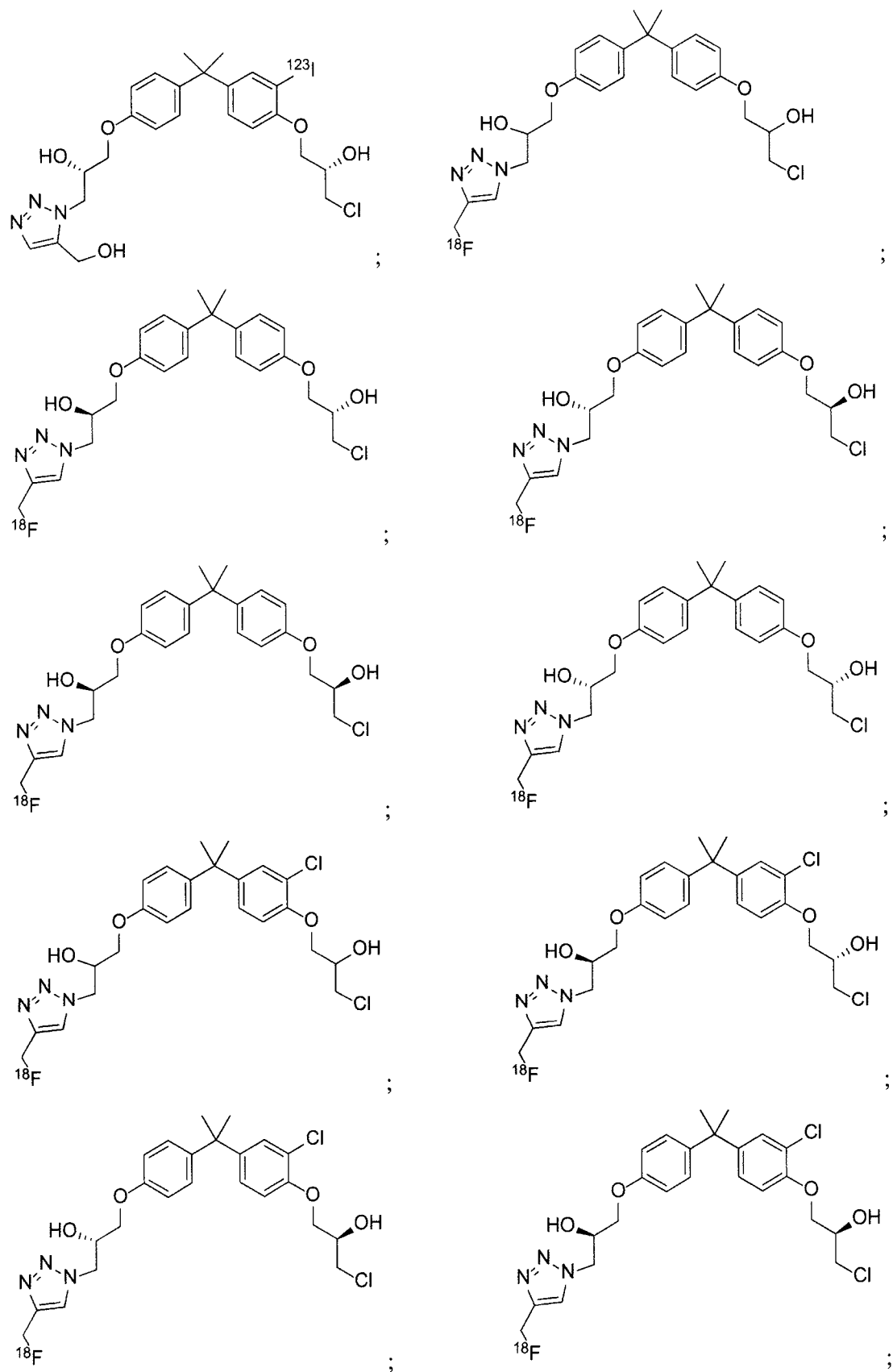
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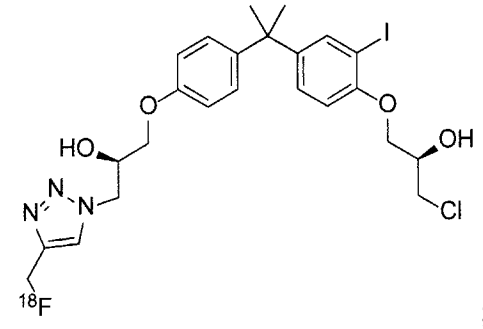
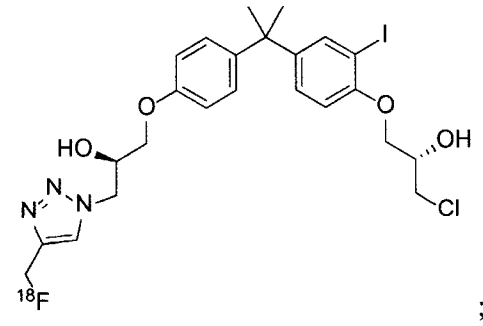
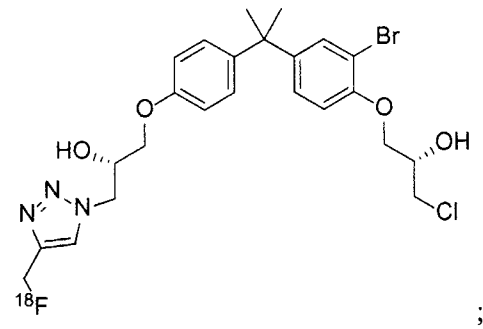
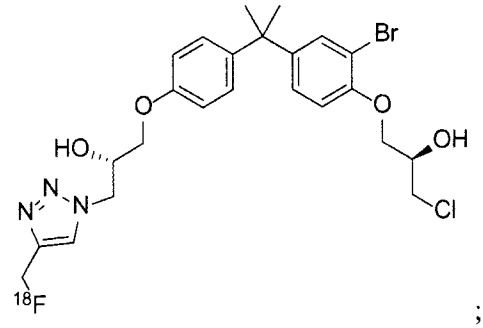
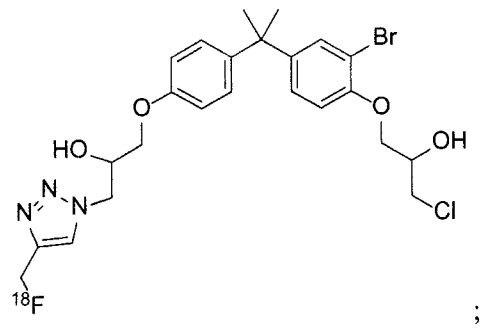


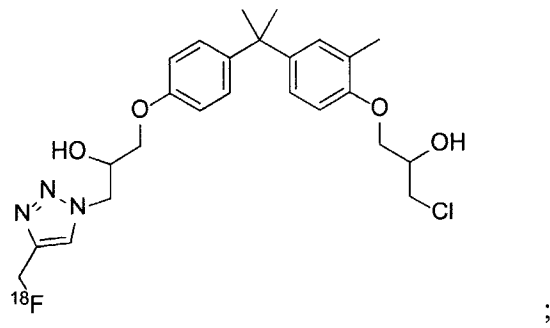
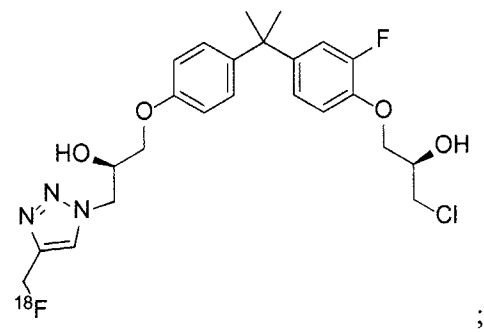
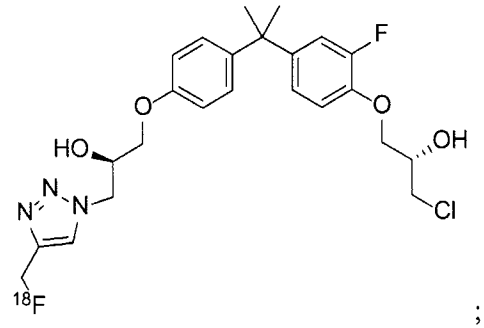
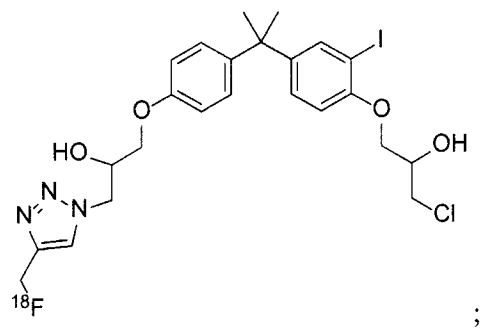


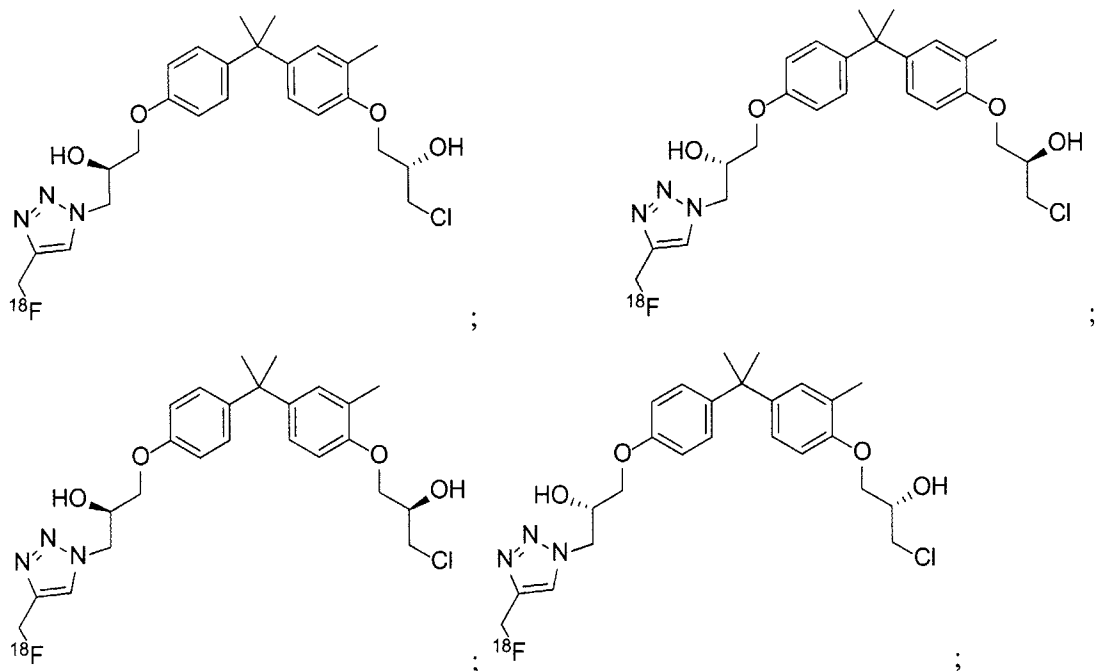








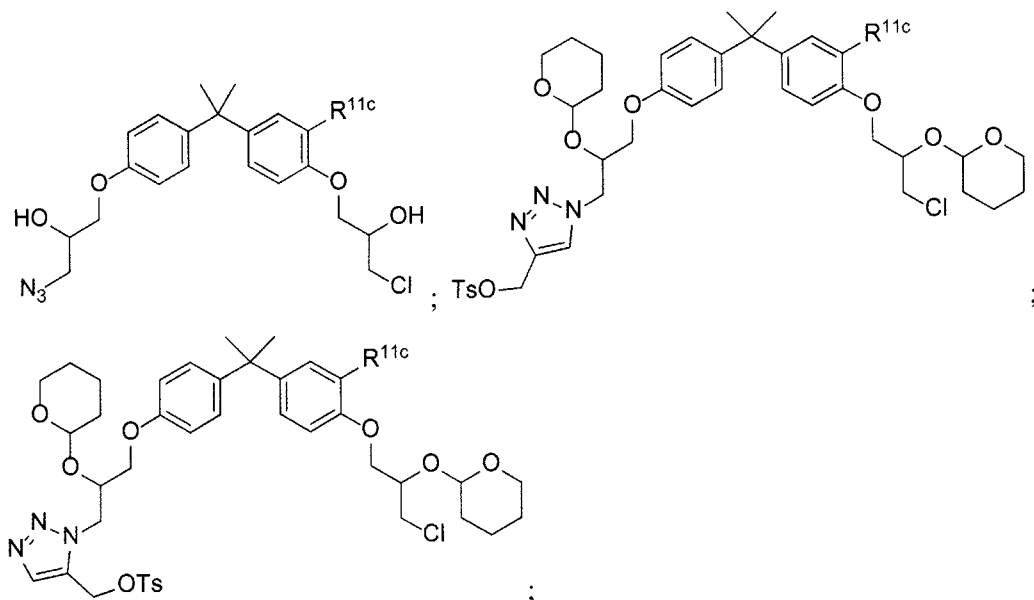




or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof.

35. A method of imaging cancer, the method comprising administering the pharmaceutical composition of claim 17 to a subject and detecting the presence or absence of cancer by use of SPECT or PET.

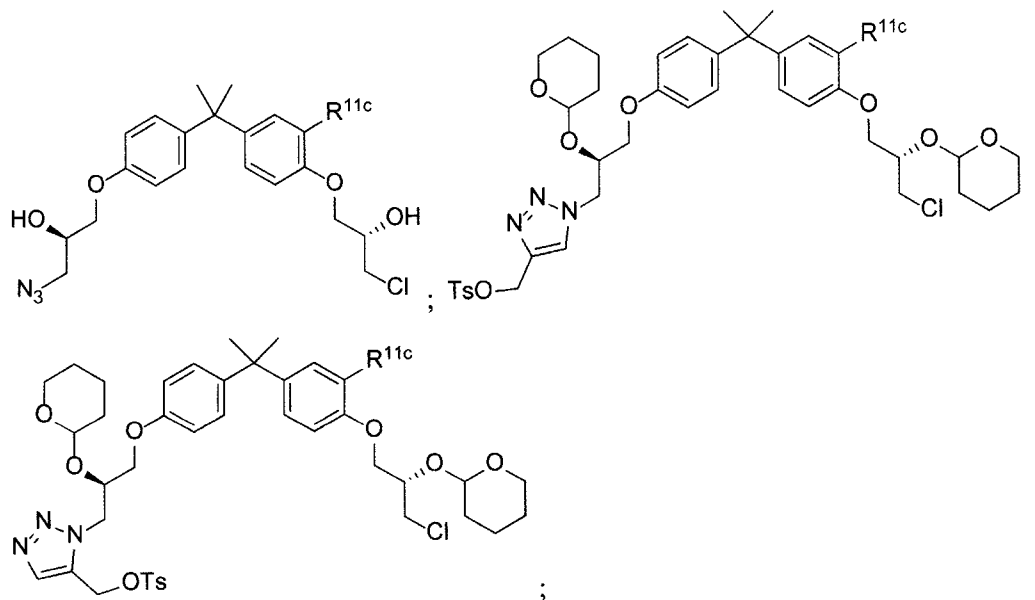
36. The compound of claim 1, wherein the compound has one of the following structures:



or a pharmaceutically acceptable salt, tautomer, or stereoisomer thereof;

wherein R^{11c} is H, Cl, Br, I, F, or C_1 - C_6 alkyl.

37. The compound of claim 1, wherein the compound has one of the following structures:



or a pharmaceutically acceptable salt thereof;

wherein R^{11c} is H, Cl, Br, I, F, or C_1 - C_6 alkyl.

FIG. 1

¹H

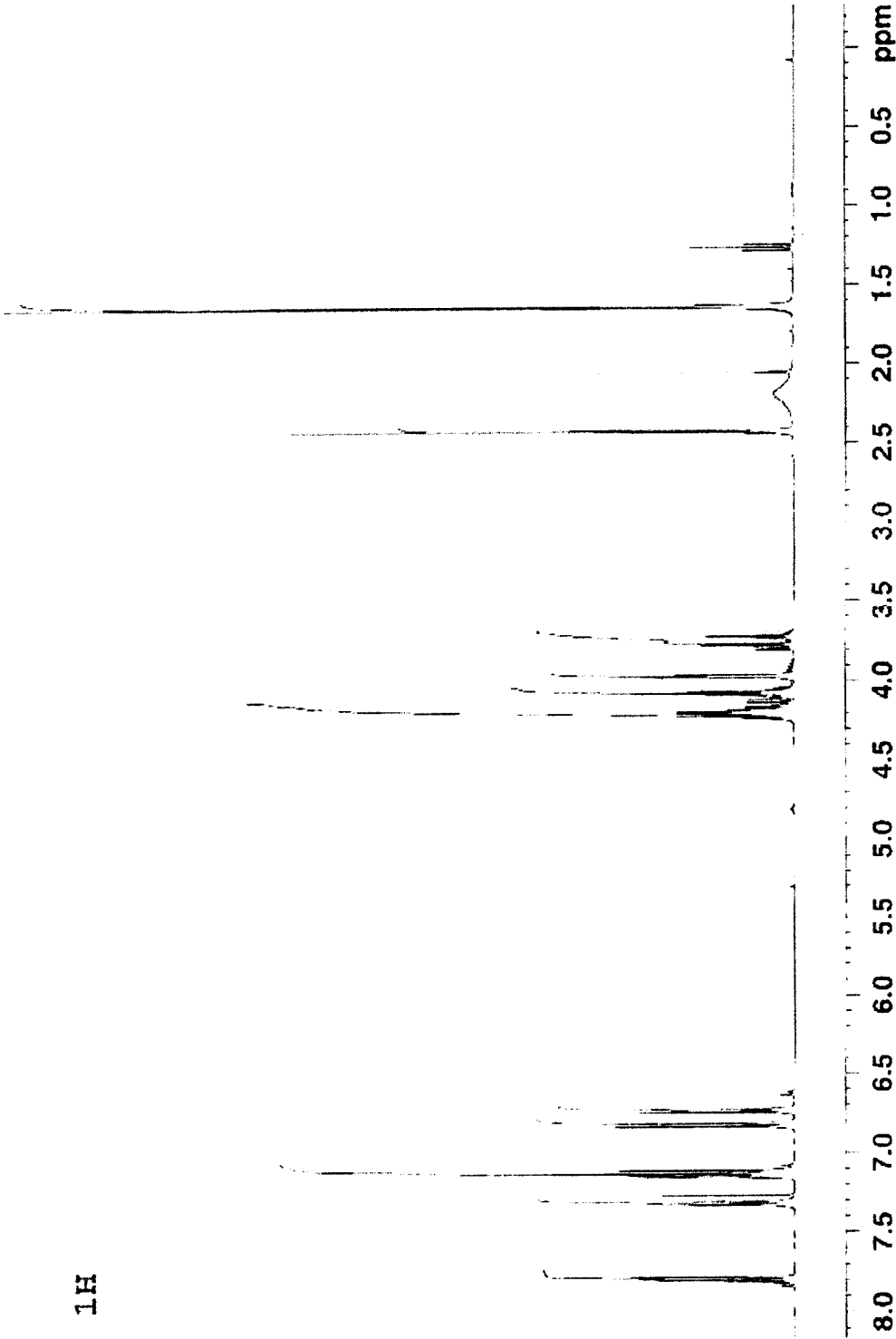


FIG. 2

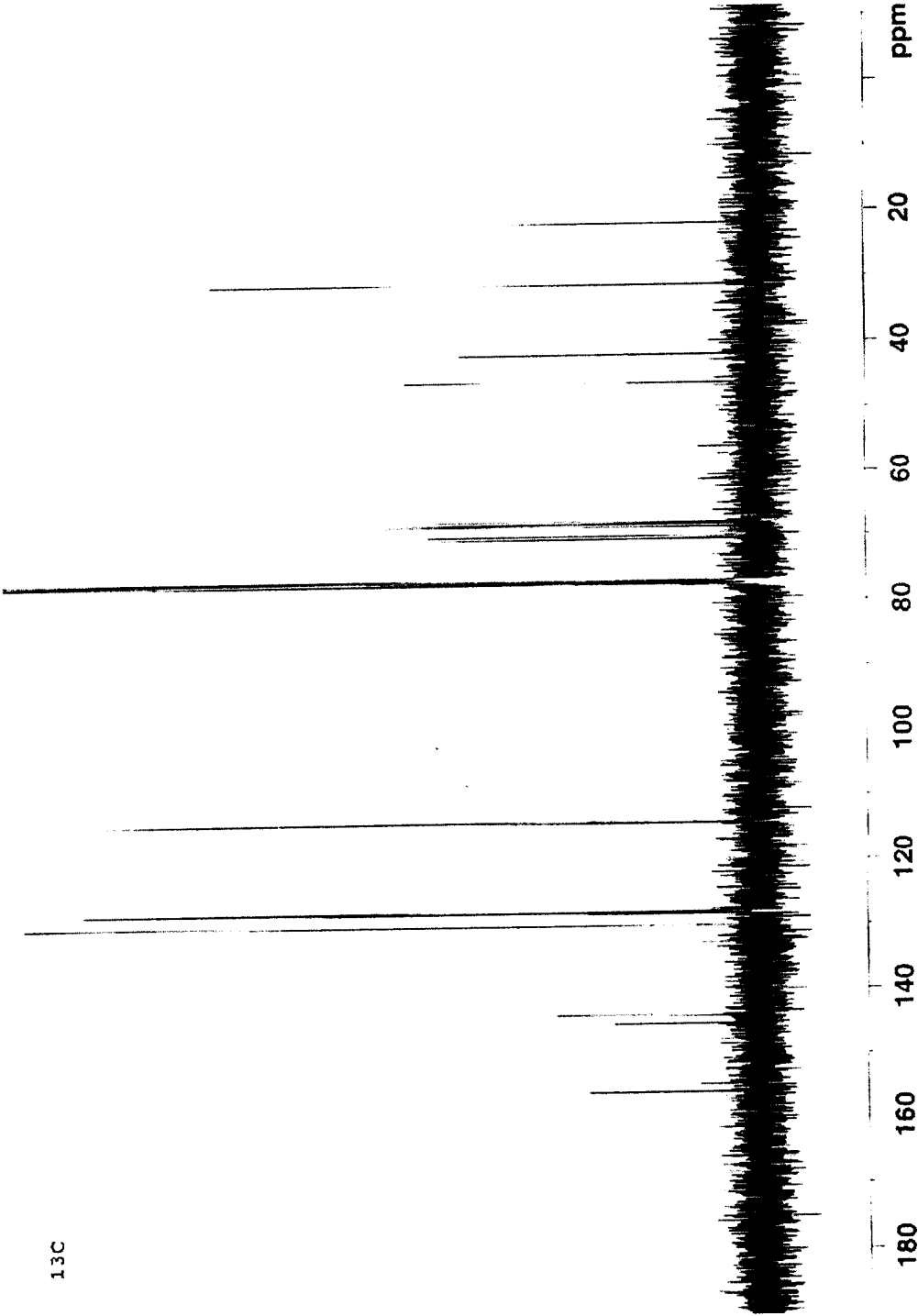


FIG. 3

¹H

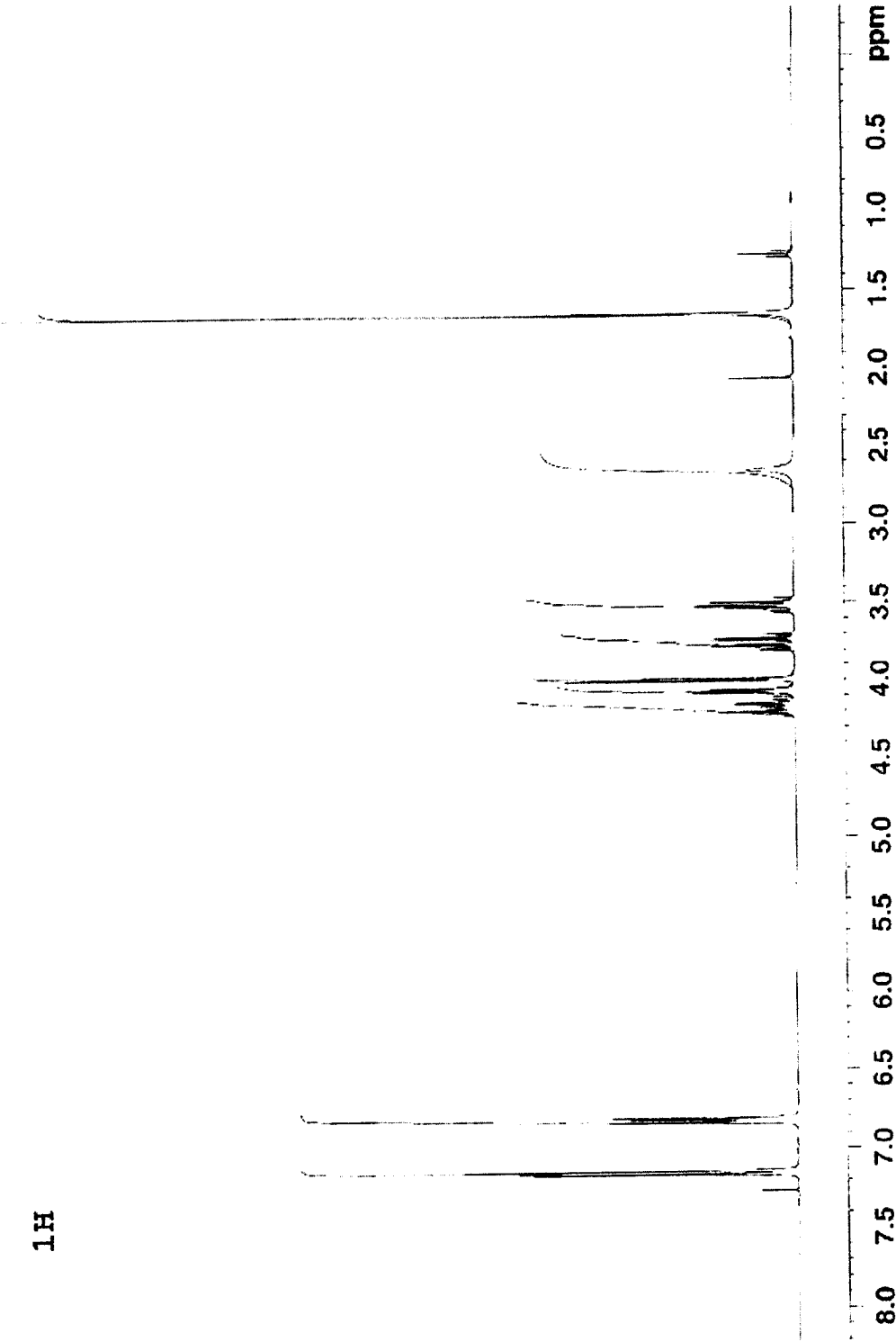


FIG. 4
¹³C

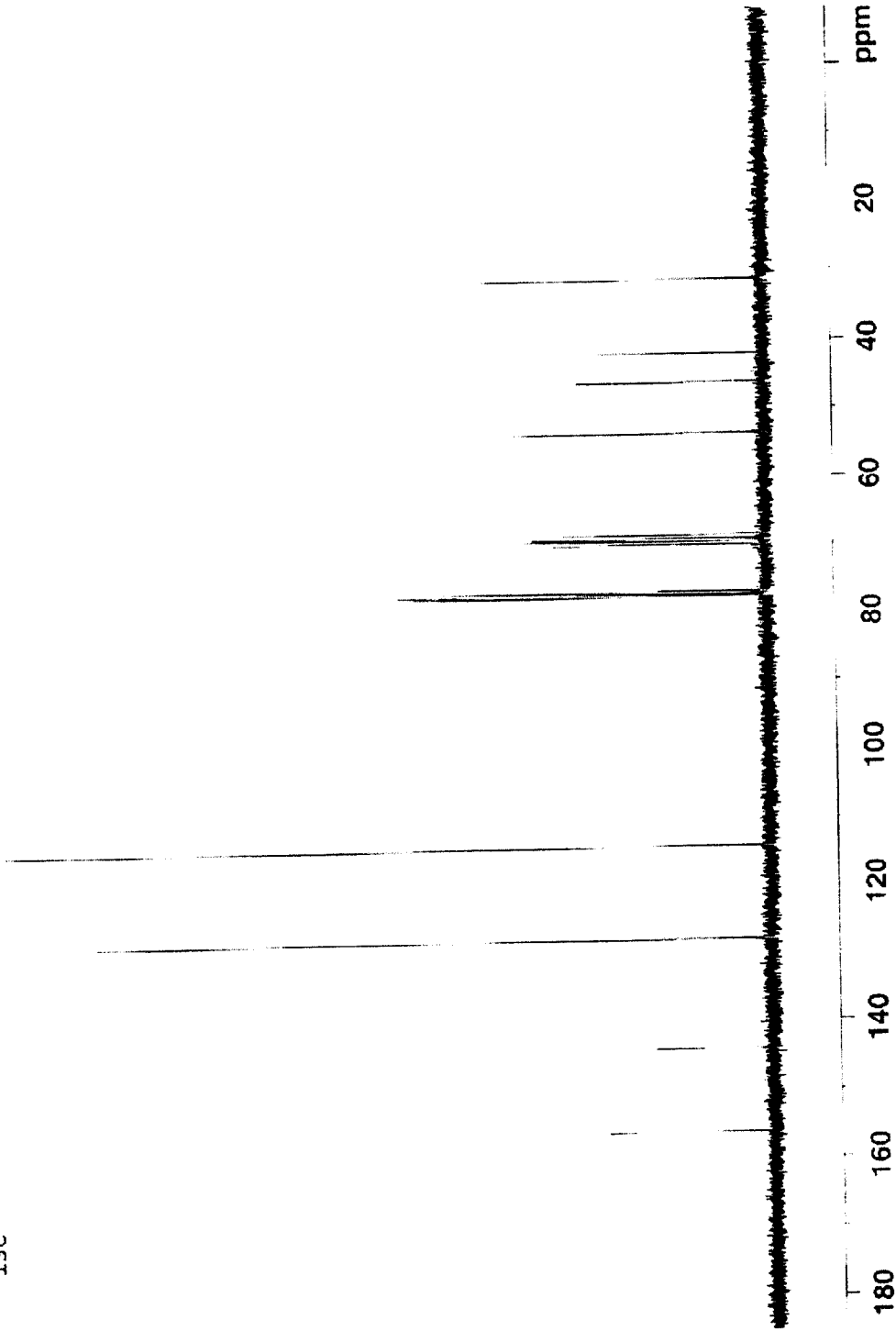


FIG. 5
1H

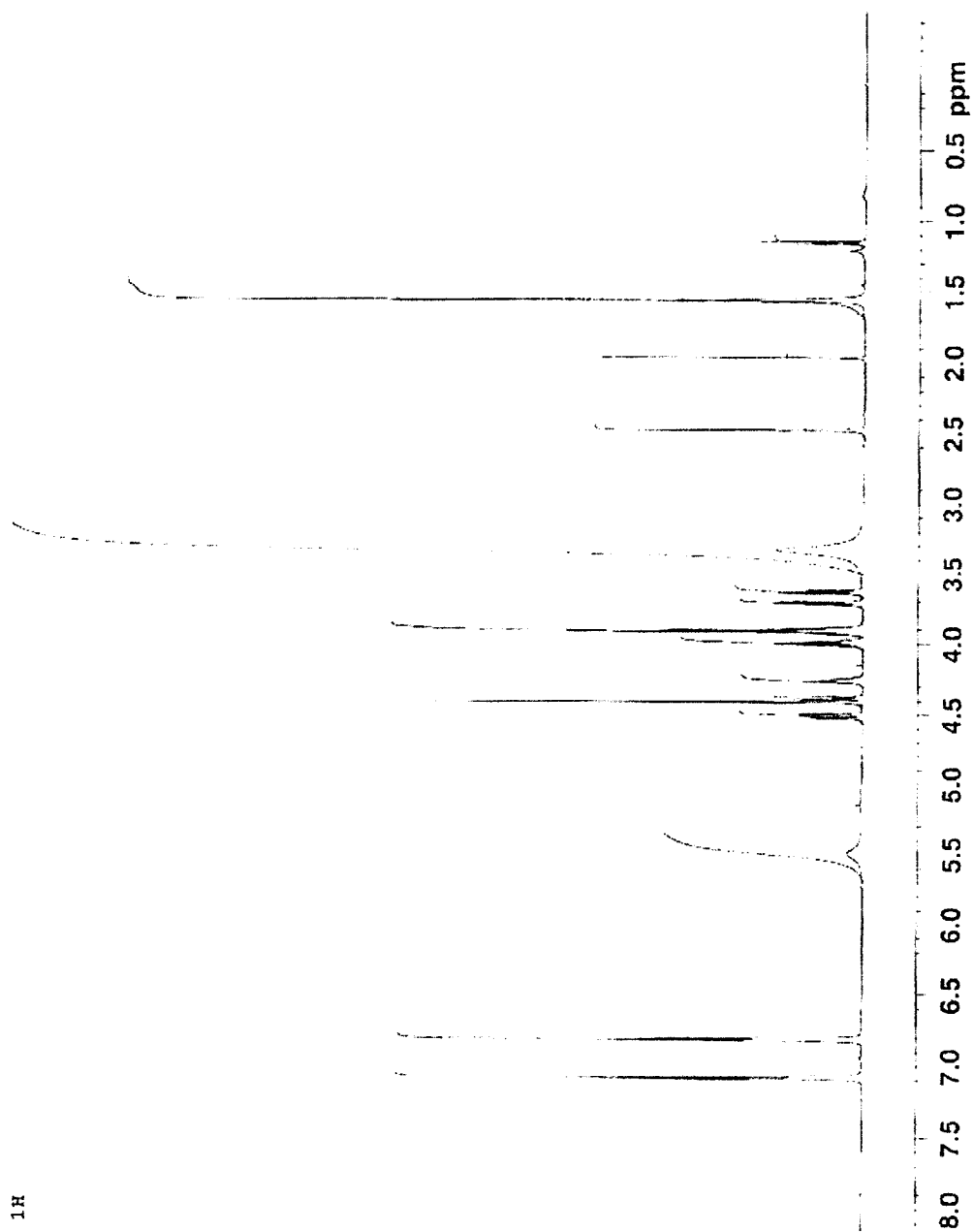
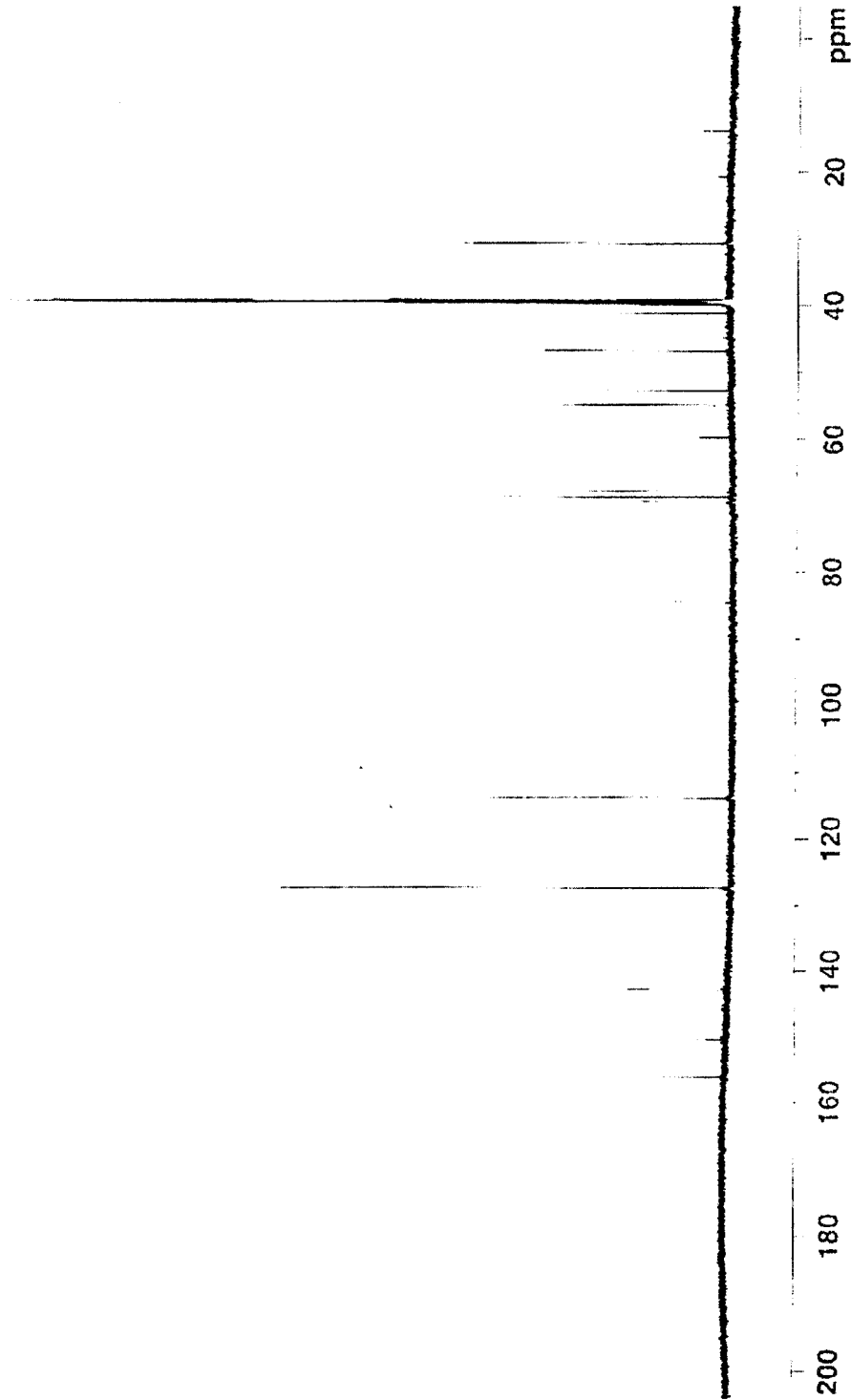


FIG. 6
13C{H}



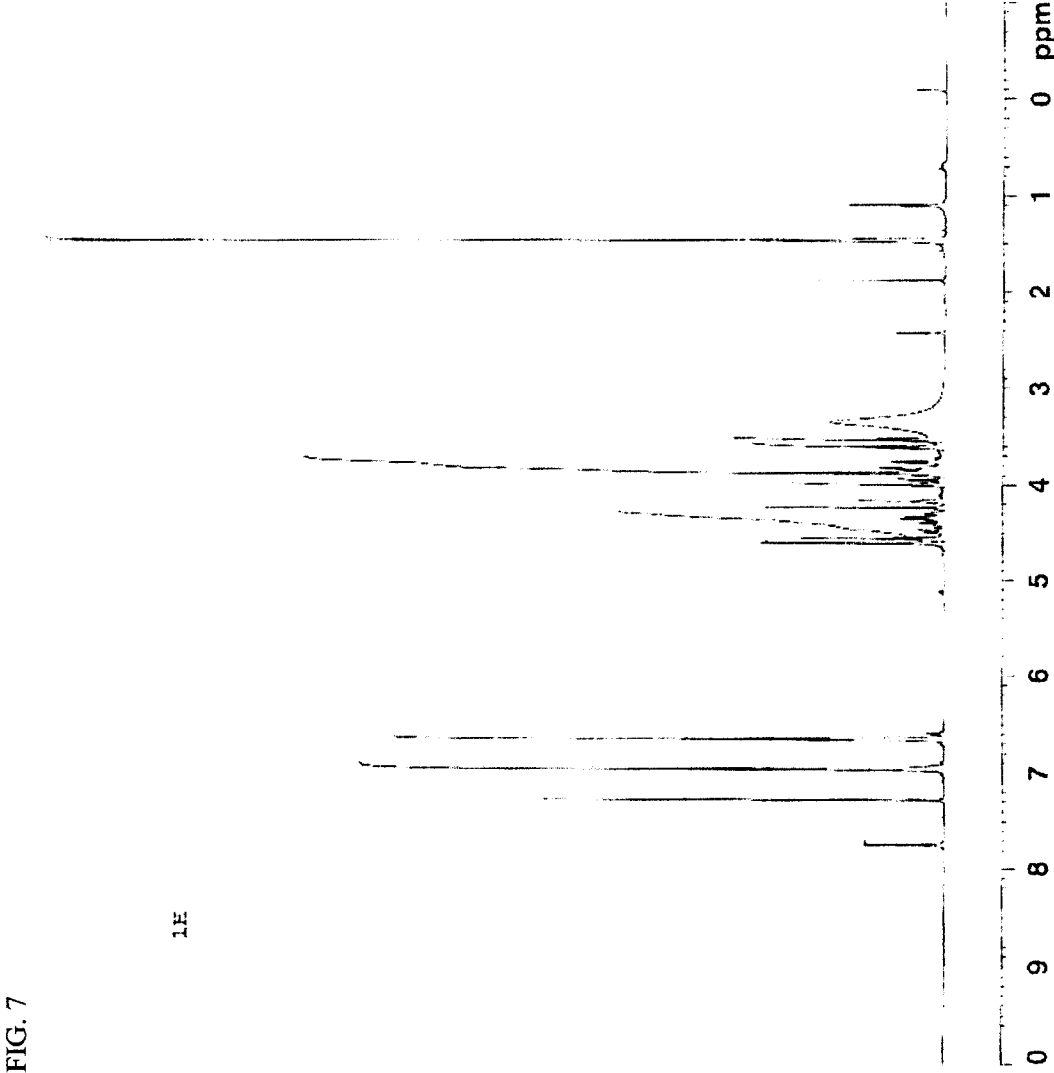
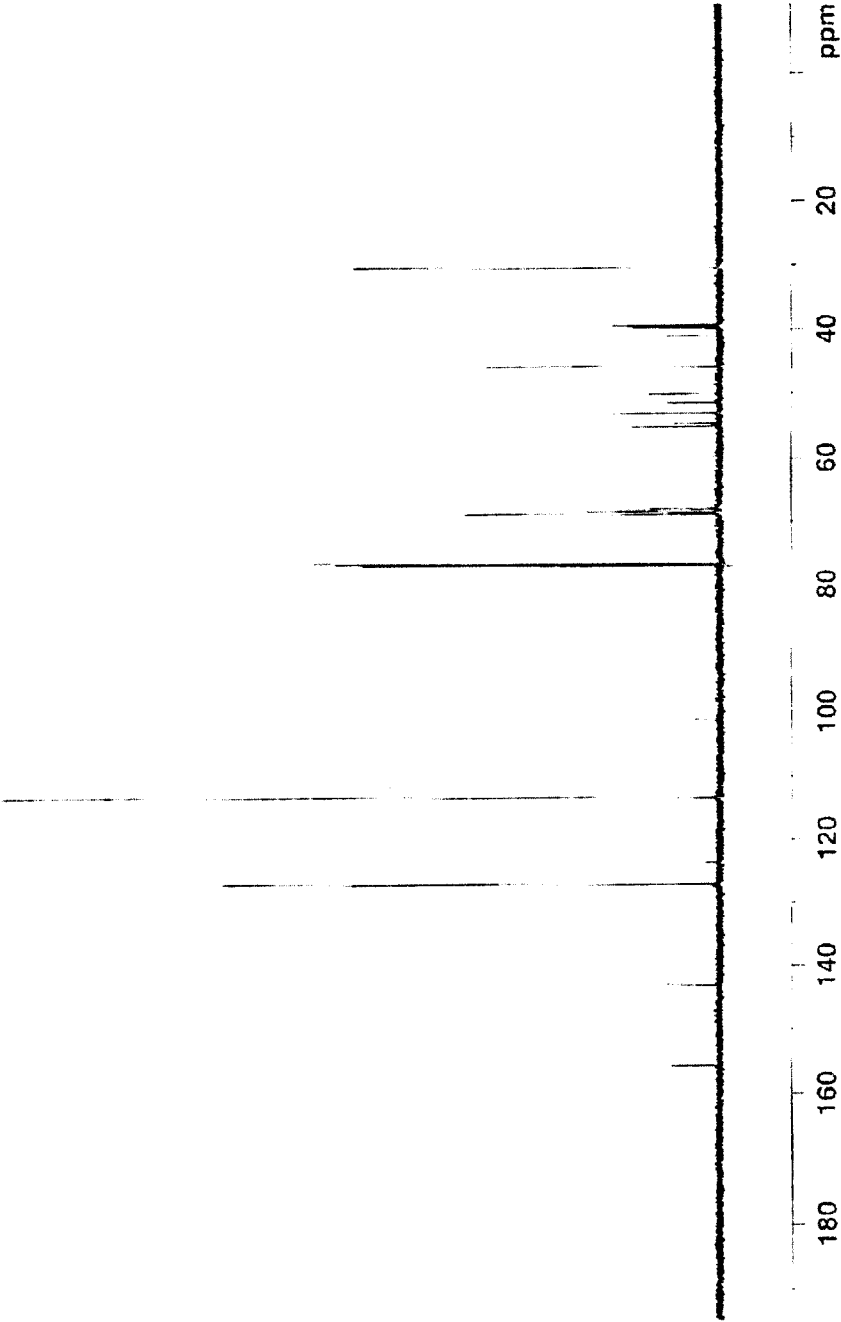


FIG. 8



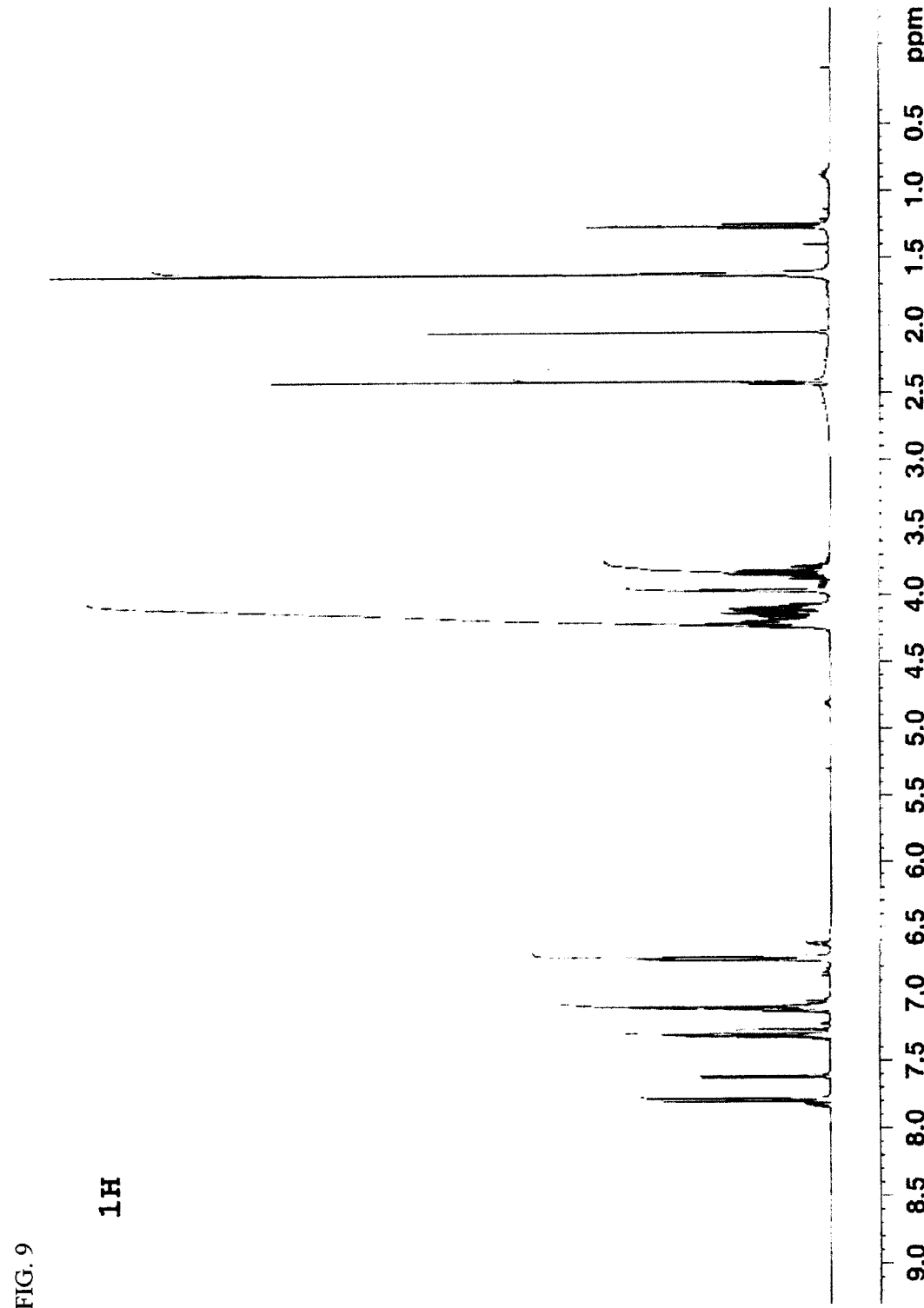


FIG. 10
¹³C

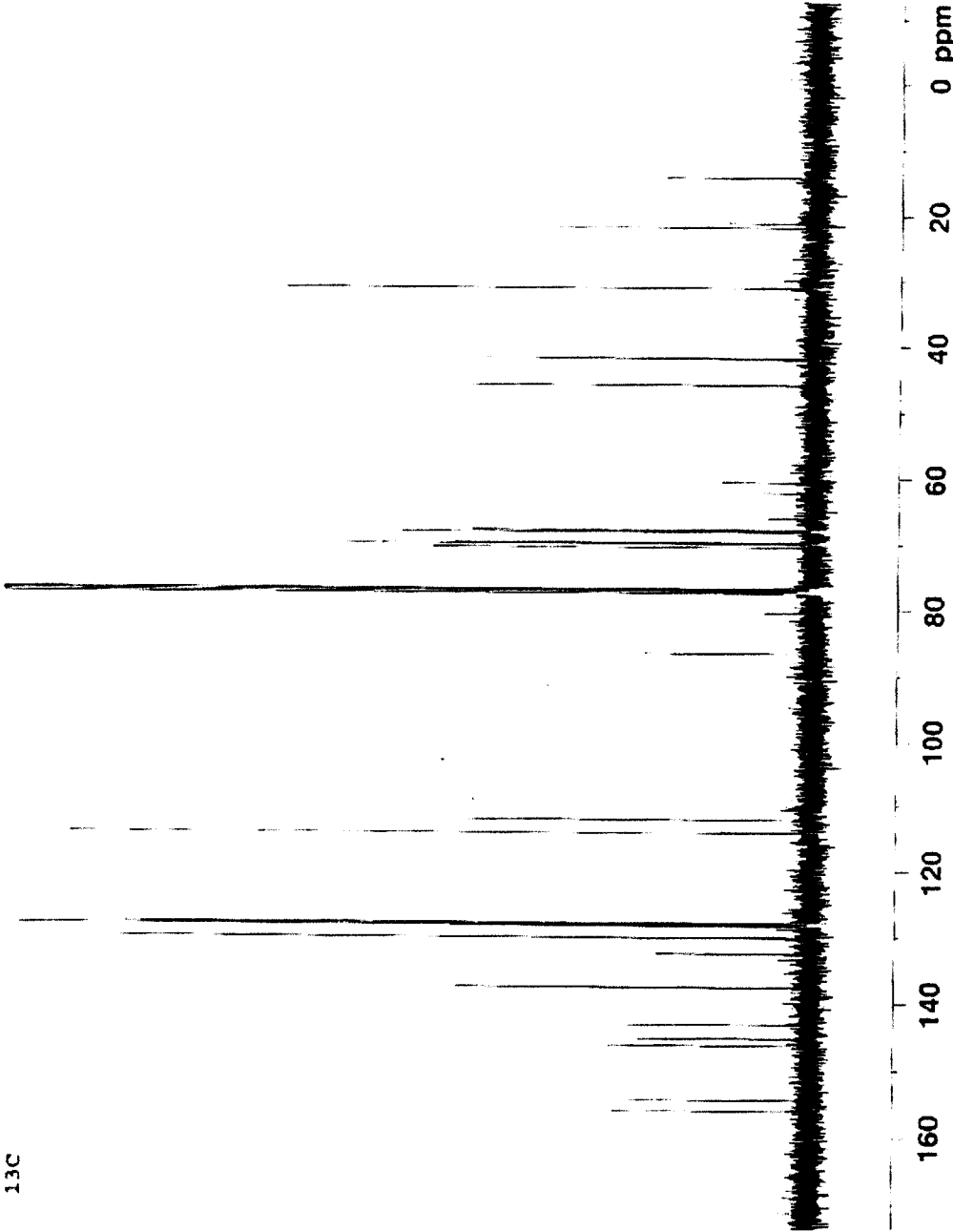


FIG. 11

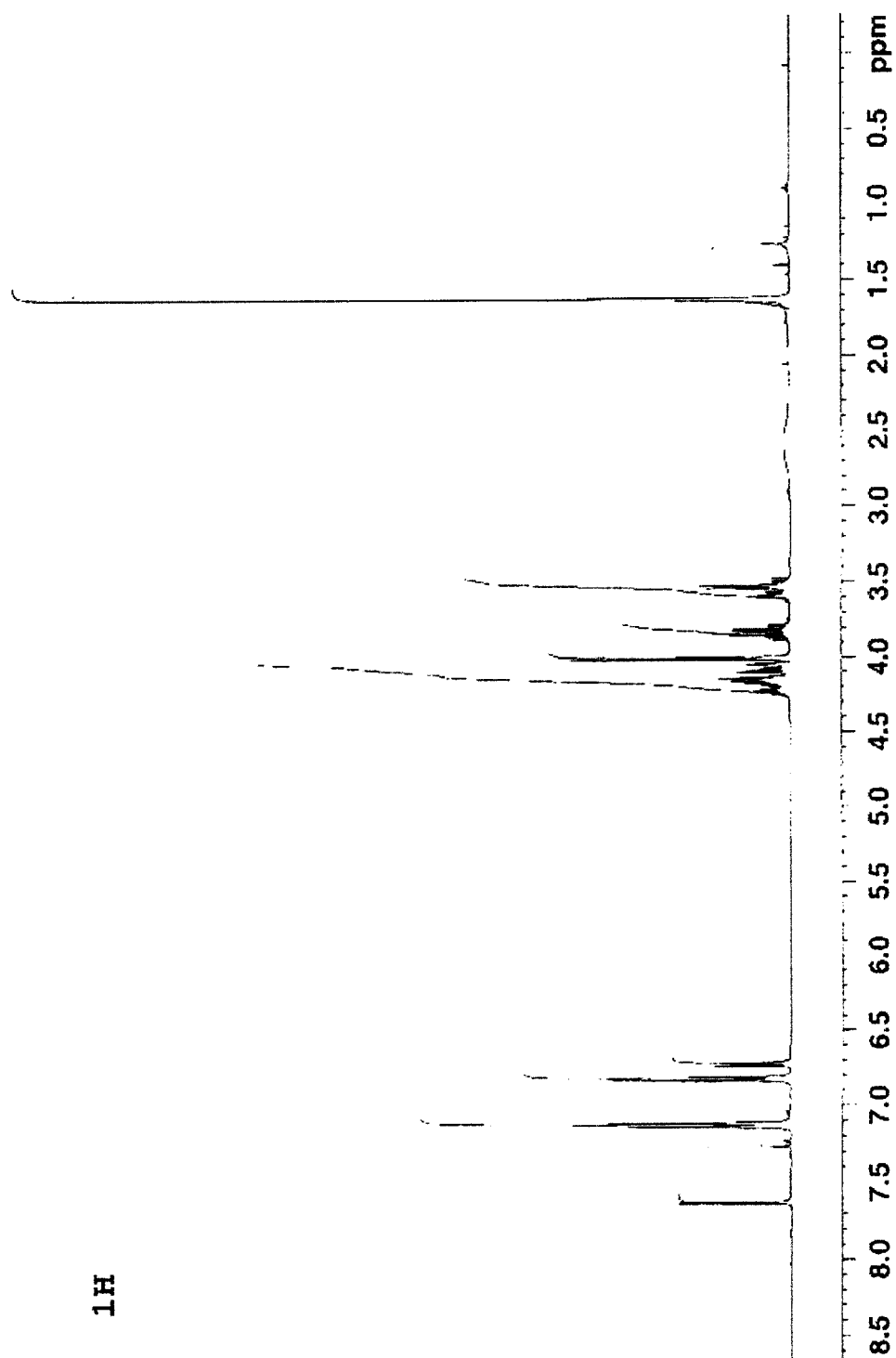
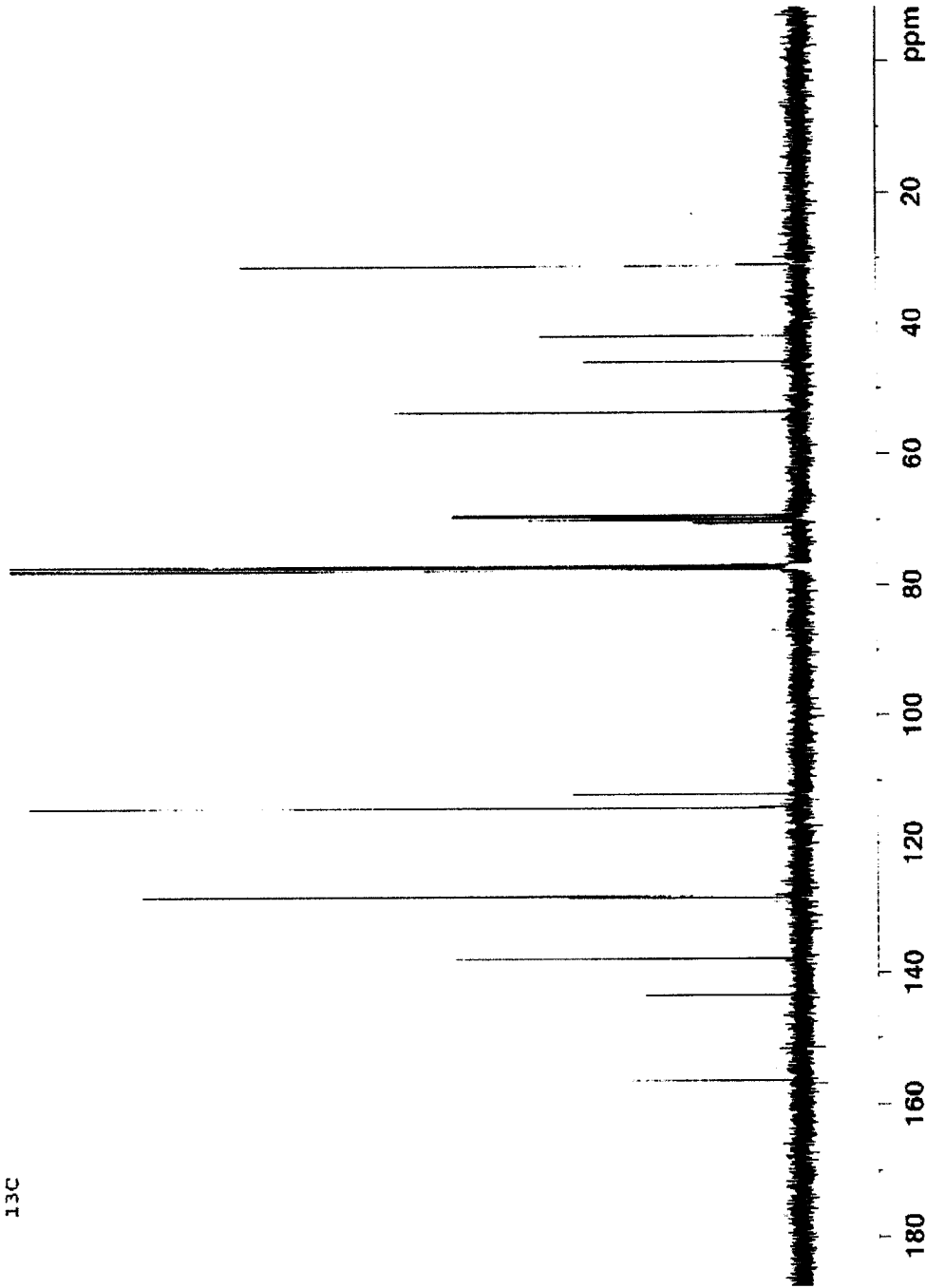
 ^1H 

FIG. 12

¹³C



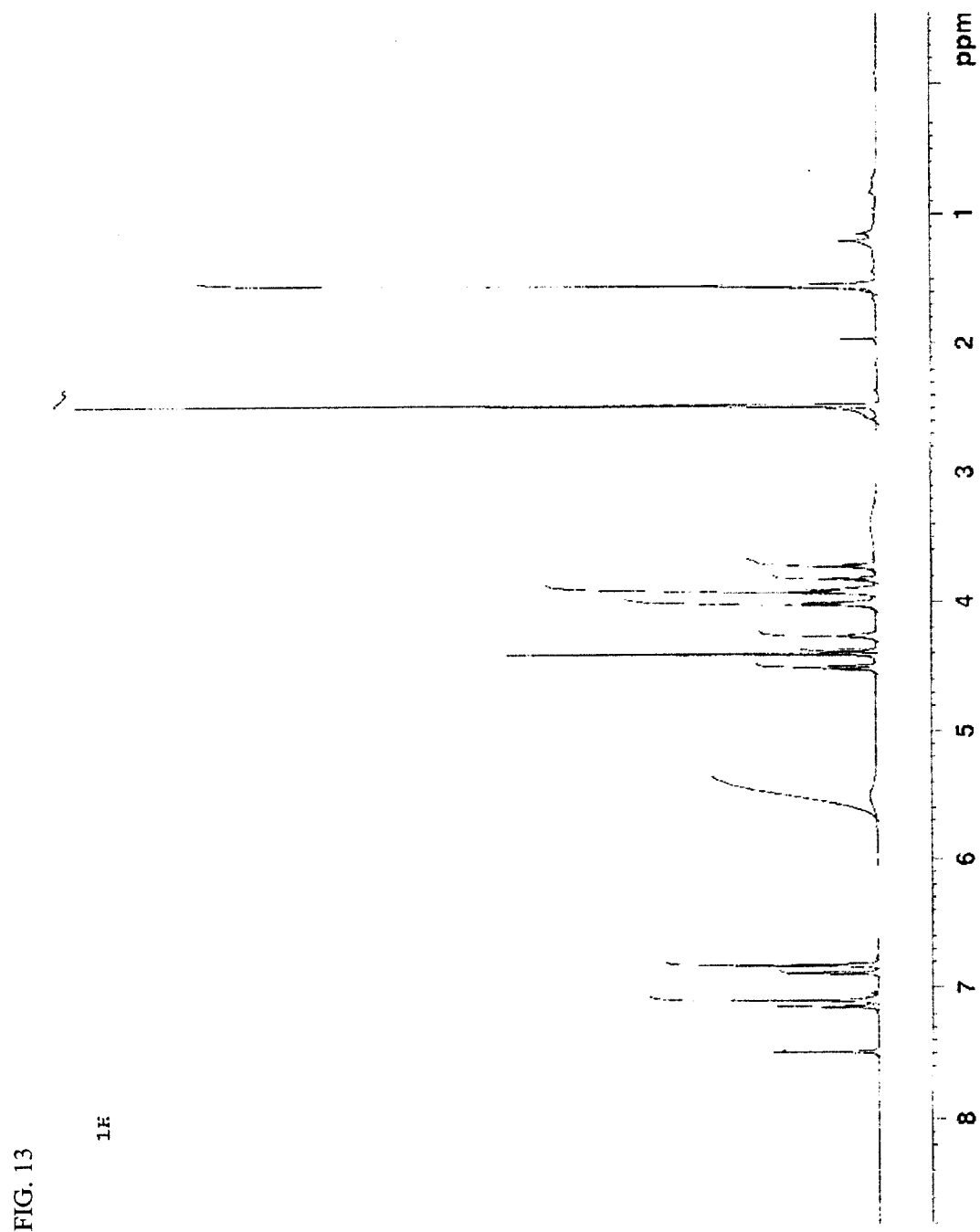


FIG. 14

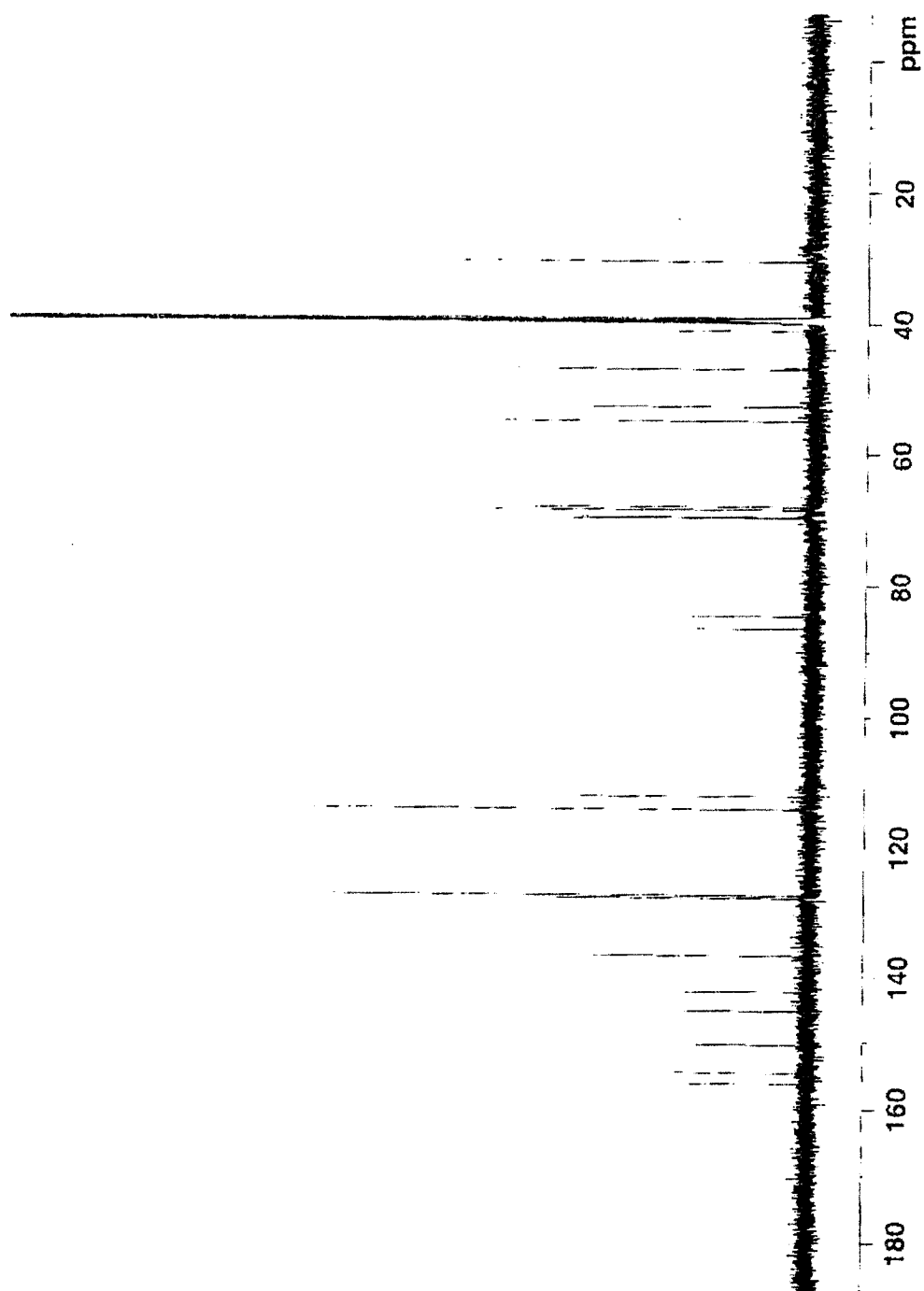


FIG. 15

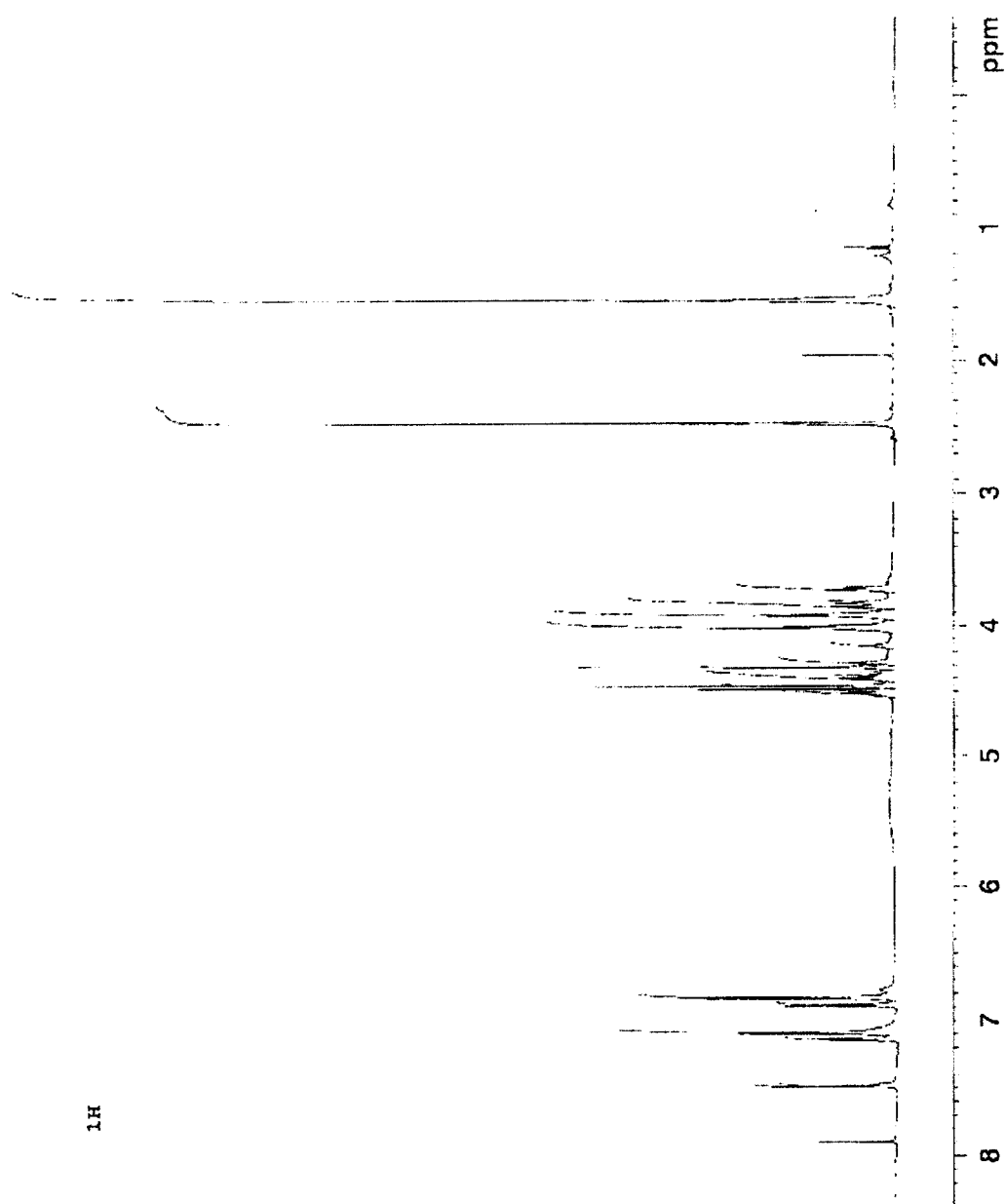
 ^1H 

FIG. 16

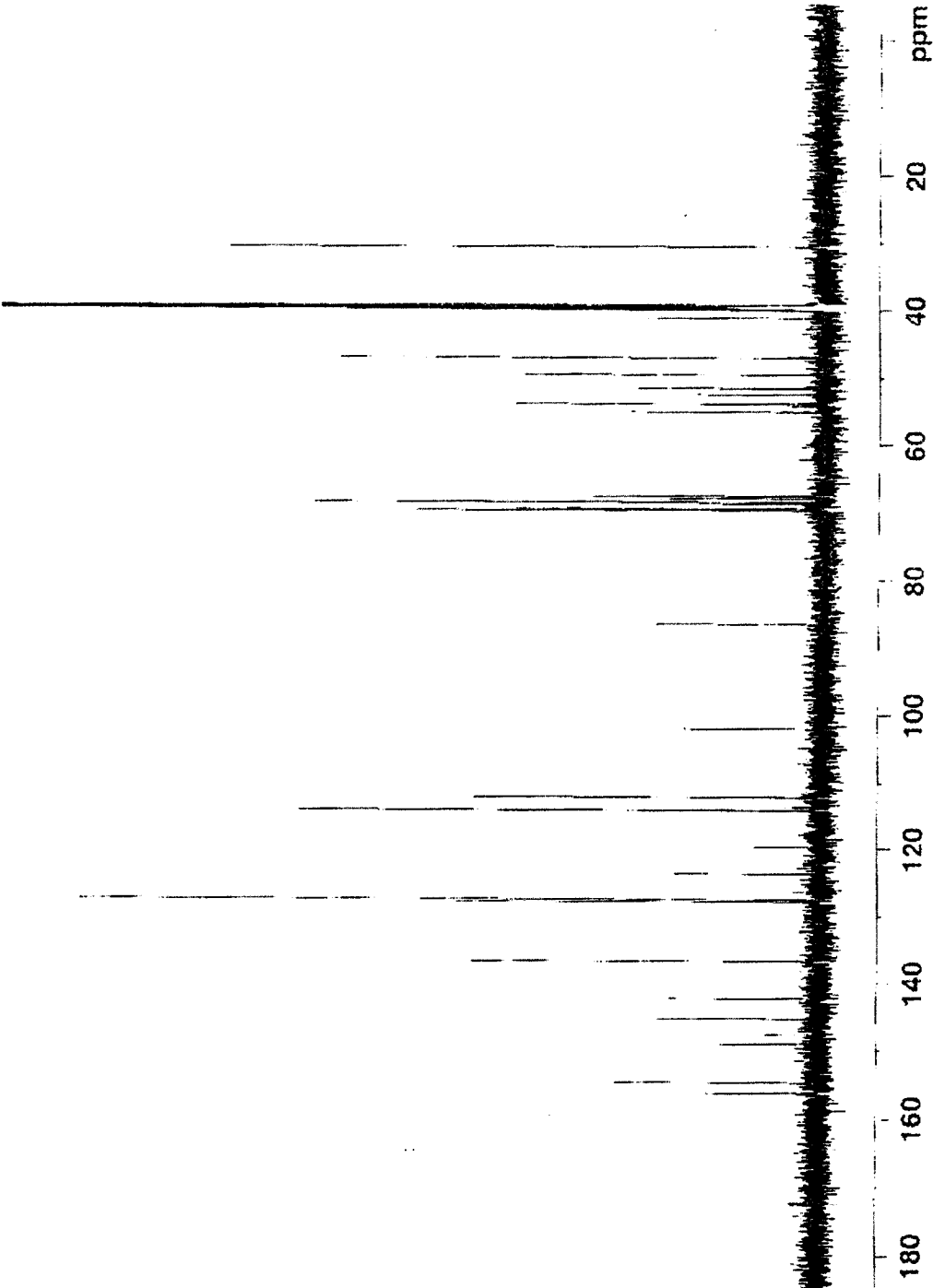
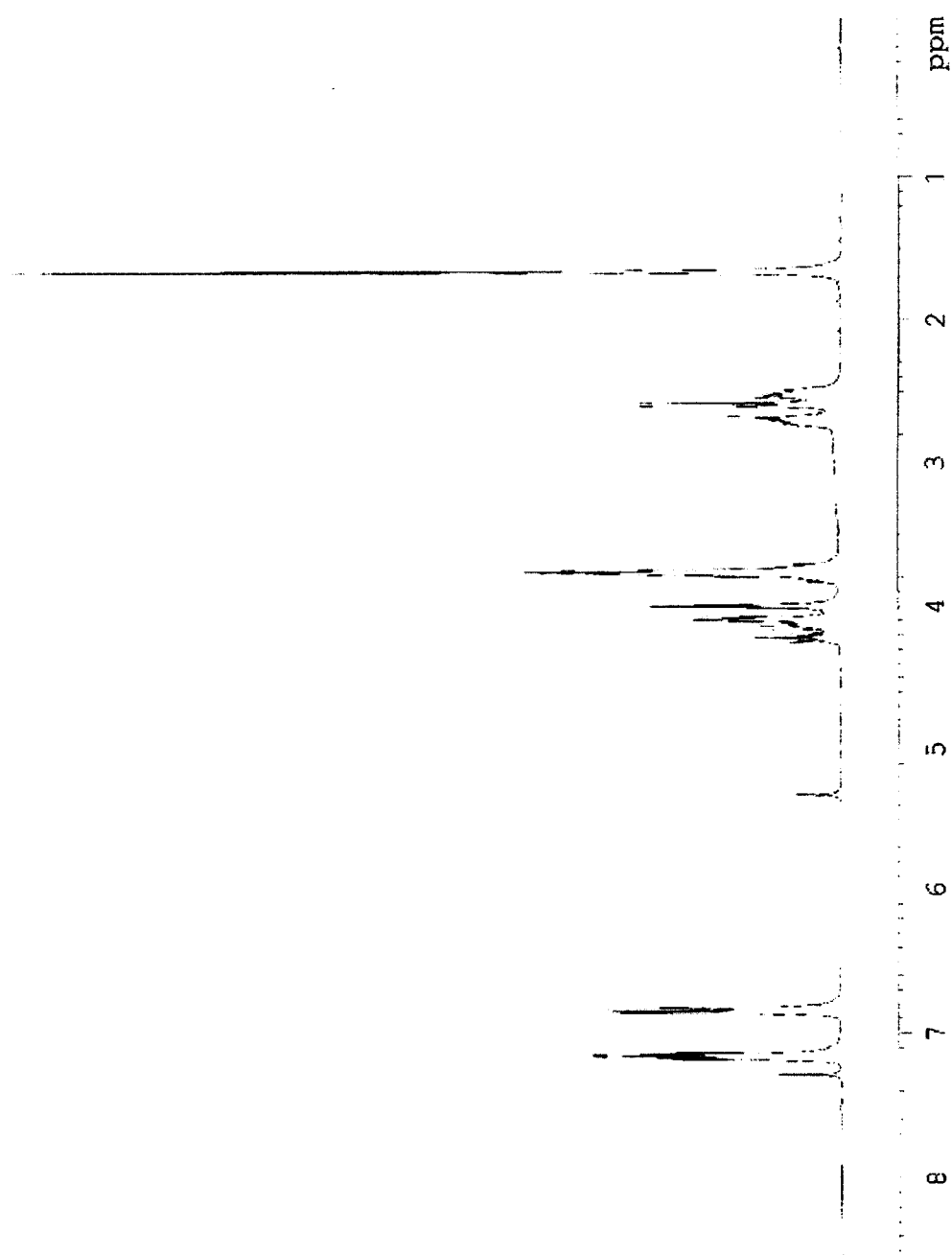


FIG. 17



INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA2016/000008

A. CLASSIFICATION OF SUBJECT MATTER

IPC: **C07D 249/04** (2006.01), **A61K 31/4164** (2006.01), **A61K 31/4192** (2006.01), **A61K 31/495** (2006.01), **A61K 31/5375** (2006.01), **A61K 31/54** (2006.01) (more IPCs on the last page)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC: **C07D 249/04** (2006.01), **A61K 31/4164** (2006.01), **A61K 31/4192** (2006.01), **A61K 31/495** (2006.01), **A61K 31/5375** (2006.01), **A61K 31/54** (2006.01), **A61K 51/04** (2006.01), **A61P 35/00** (2006.01), **C07D 233/64** (2006.01), **C07D 295/088** (2006.01)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic database(s) consulted during the international search (name of database(s) and, where practicable, search terms used)

STN Structure Search

Questel-Orbit (CANCER)/TI/AB/IW/OBJ/ADB/ICLM AND (((SADAR MARIANNE) OR (MAWJI NASRIN) OR (BANUELOS CARMEN) OR (ANDERSEN RAYMOND) OR (FERNANDEZ JAVIER) OR (JIAN KUNZHONG))/IN/OIN/INH/INV)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2012/139039 A2 Andersen, R. J. et al. 11 October 2012 (11-10-2012) *whole document, specifically examples 1, 2, 27, 29*	1-4, 14, 17-21
X	US 2015/010469 A1 Andersen, R. J. et al. 08 January 2015 (08-01-2015) *whole document, specifically compounds 26, 34, 73 and 81*	1-21 and 36-37
X	CA 2226469 A1 Bohlmann, R. et al. 30 January 1997 (30-01-1997) *example 18*	1 and 14
X	CA 2206422 A1 Bitonti, A. J. et al. 06 June 1996 (06-06-1996) *example 2, 10 and 18*	1 and 14

☒ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* "A" "E" "L" "O" "P"	Special categories of cited documents: document defining the general state of the art which is not considered to be of particular relevance earlier application or patent but published on or after the international filing date document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) document referring to an oral disclosure, use, exhibition or other means document published prior to the international filing date but later than the priority date claimed	"T" "X" "Y" "&"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art document member of the same patent family
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Date of the actual completion of the international search
03 March 2016 (03-03-2016)

Date of mailing of the international search report
15 March 2016 (15-03-2016)

Name and mailing address of the ISA/CA
Canadian Intellectual Property Office
Place du Portage I, C114 - 1st Floor, Box PCT
50 Victoria Street
Gatineau, Quebec K1A 0C9
Facsimile No.: 819-953-2476

Authorized officer

Karla Randell (819) 635-5133

INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA2016/000008**Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of the first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claim Nos.: 22-35
because they relate to subject matter not required to be searched by this Authority, namely:
Claims 22-29 are directed to a method for treatment of the human or animal body by surgery or therapy, which the International Searching Authority is not required to search under PCT Rule 39.1(iv). Claims 30-35 are directed to a diagnostic method, which the International Searching Authority is not required to search under PCT Rule 39.1(iv). However, this Authority has carried out a search based on the alleged effect or purpose/use of the product defined in claims 22-35.
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claim Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claim Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA2016/000008

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2014/179867 A1 Mawji, N. R. et al 13 November 2014 (13-11-2014)	1-21 and 36-37
A	WO 2013/028791 A1 Sadar, M. D. et al. 28 February 2013 (28-02-2013)	1-21 and 36-37
A	WO 2011/082487 A1 Sadar, M. D. et al. 14 July 2011 (14-07-2011)	1-21 and 36-37
A	WO 2011/082488 A1 Sadar, M. D. et al. 14 July 2011 (14-07-2011)	1-21 and 36-37
A	WO 2010/000066 A1 Sadar, M. D. et al. 07 January 2010 (07-01-2010)	1-21 and 36-37

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.
PCT/CA2016/000008

Patent Document Cited in Search Report	Publication Date	Patent Family Member(s)	Publication Date
WO2012139039A2	11 October 2012 (11-10-2012)	WO2012139039A2 WO2012139039A3 EP2693875A2 EP2693875A4 US2014248263A1	11 October 2012 (11-10-2012) 25 April 2013 (25-04-2013) 12 February 2014 (12-02-2014) 22 October 2014 (22-10-2014) 04 September 2014 (04-09-2014)
US2015010469A1	08 January 2015 (08-01-2015)	None	
CA2226469A1	30 January 1997 (30-01-1997)	CA2226469A1 AU6519796A DE19526146A1 EP0839130A1 IL122858D0 US5807899A WO9703046A1 ZA9605752A	30 January 1997 (30-01-1997) 10 February 1997 (10-02-1997) 09 January 1997 (09-01-1997) 06 May 1998 (06-05-1998) 16 August 1998 (16-08-1998) 15 September 1998 (15-09-1998) 30 January 1997 (30-01-1997) 19 August 1997 (19-08-1997)
CA2206422A1	06 June 1996 (06-06-1996)	CA2206422A1 AU707290B2 AU712711B2 AU1729799A AU4196796A CN1167437A EP0794771A1 FI972235A HUT77603A IL116142D0 IL129431D0 JPH10509962A MX9703944A NO972426D0 NO972426A US5691384A US5693674A WO9616646A1 ZA9510062A	06 June 1996 (06-06-1996) 08 July 1999 (08-07-1999) 11 November 1999 (11-11-1999) 29 April 1999 (29-04-1999) 19 June 1996 (19-06-1996) 10 December 1997 (10-12-1997) 17 September 1997 (17-09-1997) 27 May 1997 (27-05-1997) 29 June 1998 (29-06-1998) 31 January 1996 (31-01-1996) 17 February 2000 (17-02-2000) 29 September 1998 (29-09-1998) 31 May 1998 (31-05-1998) 28 May 1997 (28-05-1997) 29 July 1997 (29-07-1997) 25 November 1997 (25-11-1997) 02 December 1997 (02-12-1997) 06 June 1996 (06-06-1996) 05 June 1996 (05-06-1996)
WO2014179867A1	13 November 2014 (13-11-2014)	WO2014179867A1 AP201508838D0 AU2014262333A1 CA2911352A1 CN105358522A KR20160013072A SG11201509038UA US2014335080A1 US9173939B2	13 November 2014 (13-11-2014) 30 November 2015 (30-11-2015) 19 November 2015 (19-11-2015) 13 November 2014 (13-11-2014) 24 February 2016 (24-02-2016) 03 February 2016 (03-02-2016) 30 December 2015 (30-12-2015) 13 November 2014 (13-11-2014) 03 November 2015 (03-11-2015)
WO2013028791A1	28 February 2013 (28-02-2013)	None	

continued on extra sheet

INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA2016/000008***Continuation of Patent Family***

Patent Document Cited in Search Report	Publication Date	Patent Family Member(s)	Publication Date
WO2011082487A1	14 July 2011 (14-07-2011)	WO2011082487A1 WO2011082487A8 AR079846A1 CA2786319A1 EP2521707A1 EP2521707A4 JP2013516435A US2013131167A1	14 July 2011 (14-07-2011) 17 January 2013 (17-01-2013) 22 February 2012 (22-02-2012) 14 July 2011 (14-07-2011) 14 November 2012 (14-11-2012) 24 July 2013 (24-07-2013) 13 May 2013 (13-05-2013) 23 May 2013 (23-05-2013)
WO2011082488A1	14 July 2011 (14-07-2011)	WO2011082488A1 AR079975A1 US2013109758A1	14 July 2011 (14-07-2011) 07 March 2012 (07-03-2012) 02 May 2013 (02-05-2013)
WO2010000066A1	07 January 2010 (07-01-2010)	WO2010000066A1 AU2009266379A1 AU2009266379B2 AU2009266379C1 BRPI0913673A2 CA2728219A1 CN102083780A CN105012282A CO6351774A2 EP2307342A1 EP2307342A4 IL210120D0 IL210120A JP2011526250A JP2015163615A KR20110044216A KR20150038226A MX2010014372A NZ589759A RU2011103538A RU2572596C2 RU2015107733A TR201011157T1 US2013245129A1 US8686050B2 US2011230556A1	07 January 2010 (07-01-2010) 07 January 2010 (07-01-2010) 15 January 2015 (15-01-2015) 17 September 2015 (17-09-2015) 13 October 2015 (13-10-2015) 07 January 2010 (07-01-2010) 01 June 2011 (01-06-2011) 04 November 2015 (04-11-2015) 20 December 2011 (20-12-2011) 13 April 2011 (13-04-2011) 20 July 2011 (20-07-2011) 28 February 2011 (28-02-2011) 30 June 2015 (30-06-2015) 06 October 2011 (06-10-2011) 10 September 2015 (10-09-2015) 28 April 2011 (28-04-2011) 08 April 2015 (08-04-2015) 20 June 2011 (20-06-2011) 21 December 2012 (21-12-2012) 10 August 2012 (10-08-2012) 20 January 2016 (20-01-2016) 27 June 2015 (27-06-2015) 21 October 2011 (21-10-2011) 19 September 2013 (19-09-2013) 01 April 2014 (01-04-2014) 22 September 2011 (22-09-2011)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA2016/000008

A61K 51/04 (2006.01) , *A61P 35/00* (2006.01) , *C07D 233/64* (2006.01) , *C07D 295/088* (2006.01)