Title: FLUORENE-9-BISPHENOL COMPOUNDS AND METHODS FOR THEIR USE

Abstract: Compounds having a structure of Formula 1: or a pharmaceutically acceptable salt or tautomer thereof, wherein Z₁, Z₂, Z₃ and Z₄ are as defined herein. Uses of such compounds for treatment of various indications, including prostate cancer as well as methods of treatment involving such compounds are also provided.
CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims the benefit under 35 U.S.C. § 119(e) of U.S. Provisional Patent Application No. 61/476,728 filed 18 April 2011, where this provisional application is incorporated herein by reference in its entirety.

FIELD OF INVENTION

This invention relates to therapeutics, their uses and methods for the treatment of various indications, including various cancers. In particular the invention relates to therapies and methods of treatment for cancers such as prostate cancer, including all stages and androgen dependent, androgen-sensitive and castration-resistant (also referred to as hormone refractory, androgen-independent, androgen deprivation resistant, androgen ablation resistant, androgen depletion-independent, castration-recurrent, anti-androgen-recurrent).

BACKGROUND OF THE INVENTION

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).


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 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.

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, 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 (Culis et al 1994 Cancer Res. 54, 5474-5478; Nazareth et al 1996 J. Biol. Chem. 271, 19900-19907; Sadar 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 2002 A J. Biol. Chem. 277, 7076-7085; and Ueda et al 2002 B J. Biol. Chem. 277, 38087-38094). The AR may 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 48, 189-193).

Available inhibitors of the AR include nonsteroidal antiandrogens such as bicalutamide (Casodex™), nilutamide, flutamide, investigational drugs MDV3100 and ARN-509, and the steroidal antiandrogen, 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 androgen-independent 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. Recent studies developing antagonists to the AR have concentrated on the C-terminus and specifically: 1) the allosteric pocket and AF-2 activity (Estebanez-Perpina et al 2007, PNAS 104,


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.

Recent advances in the development of compounds that modulate AR include the bis-phenol compounds disclosed in published PCT WO 2010/000066 to the British Columbia Cancer Agency Branch and The University of British Columbia. While such compounds appear promising, there remains a need in the art for additional and/or improved compounds that modulate the AR, and which provide treatment for conditions that benefit from such modulation.

**BRIEF SUMMARY**

The compounds described herein may 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 the androgen receptor). Furthermore, these compounds may be 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.

This invention is also based in part on the surprising discovery that the compounds described herein, may also be used to modulate AR activity either in vivo or in vitro for both research and therapeutic uses. The compounds may be used in an effective amount so that androgen receptor activity may be modulated. The AR may be mammalian. Alternatively, the androgen receptor may be human. In particular, the compounds may be used to inhibit the AR. The compounds modulatory activity may be used in either an in vivo or an in vitro model for the study of at least one of the following indications: prostate cancer, breast cancer, ovarian cancer, salivary gland carcinoma, endometrial cancer, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty (testotoxicosis), spinal and bulbar muscular atrophy (SBMA, Kennedy's disease), and age-related macular degeneration. Furthermore, the compounds modulatory activity may be used for the treatment of at least one of the following indications: 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. The indication for treatment may be prostate cancer. The prostate cancer may be castration-resistant prostate cancer. The prostate cancer may be androgen-dependent prostate cancer. In other examples the indication is Kennedy's disease.

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

\[ \text{I} \]

or a pharmaceutically acceptable salt or tautomer thereof, wherein \( Z^1, Z^2, Z^3 \) and \( Z^4 \) are as defined herein.

In other embodiments, the present disclosure provides the use of a compound of Formula I, for modulating androgen receptor (AR) activity. Methods for
modulating AR, as well as pharmaceutical compositions comprising a compound of Formula I and a pharmaceutically acceptable excipient are also provided.

In addition, the present disclosure provides combination therapy treatments for any of the disease states disclosed herein, for example prostate cancer or Kennedy's disease. The disclosed therapies include use of a pharmaceutical composition comprising a compound of Formula I, an additional therapeutic agent and a pharmaceutically acceptable excipient. Methods and compositions related to the same are also provided.

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.

**DETAILED DESCRIPTION**

As used herein, the phrase "C_x-C_y alkyl" is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that has a carbon skeleton or main carbon chain comprising a number from x to y (with all individual integers within the range included, including integers x and y) of carbon atoms. For example a "Ci-Cio alkyl" is a chemical entity that has 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 carbon atom(s) in its carbon skeleton or main chain. Non-limiting examples of saturated Ci-Ci_0 alkyl include methyl, ethyl, n-propyl, i-propyl, sec-propyl, n-butyl, i-butyl, sec-butyl, t-butyl and n-penty, n-hexyl, n-heptane, and the like. Non-limiting examples of C_2-Cio alkenyl include vinyl, allyl, isopropenyl, 1-propene-2-yl, 1-butene-1-yl, 1-butene-2-yl, 1-butene-3-yl, 2-butene-1-yl, 2-butene-2-yl, pentenyl, hexenyl, and the like. Non-limiting examples of C_2-Cio alkynyl include ethynyl, propynyl, butynyl, pentynyl, hexynyl, and the like. Unless stated otherwise specifically in the specification, an alkyl group may be optionally substituted (i.e., a hydrogen atom in the alkyl group may be replaced with an optional substituent).

As used herein, the term "cyclic C_x-C_y alkyl" is used as it is normally understood to a person of skill in the art and often refers to a compound or a chemical entity in which at least a portion of the carbon skeleton or main chain of the chemical entity is bonded in such a way so as to form a 'loop', circle or ring of atoms that are bonded together. The atoms do not have to all be directly bonded to each other, but
rather may be directly bonded to as few as two other atoms in the 'loop'. Non-limiting examples of cyclic alkyls include benzene, toluene, cyclopentane, bisphenol and 1-chloro-3-ethylcyclohexane.

As used herein, the term "branched" is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that comprises a skeleton or main chain that splits off into more than one contiguous chain. The portions of the skeleton or main chain that split off in more than one direction may be linear, cyclic or any combination thereof. Non-limiting examples of a branched alkyl are tert-butyl and isopropyl.

As used herein, the term "unbranched" is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that comprises a skeleton or main chain that does not split off into more that one contiguous chain. Non-limiting examples of unbranched alkyls are methyl, ethyl, n-propyl, and n-butyl.

As used herein, the term "substituted" is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that has one chemical group replaced with a different chemical group that contains one or more heteroatoms. Unless otherwise specified, a substituted alkyl is an alkyl in which one or more hydrogen atom(s) is/are replaced with one or more atom(s) that is/are not hydrogen(s). For example, chloromethyl is a non-limiting example of a substituted alkyl, more particularly an example of a substituted methyl. Aminoethyl is another non-limiting example of a substituted alkyl, more particularly it is a substituted ethyl.

As used herein, the term "unsubstituted" is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that is a hydrocarbon and/or does not contain a heteroatom. Non-limiting examples of unsubstituted alkyls include methyl, ethyl, tert-butyl, and pentyl.

As used herein, the term "saturated" when referring to a chemical entity is used as it is normally understood to a person of skill in the art and often refers to a chemical entity that comprises only single bonds. Non-limiting examples of saturated chemical entities include ethane, tert-butyl, and N-H.

As used herein, Ci-Cio alkyl may include, for example, and without limitation, saturated Ci-Cio alkyl, C2-C10 alkenyl and C2-C10 alkynyl. Non-limiting examples of saturated C1-C10 alkyl may include methyl, ethyl, n-propyl, i-propyl, sec-
propyl, n-butyl, i-butyl, sec-butyl, t-butyl, n-pentyl, i-pentyl, sec-pentyl, t-pentyl, n-hexyl, i-hexyl, 1,2-dimethylpropyl, 2-ethylpropyl, 1-methyl-2-ethylpropyl, 1-ethyl-2-methylpropyl, 1,1,2-trimethylpropyl, 1,1,2-triethylpropyl, 1,1-dimethylbutyl, 2,2-dimethylbutyl, 2-ethylbutyl, 1,3-dimethylbutyl, 2-methylpentyl, 3-methylpentyl, sec-hexyl, t-hexyl, n-heptyl, i-heptyl, sec-heptyl, t-heptyl, n-octyl, i-octyl, sec-octyl, t-octyl, n-nonyl, i-nonyl, sec-nonyl, t-nonyl, n-decyl, i-decyl, sec-decyl and t-decyl. Non-limiting examples of C_2-C_10 alkynyl may include vinyl, allyl, isopropenyl, 1-propene-2-yl, 1-butene-1-yl, 1-butene-2-yl, 2-butene-3-yl, 2-butene-2-yl, octenyl and decenyl. Non-limiting examples of C_2-C_10 alkynyl may include ethynyl, propynyl, butynyl, pentynyl, hexynyl, heptynyl, octynyl, nonynyl and decynyl. Saturated C_1-C_10 alkyl, C_2-C_10 alkenyl or C_2-C_10 alkynyl may be, for example, and without limitation, interrupted by one or more heteroatoms which are independently nitrogen, sulfur or oxygen.

As used herein, cyclic C_3-C_10 alkyl may include, for example, and without limitation, saturated C_3-C_10 cycloalkyl, C_3-C_10 cycloalkenyl, C_3-C_10 cycloalkynyl, C_{6,10} aryl, C_{6,9} aryl-C_{1-4} alky, C_{6,8} aryl-C_{2-4} alkenyl, C_{6,8} aryl-C_{2-4} alkynyl, a 4- to 10-membered non-aromatic heterocyclic group containing one or more heteroatoms which are independently nitrogen, sulfur or oxygen, and a 5- to 10-membered aromatic heterocyclic group containing one or more heteroatoms which are independently nitrogen, sulfur or oxygen. Non-limiting examples of the saturated C_3-C_10 cycloalkyl group may include cyclopropanyl, cyclobutanyl, cyclopentany1, cyclohexanyl, cycloheptanyl, cyclooctanyl, cyclononanyl and cyclodecananyl. Non-limiting examples of the C_3-C_10 cycloalkenyl group may include cyclopropanenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, cyclpentenyl, cycloctenyl, cyclononanenyl and cyclodecanenyl. Non-limiting examples of the C_{6,10} aryl group may include phenyl (Ph), pentalenyl, indenyl, naphthyl, and azulenyl. The C_{6,9} aryl-C_{1-4} alkyl group may be, for example, and without limitation, a C_{1-4} alkyl group as defined anywhere above having a C_{6,9} aryl group as defined anywhere above as a substituent. The C_{6,8} aryl-C_{2-4} alkenyl group may be, for example, and without limitation, a C_{2-4} alkenyl as defined anywhere above having a C_{6,8} aryl group as defined anywhere above as a substituent. The C_{6,8} aryl-C_{2-4} alkynyl group may be, for example, and without limitation, a C_{2-4} alkynyl group as defined anywhere above having a C_{6,8} aryl group as defined anywhere above as a substituent. Non-limiting examples of the 4- to 10-
membered non-aromatic heterocyclic group containing one or more heteroatoms which are independently nitrogen, sulfur or oxygen may include pyrrolidinyl, pyrrolinyl, piperidinyl, piperazinyl, imidazolinyl, pyrazolidinyl, morpholinyl, tetrahydropyranyl, azetidinyl, oxetanyl, oxathiolanyl, phthalimide and succinimide. Non-limiting examples of the 5- to 10-membered aromatic heterocyclic group containing one or more heteroatoms which are independently nitrogen, sulfur or oxygen may include pyrrolyl, pyridinyl, pyridazinyl, pyrimidinyl, pirazinyl, imidazolyl, thiazolyl and oxazolyl.

Non-limiting examples of one to ten carbon substituted or unsubstituted acyl include acetyl, propionyl, butanoyl and pentanoyl. Non-limiting examples of Ci-Cio alkoxy include methoxy, ethoxy, propoxy and butoxy.

As used herein, the symbol "\( \text{\_\_} \)" (hereinafter may 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 example, "\( \text{\_\_} \)" 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 may be specified by inference. For example, the compound CH\(_3\)-R\(^3\), wherein R\(^3\) is H or "\( \text{\_\_\_\_\_} \)" 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\(_3\).

"Halo" refers to fluoro (F), chloro (Cl), bromo (Br) or iodo (I). Radioisotopes are included within the definition of halo. Accordingly, compounds comprising fluoro, chloro, bromo or iodo may also comprise radioisotopes of the same.

As noted above, the present disclosure provides a compound having a structure of Formula I:
or a pharmaceutically acceptable salt or tautomer thereof, wherein:

at least one \( Z_1 \) is independently \( C-Q \);

at least one \( Z_2 \) is independently \( C-T \), \( CCH_3 \), \( CF \), \( CC1 \), \( CBr \), \( Cl \), \( COH \), \( CG_1 \), \( COG_1 \), \( CNH_2 \), \( CNHG_1 \), \( CN(G^1)_2 \), \( COS0_3H \), \( COP0_3H_2 \), \( CSG_1 \), \( CSOG_1 \), or \( CSOzG_1 \);

\( Z_3 \), \( Z_4 \) and each remaining \( Z_1 \) and \( Z_2 \) are, at each occurrence, independently \( C-T \), \( N \), \( CH \), \( CCH_3 \), \( CF \), \( CC1 \), \( CBr \), \( Cl \), \( COH \), \( CG_1 \), \( COG_1 \), \( CNH_2 \), \( CNHG_1 \), \( CNG_1 \), \( COS0_3H \), \( COPO2H_2 \), \( CSG_1 \), \( CSOG_1 \), or \( CSOzG_1 \);

\( Q \) is \( G^1 \), \( O \), \( CH_2 \), \( CHG_1 \), \( C(G^1)_2 \), \( S \), \( NH \), \( NG^1 \), \( SO \), \( S0_2 \), or \( NR \);

\( J \) is \( G^1 \), \( O \), \( CH_2 \), \( CHG_1 \), \( C(G^1)_2 \), \( S \), \( NH \), \( NG^1 \), \( SO \), \( S0_2 \), or \( NR \);

\( M \) is \( H \), \( F \), \( Cl \), \( Br \), \( CH_2OH \), \( CH_2OD \), \( CH_2OG^1 \), \( CH_2F \), \( CH_2Cl \), \( CHC1_2 \), \( CC1_3 \), \( CH_2Br \), \( CHBr_2 \), \( CBr_3 \) or \( C=CH \);

\( L \) is \( H \) or \( A-D \);

\( A \) is \( O \), \( S \), \( NH \), \( NG^1 \), \( N^+H_2 \) or \( N^+HG^1 \);

\( D \) is, at each occurrence, independently \( H \), \( G^1 \), \( R \),

or a moiety from TABLE 1;

each of \( q \), \( r \) and \( t \) may independently be 0, 1, 2, 3, 4, 5, 6 or 7;

\( n \) is 0, 1, 2, 3, 4, 5, 6, 7 or 8;

\( T \) is, at each occurrence, independently
\( J^2 \) is, at each occurrence, independently \( \text{G}^1, \text{O}, \text{CH}_2, \text{CHG}^1, \text{C}(\text{G}^1)_2, \text{S}, \text{NH}, \text{NG}^1, \text{SO}, \text{S0}_2 \), or \( \text{NR} \);

\( \text{M}^2 \) is, at each occurrence, independently \( \text{H}, \text{CH}_3, \text{F}, \text{Cl}, \text{Br}, \text{CH}_2\text{F}, \text{CH}_2\text{Cl}, \text{CHCl}_2, \text{CCl}_3, \text{CH}_2\text{Br}, \text{CHBr}_2, \text{CBr}_3, \text{CH}_2\text{OH}, \text{CH}_2\text{OD}^2, \text{CH}_2\text{OJ}^\prime, \text{G}^1, \text{CH}_2\text{OG}^1, \text{CH}_2\text{OR}, \text{CH}_2\text{OJ}^\prime \text{OG}^1, \text{G}^\prime \text{G}^1, \text{G}^\prime \text{G}^\prime \text{OG} \text{J}^\prime \prime, \text{CH}_2\text{SG}^1, \text{CH}_2\text{NH}_2, \text{CH}_2\text{NHG}^1, \text{CH}_2\text{NG}^1, \text{or} \text{C}=\text{CH} ;

\( \text{L}^2 \) is, at each occurrence, independently \( \text{H} \) or \( \text{A}^2-\text{D}^2 \);

\( \text{A}^2 \) is, at each occurrence, independently \( \text{O}, \text{S}, \text{SO}, \text{S0}_2, \text{NH}, \text{NG}^1, \text{N}^+\text{H}_2, \) or \( \text{N}^+\text{HG}^1 \);

\( \text{D}^2 \) is, at each occurrence, independently \( \text{H, G}^1, \text{R}, \text{OH}, \text{OG}^1, \text{OR} \), or a moiety selected from \( \text{TABLE 1} \);

each of \( \text{u}, \text{y} \) and \( \text{j} \) are, at each occurrence, independently \( 0, 1, 2, 3, 4, 5, 6 \) or \( 7 \);

\( \text{m} \) is, at each occurrence, independently \( 0, 1, 2, 3, 4, 5, 6, 7 \) or \( 8 \);

\( \text{J}^\prime \) and \( \text{J}^\prime \prime \) are, at each occurrence, independently a moiety selected from \( \text{TABLE 1} \);

\( \text{G}^1, \text{G}^1 \) and \( \text{G}^1 \) are, at each occurrence, independently a linear or branched, aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated \( \text{C}_1-\text{C}_{10} \) alkyl, wherein the optional substituents for the \( \text{C}_1-\text{C}_{10} \) alkyl are \( \text{oxo, CH}_2\text{CO}_3 \text{R}^1, \text{OJ}^\prime \text{J}^\prime, \text{COOH}, \text{R}^1, \text{OH}, \text{OR}^1, \text{F, Cl, Br, I, NH}_2, \text{NHR}^1, \text{N(R}^1)_2, \text{CN, SH, SR}^1, \text{SO}_3 \text{H, SO}_2 \text{R}^1, \text{OSO}_2 \text{R}^1, \text{OR}^2, \text{CO}_2 \text{R}^1, \text{CONH}_2, \text{CONHR}^1, \text{CONHR}^2, \text{CON(R}^1)_2, \text{NHR}^2, \text{OPo}_3 \text{H}_3, \text{CON^AR}^2, \text{NR}^1 \text{R}^2 \) or \( \text{N0}_2 \);

each \( \text{R}^1 \) is independently \( \text{H, linear or branched, aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C}_1-\text{C}_{10} \) alkyl or a metal counter ion, wherein the metal counter ion is \( \text{Li, Na, K, Mg or Ca} \);

each \( \text{R}^1 \) is independently unsubstituted \( \text{C}_1-\text{C}_{10} \) alkyl; and

each \( \text{R} \) and \( \text{R}^2 \) are independently \( \text{C}_1-\text{C}_{10} \) acyl. For example, in some embodiments at least one \( Z^2 \) is \( \text{C}-\text{T} \).

In other embodiments, the compound has a structure of Formula II:
In yet other embodiments, the compound has a structure of Formula III:

wherein:

\[ R_3, R_4, R_5, R_6, R_7, R_8, R_9 \text{ and } R_{10} \text{ are each independently hydrogen, halo, or linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cio alkyl.} \]

In still other exemplary embodiments, the compound has a structure of Formula IV:

In yet other embodiments, the compound has a structure of Formula V:
For example, in some further embodiments, the compound has a structure of Formula Va, Vb, Vc or Vd:

In other further embodiments, the compound has one of the following structures:
wherein:

R<sub>i</sub> is hydrogen or linear or branched, substituted or unsubstituted, saturated or unsaturated C<sub>i</sub>-C<sub>i</sub>0 alkyl;

Y is Cl or OH;

q is 0, 1, 2, 3, 4, 5, 6 or 7; and

m is 0, 1, 2, 3, 4, 5, 6, 7 or 8.

In other embodiments of the foregoing compound, J is G<sup>1</sup>, O, CH<sub>2</sub>, CHG<sup>1</sup>, CG<sup>+</sup>, NH, SO, or NR. For example in some embodiments, J is O.

In other embodiments, M is Cl, Br, CH<sub>2</sub>OH, CH<sub>2</sub>OD, CH<sub>2</sub>OG<sup>1</sup>, CH<sub>2</sub>F, CH<sub>2</sub>Cl, CHCl<sub>2</sub>, CC<sub>1</sub>3, CH<sub>2</sub>Br, CHBr<sub>2</sub>, CBr<sub>3</sub>, or C≡CH. In still other embodiments, M is CH<sub>2</sub>OH, CH<sub>2</sub>F, C≡CH, CH<sub>2</sub>OCH<sub>2</sub>C≡CH, CH<sub>2</sub>OCH<sub>3</sub>, CH<sub>2</sub>0-/-propyl, CH<sub>2</sub>0-n-butyl, or
CH$_2$OD, wherein D is CH$_2$OH or CHG$^1$. For example in some embodiments, M is CH$_2$OH, and in other embodiments, M is H.

In other embodiments, L is H, and in other embodiments, L is A-D.

In yet other embodiments, A is O while in other embodiments D is H, R, or a moiety from TABLE 1; and each of q, r and t is independently 0, 1, 2, 3, 4, 5, 6 or 7. For example in some embodiments, D is H, and in other embodiments D is R.

In certain embodiments, D is a moiety selected from TABLE 1. For example in some embodiments, the moiety from TABLE 1 is or 

In other embodiments, n is 0, and in some other embodiments n is 1, 2, 3, 4, or 5. For example in some embodiments, n is 1.

In certain embodiments, J$^2$ is G$^1$, O, CH$_2$, CHG$^1$, CG$^A$, NH, SO, or NR. For example in some embodiments, J$^2$ is O.

In other embodiments, M$^2$ is H, CH$_2$F, CH$_2$Cl, CH$_2$Br, CH$_2$OH, CH$_2$OJ", CH$_2$OG$^1$, or C≡CH. For example in some embodiments, M$^2$ is CH$_2$F. In other embodiments, M$^2$ is CH$_2$Cl. In still other embodiments, M$^2$ is CH$_2$Br. In yet other embodiments, M$^2$ is CH$_2$OH, and in some embodiments M$^2$ is H. In other embodiments, M$^2$ is C≡CH.

In still other embodiments, L$^2$ is H, and in other examples L$^2$ is A$^2$-D$^2$.

In certain embodiments, A$^2$ is O, and in other certain embodiments D$^2$ is H, R, or a moiety from TABLE 1; and each of u, v and j is independently 0, 1, 2, 3, 4, 5, 6 or 7. For example, in some embodiments D$^2$ is H, and in other embodiments D$^2$ is R. In still
other embodiments, D² is a moiety from TABLE 1. For example in some
embodiments, the moiety from TABLE 1 is

\[
\begin{align*}
\text{O} & \quad \text{NH}_2 \\
\text{O} & \quad \text{NH}_3^+
\end{align*}
\]

In other embodiments of the foregoing compound, m is 0. In still other
embodiments, m is 1, 2, 3, 4, or 5. For example in some embodiments, m is 1.

In other examples, M is CH₂OH and L is OH. In still other examples,
M² is CH₂Cl and L is OH. In even further examples, M is CH₂OH, M² is CH₂Cl, L is
OH and L² is OH. In even further examples, M is CH₂F, M² is CH₂Cl, L is OH and L²
is OH.

In other embodiments, n and m are each 1.

In still other embodiments, at least one of R¹, R², R³ or R⁴ is methyl. For
example in some embodiments, each of R¹, R², R³ and R⁴ is methyl.

In other embodiments, at least one of R¹, R², R³ or R⁴ is hydrogen. For
example in some embodiments, each of R¹, R², R³ and R⁴ is hydrogen.

In other examples, R¹ and R² or R³ and R⁴ join to form substituted or
unsubstituted cyclohexyl. For example in some embodiments, each of R¹ and R² and R³
and R⁴ join to form substituted or unsubstituted cyclohexyl.

In still other embodiments, at least one of R⁷, R⁸, R⁹, R¹⁰, R¹¹ or R¹² is
hydrogen. For example in some embodiments, each of R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are
hydrogen.

In certain other embodiments, at least one of R⁷, R⁸, R⁹, R¹⁰, R¹¹ or R¹² is
methyl. For example, each of R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are methyl in some
embodiments. In other embodiments, each of R⁷, R⁸, R⁹, and R¹² are methyl.

In still other embodiments, at least one of R⁷, R⁸, R⁹, R¹⁰, R¹¹ or R¹² is
fluoro. For example in some embodiments, each of R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are
fluoro. In certain further embodiments, each of R⁷, R⁸, R⁹, and R¹² are fluoro.

In other aspects, at least one of R⁷, R⁸, R⁹, R¹⁰, R¹¹ or R¹² is chloro. For
example in some embodiments, each of R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are chloro, while in
other examples each of R⁷, R⁸, R⁹, and R¹² are chloro.

In other embodiments, at least one of R⁷, R⁸, R⁹, R¹⁰, R¹¹ or R¹² is
bromo. For example in some embodiments, each of R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are
bromo, and in other embodiments each of R⁷, R⁸, R⁹, and R¹² are bromo.
In other embodiments of the foregoing compound, 

\[
\begin{align*}
\text{Q} &
\end{align*}
\]
; n is 0, 1, 2, 3, 4, 5, 6, 7 or 8; each of q, r, and t is independently 0, 1, 2, 3, 4, 5, 6 or 7; m is 0, 1, 2, 3, 4, 5, 6, 7 or 8; each of u, j and y is independently 0, 1, 2, 3, 4, 5, 6 or 7 and each G is independently linear or branched, substituted or unsubstituted, saturated or unsaturated C1-C10 alkyl, wherein the optional substituents are selected from oxo, OJ', COOH, OH, F, Cl, Br, I, NH2, CN, SH, S02H, CONH2, OPO3H3 and NO2.

In still other embodiments, Q is ;
n is 0, 1, 2, 3, 4, 5, 6, 7 or 8; each of q, r, and t is independently 0, 1, 2, 3, 4, 5, 6 or 7; m is 0, 1, 2, 3, 4, 5, 6, 7 or 8; each of u, j and y is independently 0, 1, 2, 3, 4, 5, 6 or 7 and each G¹ is independently linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cio alkyl, wherein the optional substituents are selected from oxo, OJ™, COOH, OH, F, Cl, Br, I, NH₂, CN, SH, SO₃H, CONH₂, OP0₃H₃, and NO₂.

In other embodiments, Q is
4, 5, 6, 7 or 8; m is 0, 1, 2, 3, 4, 5, 6, 7 or 8; and each $G^1$ is independently linear or branched, substituted or unsubstituted, saturated or unsaturated $C_1-C_i$ alkyl, wherein the optional substituents are selected from oxo, OJ', COOH, OH, F, Cl, Br, I, NH$_2$, CN, SH, S0$_3$H, CONH$_2$, OPO$_3$H$_3$, and NO$_2$.

In still other embodiments, Q is

$n$ is 0, 1, 2, 3, 4, 5, 6, 7 or 8; m is 0, 1, 2, 3, 4, 5, 6, 7 or 8; and each $G^1$ is
independently linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cio alkyl, wherein the optional substituents are selected from oxo, OJ", COOH, OH, F, Cl, Br, I, NH₂, CN, SH, SO₃H, CONH₂, OP0₃H₃, and NO₂.

In yet other embodiments, q is

In other embodiments, Q is
In certain other embodiments of the foregoing compound, T is
and n, m and q are each independently 0, 1, 2, 3, 4, 5, 6, 7 or 8.

In still other embodiments, T is
In other examples, Q is

![Chemical structures](image)

and T is

![Chemical structure](image)
In certain further embodiments, $Q$ is

\[
\text{or} \quad \text{and } T \text{ is}
\]

In other embodiments, $Z^1, Z^4$ and each remaining $Z^1$ and $Z^2$ is, at each occurrence, independently $\text{CH}_3; \text{CH}; \text{CF}, \text{CCl}$ or $\text{CBr}$. For example in some embodiments, $Z^1, Z^4$ and each remaining $Z^1$ and $Z^2$ is, at each occurrence, $\text{CH}$.

In some other embodiments, one or more of the $\text{OH}$ groups of any one of the foregoing compounds of Formula I is substituted to replace the $\text{H}$ with a moiety from TABLE 1. For example in some embodiments, the moiety from TABLE 1 is

\[
\text{or} \quad \text{or}
\]

In another embodiment, the present disclosure provides a compound having one of the following structures:
For example, in some other embodiments, one or more of the OH groups of the foregoing compounds is substituted to replace the H with a moiety from TABLE 1. For example in some embodiments, the moiety from TABLE 1 is or .

In another embodiment, the present disclosure provides the use of any one of the foregoing compounds for modulating androgen receptor (AR) activity. For example in some embodiments, modulating androgen receptor (AR) activity is in a mammalian cell.

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 muscular atrophy, and age-related macular degeneration. For example in some embodiments, the indication is prostate cancer. In other embodiments,
the prostate cancer is castration resistant prostate cancer. 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 other embodiments, the present disclosure provides a method of modulating androgen receptor (AR) activity, the method comprising administering any one of the foregoing compounds, or pharmaceutically acceptable salt thereof to a subject in need thereof.

In other further embodiments of the foregoing method, modulating androgen receptor (AR) activity is for the treatment of one or more of the following: 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. For example in some embodiments, the prostate cancer is castration resistant prostate cancer. In other embodiments, the prostate cancer is androgen-dependent prostate cancer, and in other embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease.

In some other embodiments, the present disclosure provides a pharmaceutical composition comprising any one of the foregoing compounds and a pharmaceutically acceptable carrier.

In yet another embodiment, the present disclosure provides a pharmaceutical composition comprising any one of the foregoing compounds, an additional therapeutic agent and a pharmaceutically acceptable carrier. For example, 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 MDV3100, TOK 001, TOK 001; ARN-509; abiraterone, bicalutamide, nilutamide, flutamide, cyproterone acetate, docetaxel, Bevacizumab (Avastin), OSU-HDAC42, VTTAXIN, sunitumib, ZD-4054, VN/124-1, Cabazitaxel (XRP-6258), MDX-010 (Ipilimumab), OGX 427, OCX 011, finasteride, dutasteride, turosteride, bexlosteride, izonsteride, FCE 28260, SKF 105,111 or a related compound thereof.
In another embodiment, the present disclosure provides the use of any of the foregoing pharmaceutical compositions for modulating androgen receptor (AR) activity. For example in some embodiments, modulating androgen receptor (AR) activity is in a mammalian cell.

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 muscular atrophy, and age-related macular degeneration. For example in some embodiments, the indication is prostate cancer. For example, in some embodiments, the prostate cancer is castration resistant prostate cancer, and in other embodiments the prostate cancer is androgen-dependent prostate cancer. In still other embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease.

In yet another embodiment, the present disclosure provides a method of modulating androgen receptor (AR) activity, the method comprising administering any one of the foregoing pharmaceutical compositions to a subject in need thereof. For example in some embodiments, modulating androgen receptor (AR) activity is for the treatment of one or more of the following: 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. In other embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease. In still other embodiments, the indication is prostate cancer. For example, in some embodiments, the prostate cancer is castration resistant prostate cancer, while in other embodiments, the prostate cancer is androgen-dependent prostate cancer.

Each J may independently be \(G^1\), \(O\), \(CH_2\), \(CHG^1\), \(CG^\wedge\), \(S\), \(NH\), \(NG^1\), \(SO\), \(SO_2\), or \(NR\). Each J may independently be \(G^1\), \(O\), \(CH_2\), \(CHG^1\), \(CG^\wedge\), \(S\), \(NH\), or \(NG^1\). Each J may independently be \(O\), \(S\), \(NH\), \(NG^1\), \(SO\), \(SO_2\), or \(NR\). Each J may independently be \(O\), \(S\), \(SO\), or \(SO_2\). Each J may independently be \(O\), \(NH\), \(NG^1\), or \(NR\). Each J may independently be \(S\), \(NH\), \(NG^1\), \(SO\), \(SO_2\), or \(NR\). Each J may independently be \(S\), \(SO\), or \(SO_2\). Each J may independently be \(NH\), \(NG^1\), or \(NR\). Each J may independently be \(G^1\), \(CH_2\), \(CHG^1\), or \(CG^\wedge\). Each J may independently be \(O\), \(CH_2\), \(S\), or \(NH\). Each J may independently be \(O\), \(CH_2\), or \(NH\). Each J may independently be \(O\), or
CH₂. Each J may independently be G₁, O, CHG₁, or NH. Each J may independently be G₁, O, or CHG₁. Each J may independently be G₁, O, or G. Each J may independently be O, or S. Each J may independently be G₁. Each J may independently be CH₂. Each J may be CHG₁. Each J may be CH₂O. Each J may be NH. Each J may be S. Each J may be O.

Each M may independently be H, Cl, Br, CH₂OH, CH₂OD, CH₂OG₁, CH₂Cl, CHCl₂, CC₁₃, CH₂Br, CHBr₂, CBr₃, or C≡CH. Each M may independently be Cl, Br, CH₂OH, CH₂OD, CH₂OG₁, CH₂F, CH₂Cl, CHCl₂, CC₁₃, CH₂Br, CHBr₂, or CBr₃. Each M may independently be Cl, CH₂Cl, CHCl₂, or CC₁₃. Each M may independently be Br, CH₂Br, CHBr₂, or CBr₃. Each M may independently be Cl, or Br. Each M may independently be CH₂Cl, or CH₂Br. Each M may independently be CHCl₂, or CHBr₂. Each M may independently be Br, CH₂Br, CHBr₂, or CBr₃. Each M may independently be Cl, or CC₁₃, or CH₂Cl, or CHCl₂. Each M may independently be Br, or CBr₃. Each M may be H. Each M may be Br. Each M may be CHCl₂. Each M may be Cl, or CC₁₃. Each M may be Br, CH₂Br, CHBr₂, or CBr₃. Each M may be C≡CH. Each M may be CH₂Cl. Each M may be CH₂F. Each M may be CH₂O. Each M may be CH₂OD. Each M may be CH₂OG₁.

Each L may independently be H or A-D. Each L may be H. Each L may be A-D.

Each A may independently be O, S, NH, NG₁, N⁺H₂, or N⁺HG₁. Each A may independently be O, NH, or N⁺H₂. Each A may independently be O, S, NH, or N⁺H₂. Each A may independently be O, S, or NH. Each A may independently be O, or NH. Each A may independently be O, or S. Each A may be S. Each A may be NH. Each A may be NG₁. Each A may be N⁺H₂. Each A may be N⁺HG₁. Each A may be O.

Each D may independently be H, G₁, R, O/₉ or a moiety selected from
TABLE 1. Each D may independently be H, G\textsuperscript{1}, or R. Each D may independently be H, or G\textsuperscript{1}, or R. Each D may independently be G\textsuperscript{1}, or R. Each D may independently be H, or G\textsuperscript{1}.

Each D may independently be

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<tr>
<th>Structure</th>
<th>Structure</th>
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<tr>
<td>\begin{align*} &amp; \text{O}<em>{q}^{\text{r}} \rightarrow \text{OH} \ &amp; \text{O}</em>{t}^{\text{r}} \rightarrow \text{OR} \end{align*}</td>
<td>\begin{align*} &amp; \text{O}<em>{q}^{\text{r}} \rightarrow \text{OH} \ &amp; \text{O}</em>{t}^{\text{r}} \rightarrow \text{OG}^{\text{1}} \end{align*}</td>
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<tbody>
<tr>
<td>\begin{align*} &amp; \text{O}<em>{q}^{\text{r}} \rightarrow \text{OH} \ &amp; \text{O}</em>{t}^{\text{r}} \rightarrow \text{OR} \end{align*}</td>
<td>\begin{align*} &amp; \text{O}<em>{q}^{\text{r}} \rightarrow \text{OH} \ &amp; \text{O}</em>{t}^{\text{r}} \rightarrow \text{OG}^{\text{1}} \end{align*}</td>
</tr>
</tbody>
</table>

, or a moiety selected from TABLE 1. Each D may be H. Each D may be G\textsuperscript{1}. Each D may be R. Each D may be G\textsuperscript{1}.

Each J\textsuperscript{2} may independently be G\textsuperscript{1}, O, CH\textsubscript{2}, CHG\textsuperscript{1}, CG\textsuperscript{\wedge}, S, NH, NG\textsuperscript{1}, SO, S0\textsubscript{2}, or NR. Each J\textsuperscript{2} may independently be G\textsuperscript{1}, O, CH\textsubscript{2}, CHG\textsuperscript{1}, CG\textsuperscript{\wedge}, S, NH, or NG\textsuperscript{1}. Each J\textsuperscript{2} may independently be O, S, NH, NG\textsuperscript{1}, SO, S0\textsubscript{2}, or NR. Each J\textsuperscript{2} may independently be O, S, SO, or S0\textsubscript{2}. Each J\textsuperscript{2} may independently be O, NH, NG\textsuperscript{1}, or NR. Each J\textsuperscript{2} may independently be S, NH, NG\textsuperscript{1}, SO, S0\textsubscript{2}, or NR. Each J\textsuperscript{2} may independently be S, SO, or S0\textsubscript{2}. Each J\textsuperscript{2} may independently be NH, NG\textsuperscript{1}, or NR. Each J\textsuperscript{2} may independently be G\textsuperscript{1}, CH\textsubscript{2}, CHG\textsuperscript{1}, or CG\textsuperscript{\wedge}. Each J\textsuperscript{2} may independently be O,
CH₂, S, or NH. Each J² may independently be O, CH₂, or NH. Each J² may independently be O, or CH₂. Each J² may independently be G¹, O, CHG¹, or NH. Each J² may independently be G¹, O, or CHG¹. Each J² may independently be G¹, or O. Each J² may independently be O, or S. Each J² may independently be G¹. Each J² may independently be CH₂. Each J² may be CHG¹. Each J² may be CG⁴. Each J² may be NR. Each J² may be S0₂. Each J² may be SO. Each J² may be NG¹. Each J² may be NH. Each J² may be S. Each J² may be O.

Each M² may independently be H, CH₃, Cl, Br, CH₂F, CH₂Cl, CHCl₂, CC₁₃, CH₂Br, CHBr₂, CBr₃, CH₂OH, CH₂O¹", G¹, CH₂OG¹, CH₂OR, CH₂OG¹OG¹, G¹OG¹, G¹OG¹OG¹", CH₂SG¹, CH₂NH₂, CH₂NHG¹, CH₂NG¹, or C=CH. Each M² may independently be H, CH₃, CH₂Cl, CH₂Br, CH₂O¹", CH₂OG, CH₂OGG', GOG', GOGG', CH₂SG, CH₂NH₂, CH₂NHG, or CH₂NG₂. Each M² may independently be H, CH₃, CH₂Cl, CH₂Br, CH₂O¹", CH₂OG, or CH₂OGG'. Each M² may independently be CH₂Cl, CH₂Br, CH₂OH, CH₂OCH₃, CH₂O(isopropyl), or CH₂OC₂H₄OC₄H₉. Each M² may independently be H, CH₃, CH₃OCH₃, CH₃OCH₂CH₃, CH₃Cl, or CH₂Br. Each M² may independently be CH₃, CH₃OCH₂CH₃, CH₂Cl, CH₂Br, CH₂OH, CH₂OCH₃, or CH₂O(isopropyl). Each M² may independently be CH₃, CH₂Cl, CH₂Br, CH₂OH, CH₃OCH₂CH₃, or CH₂OCH₃. Each M² may independently be CH₃, CH₂Cl, CH₂Br, CH₂OH, or CH₂OCH₃. Each M² may independently be CH₃, CH₂OH, CH₂OCH₃, or CH₂OCH₂CH₃. Each M² may independently be CH₂Cl, or CH₂Br. Each M² may independently be H, Cl, Br, CH₂Cl, CHCl₂, CCl₃, CH₂Br, CHBr₂, CBr₃, or C=CH. Each M² may independently be Cl, Br, CH₂Cl, CHCl₂, CCl₃, CH₂Br, CHBr₂, or CBr₃. Each M² may independently be Cl, CH₂Cl, CHCl₂, CCl₃, CH₂Br, CHBr₂, or CBr₃. Each M² may independently be Cl, CH₂Cl, CHCl₂, or CCl₃. Each M² may independently be Br, CH₂Br, CHBr₂, or CBr₃. Each M² may independently be Cl, or Br. Each M² may independently be CH₂Cl, or CH₂Br. Each M² may independently be CHCl₂, or CHBr₂. Each M² may independently be CCl₃, or CBr₃. Each M² may independently be CH₂Cl, CHCl₂, or CCl₃. Each M² may independently be CH₂Br, CHBr₂, or CBr₃. Each M² may independently be Br, CH₂Br, or CHBr₂. Each M² may independently be CH₂Cl, or CHCl₂. Each M² may independently be Br, CH₂Br, or CHBr₂. Each M² may independently be CH₂Cl, or CHCl₂.
independently be Br, or CBr₃. Each M² may be H. Each M² may be CH₃. Each M² may be Cl. Each M² may be Br. Each M² may be CH₂Cl. Each M² may be CHCl₂. Each M² may be CC₁₂. Each M² may be CH₂Br. Each M² may be CHBr₂. Each M² may be CBr₃. Each M² may be CH₂OH. Each M² may be CH₂O₂. Each M² may be CH₂OR. Each M² may be CH₂OG₁. Each M² may be CH₂OG₁O₂. Each M² may be G⁺G⁻. Each M² may be G⁺G⁺OG⁻. Each M² may be CH₂SG₁. Each M² may be CH₂NH₂. Each M² may be CH₂NHG₁. Each M² may be CH₂NG₂. Each M² may be C≡CH. Each M² may be CH₂F.

Each L² may independently be H or A²-D². Each L² may be H. Each L² may be A²-D².

Each A² may independently be O, S, SO, S₀₂, NH, NG₁, N⁺H₂, or N⁺H⁺G⁺. Each A² may independently be O, S, SO, or S₀₂. Each A² may independently be O, NH, NG₁, N⁺H₂, or N⁺H⁺G⁺. Each A² may independently be S, SO, S₀₂, NH, NG₁, N⁺H₂, or N⁺H⁺G⁺. Each A² may independently be O, S, SO, S₀₂, NH, or N⁺H₂. Each A² may independently be S, SO, or S₀₂. Each A² may independently be NH, NG₁, N⁺H₂, or N⁺H⁺G⁺. Each A² may independently be NH, or N⁺H⁺G⁺. Each A² may independently be O, S, NH, NG₁, N⁺H₂, or N⁺H⁺G⁺. Each A² may independently be O, NH, or N⁺H⁺G⁺. Each A² may independently be O, S, NH, or N⁺H₂. Each A² may independently be O, S, NH. Each A² may independently be O, or NH. Each A² may independently be O, S, or NH. Each A² may independently be O, or NH. Each A² may independently be O, S, or NH. Each A² may be S₀₂. Each A² may be NH. Each A² may be NG₁. Each A² may be N⁺H₂. Each A² may be N⁺H⁺G⁺. Each A² may be O.

Each D² may independently be H, G₁, R, , or a moiety selected from TABLE 1. Each D² may independently be H, G₁, or R. Each D² may independently be H, or R. Each D² may independently be G₁, or R. Each D² may independently be H, or G₁. Each D² may independently be , or , or , or .
or a moiety selected from TABLE 1. Each D\(^2\) may independently be

Each D\(^2\) may be H. Each D\(^2\) may be G\(^1\). Each D\(^2\) may be R. Each D\(^2\) may be

Each D\(^2\) may be a moiety selected from TABLE 1.

Each Q may independently be
Each $Q$ may independently be:

- $\text{Cl}$
- $\text{OH}$
- $\text{OR}$
- $\text{OG}^i$
- $\text{Br}$
- $\text{OH}$
- $\text{OR}$
- $\text{OG}^i$
Each Q may independently be

Each Q may independently be

Each Q may independently be

Each Q may independently be
Each Q may independently be

or

Each may independently be

or

Each may independently be

Each may independently be

or

Each may independently be

Each may independently be

or

Each may independently be

Each may independently be

or
Each Q may independently be
Each may independently be

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Each Q may independently be 
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B r c i O "Cl Ό' 
may independently be 
Each Q may independently
Each Q may independently
be
or Cl. Each Q may independently be
or Cl. Each Q may independently be
or Cl. Each Q may independently be
or Cl. Each Q may independently be
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or Cl. Each Q may independently be
or Cl. Each Q may independently be
or Cl. Each Q may independently be
or Cl. Each Q may independently be
or Cl.
Each $Q$ may independently be

$O(H)$

Each $Q$ may independently be

$O^iG$

Each $Q$ may independently be

$O^iR$

Each $Q$ may independently be

$O^iF$

Each $Q$ may independently be

$O(H)$

Each $Q$ may independently be

$O^iG$

Each $Q$ may independently be

$O^iR$

Each $Q$ may independently be

$O^iF$

Each $T$ may independently be

$O^iCl$

Each $T$ may independently be

$O^iCl$

Each $T$ may independently be

$O^iCl$

Each $T$ may independently be

$O^iCl$

Each $T$ may independently be

$O^iCl$

Each $T$ may independently be

$O^iCl$
Each T may independently be
Each T may independently be
Each T may independently be

or

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independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each q may independently be 3
to 4, 3 to 5, 3 to 6, or 3 to 7. Each q may be 0. Each q may be 1. Each q may be 2. Each
q may be 3. Each q may be 4. Each q may be 5. Each q may be 6. Each q may be 7.

Each r may independently be 0, 1, 2, 3, 4, 5, 6 or 7. Each r may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, 0 to 7. Each r may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, or 1 to 7. Each r may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each r may independently be 3 to 4, 3 to 5, 3 to 6, or 3 to 7. Each r may be 0. Each r may be 1. Each r may be 2. Each r may be 3. Each
r may be 4. Each r may be 5. Each r may be 6. Each r may be 7.

Each t may independently be 0, 1, 2, 3, 4, 5, 6 or 7. Each t may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, or 0 to 7. Each t may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, or 1 to 7. Each t may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each t may independently be 3 to 4, 3 to 5, 3 to 6, or 3 to 7. Each t may be 0. Each t may be 1. Each t may be 2. Each t may be 3. Each t
may be 4. Each t may be 5. Each t may be 6. Each t may be 7.

Each n may independently be 0, 1, 2, 3, 4, 5, 6, 7 or 8. Each n may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, 0 to 7, or 0 to 8. Each n may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, 1 to 7, or 1 to 8. Each n may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, 2 to 7, or 2 to 8. Each n may independently be 3 to 4, 3 to 5, 3 to 6, 3 to 7, or 3 to 8. Each n may be 0. Each n may be 1. Each n may be 2. Each n may be 3. Each n may be 4. Each n may be 5. Each n may be 6. Each n may be 7. Each n may be 8.

Each u may independently be 0, 1, 2, 3, 4, 5, 6 or 7. Each u may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, 0 to 7. Each u may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, or 1 to 7. Each u may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each u may independently be 3 to 4, 3 to 5, 3 to 6, or 3 to 7. Each u may be 0. Each u may be 1. Each u may be 2. Each
u may be 3. Each u may be 4. Each u may be 5. Each u may be 6. Each u may be 7.

Each y may independently be 0, 1, 2, 3, 4, 5, 6 or 7. Each y may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, or 0 to 7. Each y may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, or 1 to 7. Each y may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each y may independently be 3
to 4, 3 to 5, 3 to 6, or 3 to 7. Each y may be 0. Each y may be 1. Each y may be 2. Each y may be 3. Each y may be 4. Each y may be 5. Each y may be 6. Each y may be 7.

Each j may independently be 0, 1, 2, 3, 4, 5, 6 or 7. Each j may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, or 0 to 7. Each j may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, or 1 to 7. Each j may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, or 2 to 7. Each j may independently be 3 to 4, 3 to 5, 3 to 6, or 3 to 7. Each j may be 0. Each j may be 1. Each j may be 2. Each j may be 3. Each j may be 4. Each j may be 5. Each j may be 6. Each j may be 7.

Each m may independently be 0, 1, 2, 3, 4, 5, 6, 7 or 8. Each m may independently be 0 to 1, 0 to 2, 0 to 3, 0 to 4, 0 to 5, 0 to 6, 0 to 7, or 0 to 8. Each m may independently be 1 to 2, 1 to 3, 1 to 4, 1 to 5, 1 to 6, 1 to 7, or 1 to 8. Each m may independently be 2 to 3, 2 to 4, 2 to 5, 2 to 6, 2 to 7, or 2 to 8. Each m may independently be 3 to 4, 3 to 5, 3 to 6, 3 to 7, or 3 to 8. Each m may be 0. Each m may be 1. Each m may be 2. Each m may be 3. Each m may be 4. Each m may be 5. Each m may be 6. Each m may be 7. Each m may be 8.

At least one \( Z^1 \) is C-Q, at least one \( Z^2 \) is C-T, CH, CCH\(_3\), CF, CC1, CBr, Cl, COH, CG\(_1\), COG\(_1\), CNH\(_2\), CNHG\(_1\), CNG\(_1\), COS0\(_3\)H, COP0\(_3\)H\(_2\), CSG\(_1\), CSOG\(_1\), or CSO\(_2\)G\(_1\), and each \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) is, at each occurrence, independently C-T, N, CH, CCH\(_3\), CF, CC1, CBr, Cl, COH, CG\(_1\), COG\(_1\), CNH\(_2\), CNHG\(_1\), CNG\(_1\), COS0\(_3\)H, COP0\(_3\)H\(_2\), CSG\(_1\), CSOG\(_1\), or CSO\(_2\)G\(_1\) or one of \( Z^3 \), \( Z^4 \) or one of the remaining \( Z^1 \) or \( Z^2 \) may independently be C-T, and \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) may, at each occurrence, independently be N, CH, CF, CC1, CBr, Cl, CG\(_1\), or COH. At least one \( Z^1 \) may independently be C-Q, at least one \( Z^2 \) may independently be COH, and each \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) may, at each occurrence, independently be N, CH, CF, CC1, CBr, Cl, CG\(_1\), or COH. At least one \( Z^1 \) may independently be C-Q, at least one \( Z^2 \) may independently be C-T, and each \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) may, at each occurrence, independently be N, CH, CF, CC1, CBr, Cl, or COH. At least one \( Z^1 \) may independently be C-Q, at least one \( Z^2 \) may independently be C-T, and each \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) may, at each occurrence, independently be N, CH, CF, CC1, CBr, Cl, or COH. At least one \( Z^1 \) may independently be C-Q, at least one \( Z^2 \) may independently be C-T, and each \( Z^3 \), \( Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) may, at each occurrence, independently be CH. At least one \( Z^1 \) may independently be C-Q, at least one \( Z^2 \) may independently be COH, and each \( Z^3 \),
Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be CH. In any of the foregoing embodiments, each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be N, CH, CF, CCl, CBr, Cl, COH, CCH_3, CNH_2, COSO_3H, or COP03H_2, each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be N, CH, CF, CCl, CBr, Cl, COH, CNH_2, COSO_3H, or COP03H_2, each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be N, CH, CF, CCl, CBr, Cl, or COH, each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be CH, CF, CCl, CBr, or Cl, each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be CH, CCl, or CBr or each Z^3, Z^4 and each remaining Z^1 and Z^2 may, at each occurrence, independently be CH.

Each of J" and J" may independently be a moiety selected from TABLE 1. Each of J", and V" may independently be an amino acid based moiety or a polyethylene glycol based moiety selected from TABLE 1. Alternatively, each of J", and V" may independently an amino acid based moiety selected from TABLE 1. Each J", and J" may be

Each G^1 G^1' and G^1" may independently be linear or branched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{10} alkyl. Each G^1, G^1' and G^1" may independently be a branched, linear, or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{10} alkyl. Each G^1, G^1' and G^1" may independently be a branched, linear, or non-aromatic cyclic, substituted or saturated or unsaturated C_1-C_{10} alkyl. Each G^1, G^1' and G^1" may independently be a branched, unbranched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{9} alkyl. Each G^1, G^1' and G^1" may independently be a branched, unbranched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{7} alkyl. Each G^1, G^1' and G^1" may independently be a branched, unbranched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{9} alkyl. Each G^1, G^1' and G^1" may independently be a branched, unbranched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C_1-C_{7} alkyl.
or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C6 alkyl. Each G1, G1' and G1'' may independently be a branched, unbranched, or aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C5 alkyl. Each G1, G1' and G1'' may independently be a branched, unbranched, or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C4 alkyl. Each G1, G1' and G1'' may independently be a branched, unbranched, or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C3 alkyl. Each G1, G1' and G1'' may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated C1-C2 alkyl.

Each G1 may independently be a non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclopropyl. Each G1 may independently be a non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclobutyl. Each G1 may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclobutyl. Each G1 may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclopentyl. Each G1 may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclohexyl. Each G1 may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cycloheptyl. Each G1 may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclooctyl. Each G1 may independently be cyclohexyl.

Each G1 may independently be substituted or unsubstituted methyl. Each G1 may independently be substituted or unsubstituted, saturated or unsaturated ethyl. Each G1 may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated propyl. Each G1 may be isopropyl. Each G1 may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated butyl. Each G1 may be n-butyl. Each G1 may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated penty1. Each G1 may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated hexyl. Each G1 may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated heptyl. Each G1 may independently be a
branched, unbranched, substituted or unsubstituted, saturated or unsaturated octyl. Each G1 may be propargyl (CH2C≡CH).

An optional substituent may be selected from the group consisting of oxo, OJ", COOH, R4, OH, OR4, F, Cl, Br, I, NH2, NHR4, NR4, CN, SH, SR4, S03H, S03R4, SO3R4, OR5, CO2R4, CONH2, CONHR4, CONHR5, CONR4, NHR5, OP03H3, CONR4R5, NR4R5, and NO2. An optional substituent may be selected from the group consisting of oxo (i.e. =0), OJ", COOH, R4, OH, OR4, F, Cl, Br, I, NH2, NHR4, NR42, CN, SH, SR4, S03H, S03R4, S02R4, OS03R4, and NO2. An optional substituent may be selected from the group consisting of oxo (i.e. =0), OJ", COOH, R4, OH, OR4, F, Cl, Br, I, NH2, and NO2. An optional substituent may be selected from the group consisting of oxo (i.e. =0), OJ", COOH, R4, OH, OR4, F, Cl, Br, and I. An optional substituent may be selected from the group consisting of oxo (i.e. =0), OJ", COOH, OH, F, Cl, Br, and I. An optional substituent may be selected from the group consisting of oxo (i.e. =0), OJ", COOH, OH, F, and Cl. Each linear or branched, or aromatic cyclic or non-aromatic cyclic, saturated or unsaturated Ci-Cio alkyl may be substituted with, for example, 1, 2, 3, 4, 5, or 6 substituents.

Each R' may independently be a non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclopropyl. Each R' may independently be a non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclobutyl. Each R' may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclobutyl. Each R' may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclopentyl. Each R' may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclohexyl. Each R' may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cycloheptyl. Each R' may independently be an aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated cyclooctyl.
Each R may independently be substituted or unsubstituted methyl. Each R may independently be substituted or unsubstituted, saturated or unsaturated ethyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated propyl. Each R may be isopropyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated butyl. Each R may be n-butyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated pentyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated hexyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated heptyl. Each R may independently be a branched, unbranched, substituted or unsubstituted, saturated or unsaturated octyl.

Each R may independently be H. Each R may independently be Li. Each R may independently be Na. Each R may independently be K. Each R may independently be Mg. Each R may independently be Ca.

Each R may independently be C1-C9 acyl. Each R may independently be C1-Cg acyl. Each R may independently be C1-C7 acyl. Each R may independently be C1-C6 acyl. Each R may independently be C1-C5 acyl. Each R may independently be C1-C4 acyl. Each R may independently be C1-C3 acyl. Each R may independently be C1-C2 acyl. Each R may independently be Ci acyl. Each R may independently be C2 acyl. Each R may independently be C3 acyl. Each R may independently be C4 acyl. Each R may independently be C5 acyl. Each R may independently be C6 acyl. Each R may independently be C7 acyl. Each R may independently be C8 acyl. Each R may independently be C9 acyl. Each R may independently be C10 acyl.

Each R1 may independently be unsubstituted C1-C10 alkyl. Each R1 may independently be unsubstituted C1-C9 alkyl. Each R1 may independently be unsubstituted Ci-Cg alkyl. Each R1 may independently be unsubstituted Ci-C7 alkyl. Each R1 may independently be unsubstituted Ci-C6 alkyl. Each R1 may independently be unsubstituted Ci-C5 alkyl. Each R1 may independently be unsubstituted Ci-C4 alkyl. Each R1 may independently be unsubstituted Ci-C3 alkyl. Each R1 may independently be unsubstituted Ci-C2 alkyl. Each R1 may independently be unsubstituted Ci alkyl. Each R1 may independently be unsubstituted C2 alkyl. Each R1 may independently be
unsubstituted C₃ alkyl. Each R₁ may independently be unsubstituted C₄ alkyl. Each R₁ may independently be unsubstituted C₅ alkyl. Each R₁ may independently be unsubstituted C₆ alkyl. Each R₁ may independently be unsubstituted C₇ alkyl. Each R₁ may independently be unsubstituted C₈ alkyl. Each R₁ may independently be unsubstituted C₉ alkyl. Each R₁ may independently be unsubstituted C₁₀ alkyl.

Each R² may independently be C₁-C₉ acyl. Each R² may independently be C₁-C₁₀ acyl. Each R² may independently be C₁-C₇ acyl. Each R² may independently be C₁-C₆ acyl. Each R² may independently be C₁-C₄ acyl. Each R² may independently be C₁-C₃ acyl. Each R² may independently be C₁-C₂ acyl. Each R² may independently be C₂ acyl. Each R² may independently be C₃ acyl. Each R² may independently be C₄ acyl. Each R² may independently be C₅ acyl. Each R² may independently be C₆ acyl. Each R² may independently be C₇ acyl. Each R² may independently be C₈ acyl. Each R² may independently be C₉ acyl. Each R² may independently be C₁₀ acyl.

R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently hydrogen, halo, or linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₁₀ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently hydrogen. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently hydrogen. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently fluoro. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently fluoro. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently chloro. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently chloro. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently bromo. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently bromo. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently methyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently methyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently ethyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently ethyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₃ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₃ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₄ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₄ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₆ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₆ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₇ alkyl. Each R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₇ alkyl. R₃,
R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₈ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₈ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₉ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₉ alkyl. R₃, R₄, R₅, R₆, R₇, R₈, R₉ and R₁₀ are each independently C₁₀ alkyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently C₁₀ alkyl.

Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently hydrogen. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently fluoro. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently fluoro. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently chloro. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently chloro. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently chloro. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently chloro.

Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently bromo. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently bromo. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently methyl. At least one of R₃, R₄, R₅, R₆, R₇, R₈, R₉ or R₁₀ is independently methyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently methyl.

Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₃ alkyl.
Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and R₃, R₄, R₅ and R₆ are each independently C₉ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₉ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and R₃, R₄, R₅ and R₆ are each independently C₁₀ alkyl. Each of R₇, R₈, R₉ and R₁₀ may be hydrogen, and at least one of R₃, R₄, R₅ or R₆ is independently C₁₀ alkyl.

Each R₁₁ may independently be hydrogen. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₉ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₁₀ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-Cₙ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₁₅ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₄ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₃ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₂ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₂ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₃ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₄ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₅ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₆ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₇ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₈ alkyl. Each R₁₁ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₉ alkyl.
unsubstituted, saturated or unsaturated C₉ alkyl. Each R¹¹ may independently be linear or branched, substituted or unsubstituted, saturated or unsaturated C₁₀ alkyl.

The compounds described herein are meant to include all racemic mixtures and all individual enantiomers or combinations thereof, whether or not they are specifically depicted herein. Alternatively, one or more of the OH groups on the above compounds may be substituted to replace the H with a moiety selected from TABLE 1.

In yet other embodiments, the present disclosure provide the use of any of the compounds disclosed herein for modulating androgen receptor (AR) activity. For example, in certain embodiments modulating androgen receptor (AR) activity is in a mammalian cell.

In other examples, 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 muscular atrophy, and age-related macular degeneration. For example, in certain embodiments the indication is prostate cancer, for example, castration resistant prostate cancer. In other examples, the prostate cancer is androgen-dependent prostate cancer. In other further embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease.

The present disclosure also provides a method of modulating androgen receptor (AR) activity, the method comprising administering any of the compounds disclosed herein, or pharmaceutically acceptable salt thereof, to a subject in need thereof. For example, in certain specific embodiments modulating androgen receptor (AR) activity is for the treatment of one or more of the following: 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. In certain embodiments, the spinal and bulbar muscular atrophy is Kennedy's disease.

The present disclosure also provides a pharmaceutical composition comprising any one or more of the compounds disclosed herein and a pharmaceutically acceptable carrier. The pharmaceutical composition may be for treating one or more of
the following: 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.

In accordance with another embodiment, there is provided a use of the compounds of Formula (I) as described anywhere herein for preparation of a medicament for modulating androgen receptor (AR).

In accordance with a further embodiment, there is provided a method of screening for androgen receptor modulating compounds, wherein the compounds screened are selected from the compounds as described anywhere herein.

The modulating of the androgen receptor (AR) activity may be in a mammalian cell. The modulating of the androgen receptor (AR) activity may be in a mammal. The mammal may be a human.

Alternatively, the administering may be to a mammal. The administering may 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, 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), and age-related macular degeneration.

The mammalian cell may be a human cell. The modulating AR activity may be for inhibiting AR N-terminal domain activity. The modulating AR activity may be for inhibiting AR activity. The modulating may be in vivo. The modulating AR activity may be 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 muscular atrophy (e.g., Kennedy's disease), and age-related macular degeneration. The indication may be prostate cancer. The prostate cancer may be castration-resistant prostate cancer. The prostate cancer may be androgen-dependent prostate cancer.
### TABLE 1

**MOIETIES**

<table>
<thead>
<tr>
<th>Amino Acid Based Moieties</th>
</tr>
</thead>
<tbody>
<tr>
<td>(aa) = any naturally occurring amino acid side chain</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Polyethylene Glycol Based Moieties</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{O} \quad \text{O} \quad \text{O} \quad \text{O} \quad \text{O} \quad \text{OH}$</td>
</tr>
<tr>
<td>$\text{O} \quad \text{S} \quad \text{O} \quad \text{O} \quad \text{OH}$</td>
</tr>
<tr>
<td>$n = 1-200$</td>
</tr>
<tr>
<td>$n = 1-200$</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Phosphate Based Moieties</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{O} \quad \text{H}$</td>
</tr>
<tr>
<td>$\text{O} \quad \text{N}^+$</td>
</tr>
<tr>
<td>MOIETIES</td>
</tr>
<tr>
<td>----------</td>
</tr>
<tr>
<td><img src="image1.png" alt="MOIETY 1" /></td>
</tr>
<tr>
<td><img src="image3.png" alt="MOIETY 3" /></td>
</tr>
<tr>
<td><img src="image5.png" alt="MOIETY 5" /></td>
</tr>
<tr>
<td><img src="image7.png" alt="MOIETY 7" /></td>
</tr>
</tbody>
</table>

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Moieties from TABLE 1 may be, for example, and without limitation, subdivided into three groups: 1) amino acid based moieties; 2) polyethylene glycol based moieties; and 3) phosphate based moieties. In the Moieties Table 1 above, the first four moieties are amino acid based moieties, the fifth and sixth are polyethylene glycol based moieties and the remaining moieties are phosphate based moieties.

The amino acid side chains of naturally occurring amino acids (as often denoted herein using "(aa)") are well known to a person of skill in the art and may be found in a variety of text books such as "Molecular Cell Biology" by James Darnell et al. Third Edition, published by Scientific American Books in 1995. Often the naturally occurring amino acids are represented by the formula \((\text{NH}_2)\text{C(COOH)}(\text{H})(\text{R})\), where the chemical groups in brackets are each bonded to the carbon not in brackets. \(\text{R}\) represents the side chains in this particular formula.

Those skilled in the art will appreciate that the point of covalent attachment of the moiety to the compounds as described herein may be, for example, and without limitation, cleaved under specified conditions. Specified conditions may include, for example, and without limitation, \textit{in vivo} enzymatic or non-enzymatic means. Cleavage of the moiety may occur, for example, and without limitation, spontaneously, or it may be catalyzed, induced by another agent, or a change in a physical parameter or environmental parameter, for example, an enzyme, light, acid,
temperature or pH. The moiety may be, for example, and without limitation, a
protecting group that acts to mask a functional group, a group that acts as a substrate for
one or more active or passive transport mechanisms, or a group that acts to impart or
enhance a property of the compound, for example, solubility, bioavailability or
localization.

In other particular embodiments of the compounds as described
anywhere herein, the following compounds in Table 2 are provided.

TABLE 2. Representative Compounds

<table>
<thead>
<tr>
<th>No.</th>
<th>Structure</th>
<th>Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td><img src="image1.png" alt="Structure" /></td>
<td>3,3’-(4,4’-(9H-fluorene-9,9-diy1)bis(4,1-phenylene))bis(oxy)bis(1-chloropropan-2-ol)</td>
</tr>
<tr>
<td>3</td>
<td><img src="image2.png" alt="Structure" /></td>
<td>3-(4-(9-(4-(3-chloro-2-hydroxypropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propane-1,2-diol</td>
</tr>
<tr>
<td>4</td>
<td><img src="image3.png" alt="Structure" /></td>
<td>3-(4-(9-(4-(2-hydroxy-3-(prop-2-ynyloxy)propoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propane-1,2-diol</td>
</tr>
<tr>
<td>No.</td>
<td>Structure</td>
<td>Name</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
<td>------</td>
</tr>
<tr>
<td>5</td>
<td><img src="image1" alt="Structure" /></td>
<td>$3,3'-(4,4'-(9H-fluorene-9,9-diyl)bis(4,1-phenylene))bis(oxy)dipropylene-1,2-diol</td>
</tr>
<tr>
<td>8</td>
<td><img src="image2" alt="Structure" /></td>
<td>$(2S,2'S)-3,3'-(4,4'-(9H-fluorene-9,9-diyl)bis(4,1-phenylene))bis(oxy)bis(1-chloropropan-2-ol)</td>
</tr>
<tr>
<td>9</td>
<td><img src="image3" alt="Structure" /></td>
<td>$(S)-1$-chboro-$3-(4-(9-(4-((R)-2-hydroxy-3-(prop-2-ynyloxy)propoxy)phenyl)-9H-fluorene-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>10</td>
<td><img src="image4" alt="Structure" /></td>
<td>$(S)-1$-chboro-$3-(4-(9-(4-((R)-2-hydroxy-3-isopropoxypropoxy)phenyl)-9H-fluorene-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>11</td>
<td><img src="image5" alt="Structure" /></td>
<td>$3,3'-(4,4'-(9H-fluorene-9,9-diyl)bis(4,1-phenylene))bis(oxy)bis(1-chloropropane-3,2-diyl)bis(2-(tert-butoxycarbonylamino)acetate)</td>
</tr>
<tr>
<td>No.</td>
<td>Structure</td>
<td>Name</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
<td>------</td>
</tr>
<tr>
<td>12</td>
<td><img src="image12.png" alt="Structure 12" /></td>
<td>2,2’-(3,3’-(4,4’-(9H-fluorene-9,9-diyl)bis(4,1-phenylene))bis(oxy)bis(1-chloropropane-3,2-diyl))bis(oxy)bis(2-oxoethanaminium) chloride</td>
</tr>
<tr>
<td>13</td>
<td><img src="image13.png" alt="Structure 13" /></td>
<td>(S)-1-chloro-3-(4-(9-(4-((R)-2-hydroxy-3-(2-(2-hydroxyethoxy)ethoxy)ethoxy)propoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>14</td>
<td><img src="image14.png" alt="Structure 14" /></td>
<td>(R)-3-(4-(9-(4-((S)-3-chloro-2-hydroxypropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propane-1,2-diol</td>
</tr>
<tr>
<td>15</td>
<td><img src="image15.png" alt="Structure 15" /></td>
<td>1-chloro-3-(4-(9-(4-(3-fluoro-2-hydroxypropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>No.</td>
<td>Structure</td>
<td>Name</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
<td>------</td>
</tr>
<tr>
<td>16</td>
<td><img src="image1.png" alt="Structure Image" /></td>
<td>(R)-1-butoxy-3-(4-(9-(4-((S)-3-chloro-2-hydroxypropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>17</td>
<td><img src="image2.png" alt="Structure Image" /></td>
<td>(S)-1-chloro-3-(4-(9-(4-((R)-3-(cyclohexyloxy)-2-hydroxypropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)propan-2-ol</td>
</tr>
<tr>
<td>18</td>
<td><img src="image3.png" alt="Structure Image" /></td>
<td>9-(4-(3-butoxypropoxy)phenyl)-9-(4-(3-chloropropoxy)phenyl)-9H-fluorene</td>
</tr>
<tr>
<td>19</td>
<td><img src="image4.png" alt="Structure Image" /></td>
<td>9-(4-(3-chloropropoxy)phenyl)-9-(4-(3-(cyclohexyloxy)propoxy)phenyl)-9H-fluorene</td>
</tr>
<tr>
<td>No.</td>
<td>Structure</td>
<td>Name</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
<td>------</td>
</tr>
<tr>
<td>20</td>
<td><img src="image1.png" alt="Structure Image" /></td>
<td>9-(4-(3-chloropropoxy)phenyl)-9-(4-(3-(prop-2-ynyloxy)propoxy)phenyl)-9H-fluorene</td>
</tr>
<tr>
<td>21</td>
<td><img src="image2.png" alt="Structure Image" /></td>
<td>9-(4-(3-chloropropoxy)phenyl)-9-(4-(prop-2ynyloxy)phenyl)-9H-fluorene</td>
</tr>
<tr>
<td>22</td>
<td><img src="image3.png" alt="Structure Image" /></td>
<td>2-(2-(2-(4-(9-(4-(3-chloropropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)ethoxy)ethoxy)ethanol</td>
</tr>
<tr>
<td>23</td>
<td><img src="image4.png" alt="Structure Image" /></td>
<td>2-(2-(2-(4-(9-(4-(3-chloropropoxy)phenyl)-9H-fluoren-9-yl)phenoxy)ethoxy)ethoxy)acetic acid</td>
</tr>
<tr>
<td>No.</td>
<td>Structure</td>
<td>Name</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
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</tr>
<tr>
<td>24</td>
<td><img src="image" alt="Structure 24" /></td>
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<td>25</td>
<td><img src="image" alt="Structure 25" /></td>
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</tr>
<tr>
<td>26</td>
<td><img src="image" alt="Structure 26" /></td>
<td>N/A</td>
</tr>
</tbody>
</table>
Prodrugs are also included within the scope of the present disclosure. For example, in one embodiment the hydrogen atom of one or more hydroxyl groups of any of the compounds of Formula I may be replaced with a moiety from Table 1. A non-limiting example of such prodrugs include glycine esters and salts thereof of compounds of Formula I as shown below.

In some embodiments, the compounds as described herein or acceptable salts thereof above may be used for systemic treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty and age-related macular degeneration. In some embodiments, the compounds as described herein or acceptable salts thereof above may be used in the preparation of a medicament or a composition for systemic treatment of an indication described herein. In some embodiments, methods of systemically treating any of the indications described herein are also provided. Some aspects of this invention, make use of compositions comprising a compound described herein and a pharmaceutically acceptable excipients or carrier. In some embodiments, the prostate cancer is
androgen-independent prostate cancer (also referred to as hormone refractory, castration
resistant, androgen deprivation resistant, androgen ablation resistant, androgen
depletion-independent, castration-recurrent, anti-androgen-recurrent). In some
embodiments the prostate cancer is androgen-dependent or androgen-sensitive.
Methods of treating any of the indications described herein are also provided. Such
methods may include administering a compound as described herein or a composition
of a compound as described herein, or an effective amount of a compound as described
herein or composition of a compound as described herein to a subject in need thereof.

Compounds as described herein may be in the free form or in the form of
a salt thereof. In some embodiments, compounds as described herein may 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 may
be, for example, formed as a pharmaceutically acceptable salt. Compounds containing
one or more basic functional groups may be capable of forming a pharmaceutically
acceptable salt with, for example, a pharmaceutically acceptable organic or inorganic
acid. Pharmaceutically acceptable salts may be derived from, for example, and without
limitation, acetic acid, adipic acid, alginic acid, aspartic acid, ascorbic acid, benzoic
acid, benzenesulfonylic 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 may 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 may 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-dimethylaminooctanol, 2-diethylaminooctanol, 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-ephenamine, N,N'-dibenzylylhexylenediamine or polyamine resins. In some embodiments, compounds as described herein may contain both acidic and basic groups and may be in the form of inner salts or zwitterions, for example, and without limitation, betaines. Salts as described herein may 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 may occur in situ during isolation and purification of the compounds or preparation of salts may 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 may 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 may be, for example, and without limitation, a pharmaceutically

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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 may include crystalline and amorphous forms, for example, polymorphs, pseudopolymorphs, conformational polymorphs, amorphous forms, or a combination thereof. Polymorphs include different crystal packing 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 may cause a single crystal form to dominate.

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 the sake of convenience.

In some embodiments, pharmaceutical compositions in accordance with this invention may 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 may 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 may 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 may be administered in a tablet, capsule or dissolved in liquid form. The tablet or capsule may 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 may 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 may, 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 may 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 may contain excipients, for example, lactose, or may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or may be oily solutions for administration in the form of nasal drops, or as a gel.

Compounds or pharmaceutical compositions in accordance with this invention or for use in this invention may be administered by means of a medical device or appliance such as an implant, graft, prosthesis, stent, etc. Also, implants may 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.

An "effective amount" of a pharmaceutical composition according to the invention includes 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 may 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 may 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 an androgen-independent form. Typically, a prophylactic dose is used in subjects prior to or at an earlier stage of disease, so that a prophylactically effective amount may be less than a therapeutically effective amount.

It is to be noted that dosage values may vary with the severity of the condition to be alleviated. For any particular subject, specific dosage regimens may be adjusted over time according to the individual need and the professional judgement 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 may be selected by medical practitioners. The amount of active compound(s) in the composition may vary according to factors such as the disease state, age, sex, and weight of the subject. Dosage regimens may be adjusted to provide the optimum therapeutic response. For example, a single bolus may be administered, several divided doses may be administered over time or the dose may be proportionally reduced or increased as indicated by the exigencies of the therapeutic situation. It may be advantageous to formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage.

In some embodiments, compounds and all different forms thereof as described herein may 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 and bulbar muscular atrophy, and age-related macular degeneration. For example, compounds and all their different forms as described herein may 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.

With respect to combination therapies, one embodiment of the present disclosure provides a combination of any one or more of a compound of Formula I with one or more currently-used or experimental pharmacological therapies which are or may be utilized to treat any of the above disease states (e.g., androgen-independent
prostate cancer or Kennedy's disease). Methods, uses and pharmaceutical compositions comprising the above combination are also provided. Combination therapies for such indications are disclosed in co-pending U.S. Provisional Application No. 61/384,628, which is hereby incorporated by reference in its entirety.

Surprisingly, it has been found that the disclosed compounds, which interfere with the AR principally through binding to the N-terminus of the AR, demonstrate beneficial synergistic therapeutic effects when used in concert with existing approved and in-development agents. That is, the biological impact of using the agents in concert with one another produces a biological and therapeutic effect which is greater than the simple additive effect of each of them separately.

Accordingly, one embodiment comprises the use of the disclosed compounds in 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 therapies which interfere with the biological production or function of androgen (hereinafter, the "Other Therapeutic Agents"). 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 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 and one or more of the Other 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 Other Therapeutic Agents for treatment of the above disease states also includes dosing by any dosing method including without limitation, intravenous 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 Other Therapeutic Agent may be administered to the patient before administration of the compound(s) of Formula I. In another embodiment, the compound(s) of Formula I may be co-administered with one or more of the Other Therapeutic Agents. In yet another aspect, the one or more Other Therapeutic Agent may be administered to the patient after administration of the compound(s) of Formula I.

It is fully within the scope of the disclosure that the ratio of the doses of compound(s) of Formula I to that of the one or more Other Therapeutic Agents may or may not equal to one and may be varied accordingly to achieve the optimal therapeutic benefit.

For greater clarity the compound(s) of Formula I that are combined with the one or more Other Therapeutic Agents for improved treatment of the above disease states may comprise, but are not limited to any compound having a structure of Formula 1, including those compounds shown in Table 2.

The Other 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 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 MDV3100 (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 AR LBD and is currently in development as a treatment for prostate cancer; the chemical entity known as TOK 001 and related compounds which appears to be a blocker of the AR LBD, and a CYP17 lyase inhibitor, and also appears to decrease overall androgen receptor levels in prostate cancer cells. TOK 001 is currently in development as a treatment for prostate cancer; the chemical entity known as ARN-509 and related compounds which appears to be a blocker of the 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) 2,3,4,7,8,9,10,1 1,12,13,14,15-dodecahydro-1H-cyclopenta[a]phenanthren-3-ol), and related molecules, which appears to block the production of androgen and is currently in development for the treatment of 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 AR LBD and which is currently used to treat prostate cancer, the chemical entity known as nilutamide (5,5-dimethyl-3-[4-nitro-3-(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 AR LBD and which is currently used to treat prostate cancer, the chemical entities known as cyproterone acetate (6-chloro-1β,2β-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 AR LBD and which is currently used to treat prostate cancer, the chemical entity known as docetaxel (Taxotere; 1β,10β-trihydroxy-9-oxo\(\text{\textsuperscript{20}}\)-epoxytax-1\(\text{\textsuperscript{1}}\)-ene-2α,4,13α-triy1 4-acetate 2-benzoate 13-((2R,3S)-3-[(tert-butoxycarbonyl)amino]-2-hydroxy-3-phenylpropanoate)) and related compounds, which 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 may be used to treat prostate cancer, the chemical entity known as OSU-HDA\(\text{\textsuperscript{42}}\) ((S)-(H-);)-N-hydroxy-4-(3-methyl-2-phenylbutyl)tryramino)-benzazirside), 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 \(\alpha\)\(\text{\textbeta}\)3 to prevent angiogenesis, and which may be used to treat prostate cancer, the chemical entity known as sunitimib (N-(2-diethylaminoethyl)-5-[(Z)-(5-fluoro-2-oxo-IH-indol-3-ylidene)methyl]-2,4-dimethyl-IH-pyrrole-3-carboxamide) and related compounds, which appears to inhibit multiple receptor tyrosine kinases (RTKs) and may be used for treatment of prostate cancer, the chemical entity known as ZD-4054 (N-(3-Methoxy-5-methyipyrazin-2-yl)-2-[4-(1,3,4-oxadiazol-2-yl)phenyl]pyridin-3-sulphonamid) and related compounds, which appears to block the edta receptor and which may be used for treatment of prostate cancer, the chemical entity known as VN/124-1 (3β-Hydroxy-17-(IH-benzimidazol-1-yI)andros(a-5,16-diene), and relaxed compounds which appears to block the production
of androgen (via inhibition of -hydroxylase/17,20 lyase) and is currently in
development for the 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 (Ipilimunab), a fully human monoclonal antibody that binds to and blocks the activity of CTLA-4 which is currently in development as an 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 Oil 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 (Proscar, Propecia; N-(1,1-dimethylethyl)-3-oxo-(5a, 17P)-4-azaandrost-1-ene-17-carboxamide), and related compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone, and may be used to treat prostate cancer; the chemical entity known as dutasteride (Avodart; 5a, 17β)-N-{2,5 bis(trifluoromethyl) phenyl} -3-oxo-4-azaandrost-1 -ene-17-carboxamide) and related molecules, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone, and may be used in the treatment of prostate cancer; the chemical entity known as turosteride ((4aR,4bS,6aS,7S,9aS,9bS,1 1aR)-1,4a,6a-trimethyl-2-oxo-N-(propan-2-yl)-N-(propan-2 ylcarbamoyl)hexadecahydro-lH-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 may 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 may 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 may 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 may be used for the treatment of prostate cancer; the chemical entity known as SKF 105,1 11, and related
compounds, which appears to be a 5-alpha reductase inhibitor that reduces levels of dihydrotestosterone and may be used for treatment of prostate cancer.

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 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 however, such as in severe disease conditions, it may be necessary to administer substantial excesses of the compositions. Some compounds of this invention may be toxic at some concentrations. Titration studies may be used to determine toxic and non-toxic concentrations. Toxicity may be evaluated by examining a particular compound's or composition's specificity across cell lines using PC3 cells as a negative control that do not express AR. Animal studies may 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 since antiandrogens and androgen insensitivity syndrome are not fatal.

Compounds as described herein may be administered to a subject. As used herein, a "subject" may be a human, non-human primate, rat, mouse, cow, horse, pig, sheep, goat, dog, cat, etc. The subject may be suspected of having or at risk for having a cancer, such as prostate cancer, breast cancer, ovarian cancer 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, or age-related macular degeneration. Diagnostic methods for various cancers, such as prostate cancer, breast cancer, ovarian cancer or endometrial cancer, and diagnostic methods for acne, hirsutism, alopecia, benign prostatic hyperplasia, ovarian cysts, polycystic ovary disease, precocious puberty, or age-related macular degeneration and the clinical delineation of cancer, such as prostate cancer, breast cancer, ovarian cancer or endometrial cancer, diagnoses and the clinical delineation of acne, hirsutism, alopecia, benign prostatic hyperplasia, ovarian cysts, polycystic ovary disease, precocious puberty, or age-related macular degeneration are known to those of ordinary skill in the art.

Compounds described herein may be used for treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian
cancer, endometrial cancer, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty and age-related macular degeneration. Compounds described herein may be used for treatment of prostate cancer. Compounds described herein may be used for treatment of androgen-independent prostate cancer. Compounds described herein may be used for treatment of androgen-dependent prostate cancer. Compounds described herein may be used for preparation of a medicament for treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty and age-related macular degeneration. Compounds described herein may be used for the preparation of a medicament for treatment of prostate cancer. Compounds described herein may be used for the preparation of a medicament for treatment of androgen-independent prostate cancer. Compounds described herein may be used for the preparation of a medicament for treatment of androgen-dependent prostate cancer. Compounds described herein may be used in a method for treatment of at least one indication selected from the group consisting of: prostate cancer, breast cancer, ovarian cancer, endometrial cancer, hair loss, acne, hirsutism, ovarian cysts, polycystic ovary disease, precocious puberty and age-related macular degeneration. The method may comprise administering to a subject in need thereof an effective amount of a compound described herein. Compounds described herein may be used in a method of treatment of prostate cancer, the method comprising administering to a subject in need thereof an effective amount of a compound described herein. Compounds described herein may be used in a method of treatment of androgen-independent prostate cancer, the method comprising administering to a subject in need thereof an effective amount of a compound described herein. Compounds described herein may be used in a method of treatment of androgen-dependent prostate cancer, the method comprising administering to a subject in need thereof an effective amount of a compound described herein.

Compounds described herein may 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 AR refers to transactivation of the 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 may inhibit both FSK and androgen (e.g. R1881) induction of ARE-luciferase (ARE-luc). Such compounds may block a mechanism that is common to both ligand-dependent and ligand-independent activation of the AR. This could involve any step in activation of the 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 may inhibit R1881 only and may 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 may be used to treat such conditions. Some compounds and compositions of this invention may only inhibit FSK induction and may be specific inhibitors to ligand-independent activation of the AR. These compounds and compositions may interfere with the cascade of events that normally occur with FSK and/or PKA activity or any downstream effects that may play a role on the AR (e.g. FSK increases MAPK activity which has a potent effect on AR activity). Examples may include an inhibitor of cAMP and or PKA or other kinases. Some compounds and compositions of this invention may induce basal levels of activity of the AR (no androgen or stimulation of the PKA pathway). Some compounds and compositions of this invention may increase induction by R1881 or FSK. Such compounds and compositions may stimulate transcription or transactivation of the AR. Some compounds and compositions of this invention may inhibit activity of the androgen receptor. Interleukin-6 (IL-6) also causes ligand-independent activation of the AR in LNCaP cells and can be used in addition to FSK.

Compounds for use in the present invention may 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 principles. For example, Auzou et al 1974 European Journal of Medicinal Chemistry 9(5), 548-554 describes suitable synthetic procedures that may be considered and suitably adapted for preparing compounds of any one of the

For example, compounds of the present invention which contain an ether moiety may be obtained with reference to the following General Reaction Scheme I:

General Reaction Scheme I

wherein R₇-OH represents an alcohol and M, L, and n are as defined anywhere herein. Bismuth triflate may be added in portions to a solution of racemic derivative A in an alcohol R₇-OH over the course of the reaction. The mixture may be stirred under suitable conditions (for example, rt for 24 h). The resulting suspension may be quenched by a suitable reagent (for example, by addition of sodium bicarbonate), extracted (for example, with ethyl acetate), dried (for example, over anhydrous magnesium sulphate), and concentrated (for example, under vacuum). The resulting residue may be purified by a suitable method (for example, flash column chromatography on silica gel - eluent: 90% hexane in ethyl acetate) to provide B. A person of skill in the art will understand that the above General Scheme I may be suitably adapted to prepare compounds of the present invention which contain any ether moiety, including polyethers (or an alcohol in the case where R₇ is H), for example a
propargyl ether moiety, for example, based on the following General Reaction Scheme II:

General Reaction Scheme II

wherein M, L, and n are as defined anywhere herein.

The General Reaction Scheme I may be suitably adapted to prepare compounds of the present invention which contain an isopropyl ether moiety, for example, based on the following General Reaction Scheme III:

General Reaction Scheme III

wherein M, L, and n are as defined anywhere herein.

The General Reaction Scheme I may be suitably adapted to prepare compounds of the present invention which contain an n-butyl ether moiety, for example, based on the following General Reaction Scheme IV:
General Reaction Scheme IV

wherein M, L, and n are as defined anywhere herein.

The General Reaction Scheme I may be suitably adapted to prepare compounds of the present invention which contain a cyclohexyl ether moiety, for example, based on the following General Reaction Scheme V:

Reaction Scheme V

wherein M, L, and n are as defined anywhere herein.

The General Reaction Scheme I may be suitably adapted to prepare compounds of the present invention which contain a chloro moiety, for example, based on the following General Reaction Scheme VI:
The General Reaction Scheme I may be suitably adapted to prepare compounds of the present invention which contain an ester moiety, for example, based on the following General Reaction Scheme VII:

For ease of illustration, the above General Reaction Schemes depict compounds of Formula I wherein each of $Z^1$, $Z^2$ $Z^3$ and $Z^4$ are C-H. However, one skilled in the art will recognize that variations of the above procedures (e.g., starting with variously substituted compounds or performing any number of aromatic substitution reactions) can yield compounds of structure I wherein each of $Z^1$, $Z^2$ $Z^3$ or $Z^4$ are optionally other than C-H. General methodologies for chemical preparation of compounds of Formula I are described in the following non-limiting exemplary schemes.

Various alternative embodiments and examples of the invention are described herein. These embodiments and examples are illustrative and should not be construed as limiting the scope of the invention.
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 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 aluminium 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 molybdatophosphoric acid, 17.5 g sulphuric acid) followed by heating (~1 min) with a heating gun (~250 °C). Organic solutions were concentrated on Buchi R-114 rotary 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 Sepaks™ were purchased from waters, Inc. Thin-layer chromatography plates were Kieselgel 60F254. All synthetic reagents were purchased from Sigma Aldrich and Fisher Scientific Canada.

Proton nuclear magnetic resonance (1H NMR) spectra were recorded at 25 °C using a Bruker 400 with inverse probe and Bruker 400 spectrometers, are reported in parts per million on the δ scale, and are referenced from the residual protium in the NMR solvent (DMSO-d₆: δ 2.50 (DMSO-d₆), CDC1₃: δ 7.24 (CHC1₃)). Carbon-13 nuclear magnetic resonance (13C NMR) spectra were recorded with a Bruker 400 spectrometer, are reported in parts per million on the δ scale, and are referenced from the carbon resonances of the solvent (DMSO-d₆: δ 39.51, CDC1₃: δ 77.00). Spectral features are tabulated in the following order: chemical shift (δ, ppm); multiplicity (s = singlet, d = doublet, t = triplet, m = multiplet, br = broad); coupling constant (J, Hz, number of protons).

LNCaP cells were employed initially for all experiments because they are well-differentiated human prostate cancer cells in which ligand-independent activation of the AR by FSK has been characterized (Nazareth et al 1996 J. Biol. Chem.
LNCaP cells express endogenous AR and secrete prostate-specific antigen (PSA) (Horoszewicz et al 1983 Cancer Res. 43, 1809-1818). LNCaP cells can be grown either as monolayers in cell culture or as tumors in the well-characterized xenograft model that progresses to androgen independence in castrated hosts (Sato et al 1996 J. Steroid Biochem. Mol. Biol. 58, 139-146; Gleave et al 1991 Cancer Res. 51, 3753-3761; Sato et al 1997 Cancer Res. 57, 1584-1589; and Sadar et al 2002 Mol. Cancer Ther. 1(8), 629-637).

R1881 was employed since it is stable and avoids problems associated with the labile physiological ligand dihydrotestosterone (DHT). Reporter specificity may be determined using several alternative reporter gene constructs. Some well characterized ARE-driven reporter gene constructs that have been used extensively are the PSA (6.1 kb) enhance/promoter which contains several AREs and is highly inducible by androgens as well as by FSK (Ueda et al 2002 A J. Biol. Chem. 277, 7076-7085) and the ARR3-thymidine kinase (tk)-luciferase, which is an artificial reporter construct that contains three tandem repeats of the rat probasin ARE1 and ARE2 regions upstream of a luciferase reporter (Snoek et al 1996 J. Steroid Biochem. Mol. Biol. 59, 243-250).
EXAMPLE 1

SYNTHESIS OF 3,3’-(4,4’-(9H-FLUORENE-9,9-DIYL)BIS (4,1-PHENYLENE))BIS(OXY)BIS (1-
CHLOROPROPAN-2-OL) (1)

To a solution of racemic derivative 2 (622 mg, 1.34 mmol, 1 equiv) in acetonitrile (5 mL) was added CeCl₃·7H₂O (1250 mg, 3.36 mmol, 2.5 equiv) and the mixture was refluxed for 19 h. The resulting white paste was filtered and washed with ethyl acetate and the clear suspension was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 2% ethyl acetate in dichloromethane to 2% methanol in dichloromethane) to provide 1 (585 mg, 81%) as a white foamy solid.

1H NMR (400 MHz, DMSO-d₆): δ 7.90 (d, J = 7.6, 2H), 7.38 (t, J = 8.0, 4H), 7.30 (t, J = 8.0, 2H), 7.01 (d, J = 8.8, 4H), 6.83 (d, J = 8.8, 4H), 5.51 (d, J = 5.2, 2H), 4.01-3.97 (m, 2H), 3.91 (d, J = 5.2, 4H), 3.73-3.69 (dd, J = 11.2, 4.8, 2H), 3.65-3.61 (dd, J = 11.2, 5.6, 2H); 13C NMR (100 MHz, DMSO-d₆): δ 157.1, 151.2, 139.3, 137.8, 128.7, 127.8, 127.5, 125.9, 120.5, 114.2, 68.8, 68.5, 63.6, 46.7; HRMS (ESI) (m/z): calc’d for C₃₁H₂₆GaCl₂ [M+Na]⁺: 557.1262, found: 557.1254.
EXAMPLE 2
SYNTHESIS OF 2,2’-(4,4’-(9H-FLUORENE-9,9-DIYL)BIS(4,1-PHENYLENE))BIS(OXY)BIS(METHYLENE)DIOXIRANE 2

Sodium hydride (60% dispersion in mineral oil, 502 mg, 12.5 mmol, 2.2 equiv) was added slowly to a stirred solution of 4,4’-(9-Fluorenylidene)diphenol (2000 mg, 5.70 mmol, 1 equiv, available from Aldrich Chemicals, catalog # 3236-71-3) in anhydrous dimethyl formamide (20 mL), at room temperature, and the contents were stirred under an atmosphere of argon for 20 min. Racemic epichlorohydrin (1340 μL, 17.1 mmol, 3.0 equiv) was added via syringe and the mixture was allowed to react at room temperature for 162 h. Then, the reaction was quenched by the addition of a saturated solution of ammonium chloride (10 mL), and the mixture was extracted with ethyl acetate (3 x 20 mL). The organic layer was washed with deionized water (20 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 10% ethyl acetate in hexane to 20% ethyl acetate to 100% dichloromethane) to provide 2 (2.53 g, 96%) as a white solid. $^1$H NMR (400 MHz, DMSO-de): δ 7.90 (d, $J = 7.6$, 2H), 7.38 (t, $J = 8.4$, 4H), 7.30 (t, $J = 7.2$, 2H), 7.02 (d, $J = 9.2$, 4H), 6.84 (d, $J = 8.8$, 4H), 4.26-4.22 (dd, $J = 11.6$, 2.4, 2H), 3.79-3.75 (dd, $J = 11.6$, 6.4, 2H), 3.29-3.25 (m, 2H), 2.81 (t, $J = 4.4$, 2H), 2.67-2.65 (dd, $J = 5.2$, 2.8, 2H); $^{13}$C NMR (100 MHz, DMSO-d$_6$): δ 156.9, 151.1, 139.3, 137.9, 128.7, 127.8, 127.5, 125.9, 120.5, 114.3, 68.9, 63.6, 49.6, 43.7; HRMS (ESI) (m/z): calc’d for C$_{31}$H$_{36}$O$_4$Na [M+Na]$^+$: 485.1729, found: 485.1741.
EXAMPLE 3

SYNTHESIS OF 3-(4-(9-(4-(3-CHLORO-2-HYDROXYPROPOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPANE-1,2-DIOL (3)

For a solution of racemic derivative 2 (511 mg, 1.30 mmol, 1 equiv) in a mixture of acetonitrile (4 mL) and water (2 mL) was added CeCl₃·7H₂O (206 mg, 0.65 mmol, 1/2 equiv) and Bismuth(III) trifluoromethanesulfonate (73 mg, 0.13 mmol, 1/10 equiv) and the mixture was heated at 80-90 °C for 51 h. Then, the reaction was quenched by the addition of a saturated solution of sodium bicarbonate (2 mL), and the mixture was extracted with ethyl acetate (3 x 10 mL). The organic layer was washed with deionized water (15 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 60% ethyl acetate in hexane to 80% ethyl acetate) to provide 3 (64 mg, 11%) as a white solid. Note: The same reaction yielded 231 mg (39%) of 1 as a white foam. 

¹H NMR (400 MHz, DMSO-de): δ 7.90 (d, J = 7.6, 2H), 7.38 (t, J = 7.6, 4H), 7.30 (t, J = 7.6, 2H), 7.02-6.99 (dd, J = 8.8, 3.6, 4H), 6.82 (d, J = 8.8, 4H), 5.51 (d, J = 5.2, 1H), 4.89 (d, J = 4.0, 1H), 4.62 (m, 1H), 4.01-3.97 (m, 1H), 3.93-3.90 (m, 3H), 3.80-3.69 (m, 3H), 3.65-3.61 (dd, J = 11.2, 5.2, 1H), 3.40 (m, 2H); ¹³C NMR (100 MHz, DMSO-d₆): δ 157.5, 157.0, 151.2, 139.3, 137.9, 137.4, 128.7, 128.7, 127.8, 127.5, 125.9, 120.5, 114.2, 114.2, 69.9, 69.5, 68.8, 68.5, 63.6, 62.7, 46.7; HRMS (ESI) (m/z): calc'd for C₃iH₂₈O₅NaCl [M+Na]⁺: 539.1601, found: 539.1600.
EXAMPLE 4

**SYNTHESIS OF 3-(4-(9-(4-(2-HYDROXY-3-(PROP-2-YNYLOXY)PROPOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPANE-1,2-DIOL (4)**

To a solution of racemic derivative 2 (200 mg, 0.43 mmol, 1 equiv) in a mixture of acetonitrile (1 mL), water (1 mL), and propargyl alcohol (1 mL) was added Bismuth(III) trifluoromethanesulfonate (28 mg, 0.043 mmol, 1/10 equiv) and the mixture was stirred at room temperature for 44 h. Then, the reaction was quenched by the addition of a saturated solution of sodium bicarbonate (1 mL), and the mixture was extracted with ethyl acetate (3 x 5 mL). The organic layer was washed with deionized water (5 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: dichloromethane to 5% methanol in dichloromethane) to provide 4 (40 mg, 17%) as a white solid. Note: The same reaction yielded 128 mg (59%) of 5 as white solid (Example 5).

**$^1$H NMR (400 MHz, DMSO-d$_6$):** δ 7.90 (d, J = 7.6, 2H), 7.40-7.36 (dd, J = 9.6, 7.6, 4H), 7.30 (t, J = 7.6, 2H), 7.01 (d, J = 8.8, 4H), 6.81 (d, J = 8.8, 4H), 5.14 (d, J = 4.8, 1H), 4.90 (d, J = 4.8, 1H), 4.62 (t, J = 5.6, 1H), 4.15 (d, J = 2.4, 2H), 3.94-3.87 (m, 3H), 3.84-3.73 (m, 3H), 3.53-3.46 (m, 2H), 3.43-3.40 (m, 3H); **$^{13}$C NMR (100 MHz, DMSO-d$_6$):** δ 157.5, 157.3, 151.2, 139.3, 137.7, 137.5, 128.7, 127.8, 127.5, 125.9, 120.4, 114.2, 114.2, 80.3, 77.2, 70.9, 69.9, 69.5, 67.8, 63.6, 62.7, 57.9; **HRMS (ESI) (m/z):** calc'd for C$_{34}$H$_{32}$O$_5$Na [M+Na]$^+$: 559.2097, found: 559.2086.
EXAMPLE 5
SYNTHESIS OF 3,3’-(4,4’-(9H-FLUORENE-9,9-DIYL)BIS(4,1-PHENYLENE))BIS(OXY)DIPROPANE-1,2-DIOL (5)

To a solution of racemic derivative 2 (200 mg, 0.43 mmol, 1 equiv) in a mixture of acetonitrile (1 mL), water (1 mL), and propargyl alcohol (1 mL) was added Bismuth(III) trifluoromethanesulfonate (28 mg, 0.043 mmol, 1/10 equiv) and the mixture was stirred at room temperature for 44 h. Then, the reaction was quenched by the addition of a saturated solution of sodium bicarbonate (1 mL), and the mixture was extracted with ethyl acetate (3 x 5 mL). The organic layer was washed with deionized water (5 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 5% methanol in dichloromethane to 30% methanol in dichloromethane) to provide 5 (128 mg, 59%) as a white solid. 

$^1$H NMR (400 MHz, DMSO-$d_6$): δ 7.90 (d, $J = 7.6$, 2H), 7.40-7.36 (m, 4H), 7.32-7.28 (m, 2H), 7.01-6.98 (m, 4H), 6.83-6.80 (m, 4H), 4.89 (d, $J = 4.8$, 2H), 4.62 (t, $J = 5.6$, 2H), 3.94-3.90 (dd, $J = 9.6$, 4.0, 2H), 3.81-3.72 (m, 4H), 3.40 (t, $J = 5.6$, 4H); $^1$C NMR (100 MHz, DMSO-$d_6$): δ 157.4, 151.3, 139.3, 137.5, 128.7, 127.8, 127.4, 125.9, 120.4, 114.2, 69.9, 69.5, 63.6, 62.7; HRMS (ESI) (m/z): calc’d for C$_{38}$H$_{36}$O$_6$Na [M+Na]$^+$: 521.1940, found: 521.1949.
EXAMPLE 6
SYNTHESIS OF (2R,2R)-2,2-(4,4-(9H-FLUORENE-9,9-DIYL)BIS(4,1-PHENYLENE))BIS(OXY)BIS(METHYLENE)DIOXIRANE (6)

Sodium hydride (60% dispersion in mineral oil, 528 mg, 13.19 mmol, 2.5 equiv) was added slowly to a stirred solution of 4,4’-(9-Fluorenylidene)diphenol (1850 mg, 5.27 mmol, 1 equiv) in anhydrous dimethyl formamide (20 mL), at room temperature, and the contents were stirred under an atmosphere of argon for 20 min. A solution of (2i?)-(−)-glycidyl tosylate 98% (3010 mg, 13.19 mmol, 2.5 equiv) in anhydrous dimethyl formamide (5 mL) was added via syringe and the mixture was allowed to react at room temperature for 94 h. Then, the reaction was quenched by the addition of a saturated solution of ammonium chloride (10 mL), and the mixture was extracted with ethyl acetate (3 x 20 mL). The organic layer was washed with deionized water (20 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 10% to 20% ethyl acetate in hexane) to provide 6 (1.91 g, 78%) as a white solid. $^1$H NMR (400 MHz, DMSO-d$_6$): δ 7.90 (d, J = 7.6, 2H), 7.38 (t, J = 8.0, 4H), 7.30 (t, J = 7.2, 2H), 7.02 (d, J = 8.8, 4H), 6.84 (d, J = 8.8, 4H), 4.26-4.22 (dd, J = 11.6, 2.8, 2H), 3.79-3.74 (dd, J = 11.2, 6.4, 2H), 3.29-3.26 (m, 2H), 2.81 (t, J = 4.8, 2H), 2.67-2.65 (dd, J = 5.2, 2.8, 2H); $^{13}$C NMR (100 MHz, DMSO-4): δ 156.9, 151.1, 139.3, 137.9, 128.7, 127.5, 125.9, 120.5, 114.3, 68.9, 63.6, 49.6, 43.7; HRMS (ESI) (m/z): calc'd for C$_{31}$H$_{26}$G$_4$Na [M+Na]$^+$: 485.1729, found: 485.1734.
EXAMPLE 7
SYNTHESIS OF (S)-1-CHLORO-3-(4-((9-((R)-OXIRAN-2-YLMETHOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPAN-2-OL (7)

To a solution of derivative 6 (1910 mg, 4.14 mmol, 1 equiv) in acetonitrile (10 mL) was added CeCl₃·7 H₂O (770 mg, 2.06 mmol, 1/2 equiv) and the mixture was refluxed for 3.5 h. The resulting white paste was filtered and washed with ethyl acetate and the clear suspension was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: dichloromethane to 2% ethyl acetate in dichloromethane to 20% ethyl acetate in dichloromethane) to provide 7 (382 mg, 19%) as a transparent foam. Note: The same reaction yielded 77 mg of 8 as a white solid together with unreacted starting material.

¹H NMR (400 MHz, DMSO-d₆): δ 7.91 (d, J = 7.2, 2H), 7.38 (t, J = 7.6, 4H), 7.30 (t, J = 7.6, 2H), 7.03-7.00 (dd, J = 8.8, 2.0, 4H), 6.85-6.82 (dd, J = 8.8, 2.8, 4H), 5.51 (d, J = 5.2, 1H), 4.26-4.22 (dd, J = 11.2, 2.8, 1H), 4.00-3.97 (m, 1H), 3.90 (d, J = 5.6, 2H), 3.79-3.69 (m, 2H), 3.65-3.61 (dd, J = 10.8, 5.2, 1H), 3.30-3.26 (m, 1H), 2.81 (t, J = 4.8, 1H), 2.68-2.66 (dd, J = 5.2, 2.8, 1H); ¹³C NMR (100 MHz, DMSO-d₆): δ 157.0, 156.9, 151.1, 139.3, 137.9, 137.8, 128.7, 127.8, 127.5, 125.9, 120.5, 114.3, 68.9, 68.8, 68.5, 63.6, 49.6, 46.7, 43.7; HRMS (ESI) (m/z): calc'd for C₃iH₂₇O₄NaCl [M+Na]⁺: 521.1496, found: 521.1508.

EXAMPLE 8
SYNTHESIS OF (2S,2S)-3,3-(4,4-((9H-FLUORENE-9,9-DIYL)BIS(4,1-PHENYLENE))BIS(OXY)BIS(1-CHLOROPROPAN-2-OL) (8)
To a solution of derivative 6 (1910 mg, 4.14 mmol, 1 equiv) in acetonitrile (10 mL) was added CeCl₃·7H₂O (770 mg, 2.06 mmol, 1/2 equiv) and the mixture was refluxed for 3.5 h. The resulting white paste was filtered and washed with ethyl acetate and the clear suspension was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: dichloromethane to 2% ethyl acetate in dichloromethane to 20% ethyl acetate in dichloromethane) to provide 8 (77 mg, 4%) as a white solid. Note: The same method applied to obtain analogue 1 can be applied for the synthesis of derivative 8 by using 6 as starting material.

**EXAMPLE 9**

**SYNTHESIS OF (S)-1-CHLORO-3-(4-(4-(R)-2-HYDROXY-3-(PROP-2-ynyloxy)propoxy)phenyl)-9H-fluoren-9-ylphenoxy)propan-2-ol (9)**

\[ \text{CeCl}_3 \times 7 \text{H}_2 \text{O} \]

\[ \text{MeCN, reflux} \]
To a mixture of derivative 7 (38 mg, 0.076 mmol, 1 equiv) and propargyl alcohol (1 mL) was added Erbium(III) trifluoromethanesulfonate (10 mg, 0.015 mmol, 1/5 equiv) and the brown solution was stirred at room temperature for 16 h. Water was added (1 mL) and the reaction was extracted with ethyl acetate (3 x 5 mL). The organic layer was washed with deionized water (5 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel Sep pak (5g) (eluent: dichloromethane to 5% ethyl acetate in dichloromethane) to provide 9 (33 mg, 79%) as a white solid. 

\[ \text{H NMR (400 MHz, DMSO-d6): } \delta 7.90 (d, J = 7.6, 2H), 7.40-7.36 (m, 4H), 7.32-7.28 (m, 2H), 7.03-7.00 (m, 4H), 6.84-6.80 (m, 4H), 5.52 (d, J = 5.2, 1H), 5.13 (d, J = 5.2, 1H), 4.15 (d, J = 2.4, 2H), 4.01-3.98 (m, 1H), 3.92-3.86 (m, 4H), 3.84-3.81 (m, 1H), 3.74-3.70 (dd, J = 11.2, 4.8, 1H), 3.66-3.61 (dd, J = 11.2, 5.2, 1H), 3.52-3.45 (m, 2H), 3.42 (t, J = 2.4, 1H); \]

\[ \text{13C NMR (100 MHz, DMSO-d6): } \delta 157.3, 157.0, 151.2, 139.3, 137.9, 137.6, 128.7, 128.7, 127.8, 127.5, 125.9, 120.5, 114.2, 80.3, 77.2, 70.9, 69.4, 68.8, 68.5, 67.8, 63.6, 57.9, 46.7; \]

\[ \text{HRMS (ESI) (m/z): calc'd for } C_{34}H_{30}i0_{3}\text{NaCl [M+Na]^+: } 577.1758, \text{ found: } 577.1747. \]

**EXAMPLE 10**

**SYNTHESIS OF (S)-1-CHLORO-3-(4-(4-((R)-2-HYDROXY-3-ISOPROPOXYPROPOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPAN-2-OL (10)**

To a solution of derivative 7 (21 mg, 0.042 mmol, 1 equiv) in isopropyl alcohol (1 mL) was added erbium(III) trifluoromethanesulfonate (6 mg, 0.008 mmol, 1/5 equiv) and the mixture was stirred at room temperature for 16 h. After the crude was concentrated under reduced pressure, the resulting residue was purified by flash column chromatography on silica gel Sep pak (5g) (eluent: dichloromethane to 5% ethyl acetate in dichloromethane) to provide 10 (23 mg, 97%) as a white foam. 

\[ \text{H} \]
NMR (400 MHz, DMSO-d$_6$): $\delta$ 7.90 (d, $J = 7.6$, 2H), 7.38 (t, $J = 7.6$, 4H), 7.30 (t, $J = 7.6$, 2H), 7.02-6.99 (dd, $J = 8.8$, 2.4, 4H), 6.84-6.80 (dd, $J = 8.4$, 7.2, 4H), 5.50 (d, $J = 5.2$, 1H), 4.97 (d, $J = 4.8$, 1H), 4.01-3.97 (m, 1H), 3.92-3.87 (m, 3H), 3.84-3.79 (m, 2H), 3.73-3.70 (dd, $J = 11.2$, 4.4, 1H), 3.65-3.61 (dd, $J = 10.8$, 7.2, 1H), 3.55-3.49 (m, 1H), 3.40-3.36 (m, 2H), 1.05 (d, $J = 6.0$, 6H); $^1$C NMR (100 MHz, DMSO-d$_6$): $\delta$ 157.4, 157.0, 151.2, 139.3, 137.9, 137.5, 128.7, 128.7, 127.5, 125.9, 120.4, 114.2, 71.1, 69.6, 69.0, 68.8, 68.5, 68.2, 63.6, 46.7, 22.0; HRMS (ESI) (m/z): calc’d for C$_{34}$H$_{35}$O$_5$NaCl [M+Na$^+$]: 581.2071, found: 581.2081

**EXAMPLE 11**

**SYNTHESIS OF 3,3'-(4,4'-(9H-FLUORENE-9,9-DIYL)BIS(4J-PHENYLENE))BIS(OXY)BIS(1-CHLOROPROPANE-3,2-DIYL) BIS(2-(TERT-BUTOXYCARBONYLAMINO)ACETATE)(11)**

![Chemical structure](image)

Boc-Gly-OH (262 mg, 1.50 mmol, 4 equiv) was dissolved in anhydrous 1,4 dioxane (2 mL), and triethylamine was added (209 $\mu$L, 1.50 mmol) at room temperature, and the contents were stirred under an atmosphere of argon for 10 min. N-(3-Dimethylaminopropyl)-N’-ethylcarbodiimide hydrochloride (579 mg, 1.50 mmol) was added in one portion followed by the slow addition of a solution of derivative 1 (200 mg, 0.37 mmol, 1 equiv) in anhydrous 1,4 dioxane (3 mL) and catalytic 4-(Dimethylamino)pyridine (~2 mg), and the solution was stirred at rt for 29 h. The reaction mixture was then evaporated to dryness and extracted with dichloromethane/water (3 x 10 mL). The organic layer was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 100% dichloromethane to 5% methanol in hexane) to provide 11 (57 mg) as a white solid. $^1$H NMR (400 MHz, CDCls): $\delta$ 7.79 (d, $J = 7.2$, 2H), 7.39 (t, $J = 7.6$, 4H), 7.30 (t, $J = 7.2$, 2H), 7.15 (d, $J = 8.4$, 4H), 6.79 (d, $J = 8.4$, 4H), 5.39-5.36 (m, 2H), 5.05-5.09 (m, 2H), 4.19-4.12 (m, 4H), 3.97 (s, 4H), 3.88-3.83 (dd, $J = 11.6$, 4.8, 2H), 3.81-3.77
(dd, J = 11.6, 5.2, 2H), 1.48 (s, 18H); \(^{13}\)C NMR (100 MHz, CDCl\(_3\)): \(\delta\) 157.0, 155.8, 151.7, 140.1, 139.2, 129.4, 127.9, 127.6, 126.1, 120.4, 114.4, 80.3, 72.2, 65.9, 64.3, 42.5, 42.3, 28.5; HRMS (ESI) (m/z): calc'd for C\(_{45}\)H\(_{50}\)N\(_2\)O\(_2\)Na\(_2\) [M+Na\(^+\): 871.2740, found: 871.2729.

EXAMPLE 12

SYNTHESIS OF 2,2'-((3,3')-(4,4'-(9H-FLUOREN-9.9-DIYL)BIS(4,1-

PHENYLENE))BIS(OXY)BIS(1-CHLOROPROPANE -3,2-DIYL))BIS(OXY)BIS(2-

OXOETHANAMINIUM) CHLORIDE (12)

11 (14 mg) was dissolved in 3 mL CH\(_2\)Cl\(_2\) and Trifluoroacetic acid (0.4 mL) was added. The solution was stirred at rt for 6 h, then concentrated to dryness. Toluene (3 mL) was added, and the solution was concentrated to dryness again. The resulting residue was dissolved in 10 mL 0.2 M HCl, freeze and lyophilized to yield 12 as the bis-HCl salt (13 mg), as a white powder.
EXAMPLE 13

SYNTHESIS OF (S)-1-CHLORO-3-(4-(9-(4-((R)-2-HYDROXY-3-(2-(2-(2-HYDROXYETHOXY)ETHOXY)ETHOXY)PROPOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPAN-2-OL (13)

To a solution of derivative 7 (21 mg, 0.042 mmol, 1 equiv) in acetonitrile (1 mL) was added Triethylene glycol (1 mL) and bismuth(III) trifluoromethanesulfonate (~6 mg, 0.008 mmol, 1/5 equiv) and the mixture was stirred at room temperature for 4 h. After the crude was concentrated under reduced pressure, the resulting residue was extracted with ethyl acetate/water (3 x 5 mL). The organic layer was washed with deionized water (5 mL), was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: dichloromethane to 5% methanol in dichloromethane) to provide 13 (26 mg, 96%) as a white foam.

$^1$H NMR (400 MHz, DMSO-d$_6$): $\delta$ 7.90 (d, J = 7.2, 2H), 7.38 (t, J = 7.6, 4H), 7.30 (t, J = 7.6, 2H), 7.02-6.99 (dd, J = 8.8, 3.2, 4H), 6.84-6.80 (dd, J = 8.8, 6.8, 4H), 5.51 (br, 1H), 5.02 (br, 1H), 4.57 (br, 1H), 4.04-3.97 (m, 1H), 3.91-3.85 (m, 4H), 3.83-3.79 (m, 1H), 3.74-3.69 (dd, J = 11.2, 4.4, 1H), 3.65-3.61 (dd, J = 11.2, 5.6, 1H), 3.55-3.42 (m, 12H), 3.40-3.37 (m, 2H); $^1$C NMR (100 MHz, DMSO-d$_6$): $\delta$ 157.4, 157.0, 151.2, 139.3, 137.9, 137.5, 128.7, 128.7, 127.8, 127.5, 125.9, 120.5, 114.2, 72.3, 72.1, 70.2, 69.8, 69.7, 69.7, 69.5, 68.5, 68.5, 67.9, 63.6, 60.2, 46.7; HRMS (ESI) (m/z): calc'd for C$_{71}$H$_{44}$O$_{10}$NaCl [M+Na]$^+$: 671.2388, found: 671.2380.
EXAMPLE 14
SYNTHESIS OF (R)-3-(4-(9-(4-((S)-3-CHLORO-2-HYDROXYPROPOXY)PHENYL)-9H-FLUOREN-9-YL)PHENOXY)PROPANE-1,2-DIOL (14)

To a solution of derivative 7 (17 mg, 0.034 mmol, 1 equiv) in a mixture of acetonitrile (2 mL) and water (1 mL) was added Bismuth(III) trifluoromethanesulfonate (2 mg, 0.0034 mmol, 1/10 equiv) and the mixture was stirred at room temperature for 77 h. Then, the reaction mixture was evaporated to dryness and extracted with ethyl acetate/water (3 x 5 mL). The organic layer was dried over anhydrous magnesium sulfate, was filtered, and was concentrated under reduced pressure. The resulting residue was purified by flash column chromatography on silica gel (eluent: 20% ethyl acetate in hexane to 70% ethyl acetate) to provide 14 (16 mg, 93%) as a white solid. $^1$H NMR (400 MHz, DMSO-d$_6$): $\delta$ 7.90 (d, $J = 7.2$, 2H), 7.38 (t, $J = 7.6$, 4H), 7.30 (t, $J = 7.6$, 2H), 7.02-6.99 (dd, $J = 8.8$, 3.6, 4H), 6.82 (d, $J = 8.4$, 4H), 5.51 (d, $J = 4.4$, 1H), 4.88 (s, 1H), 4.60 (s, 1H), 4.01-3.97 (m, 1H), 3.94-3.90 (m, 3H), 3.81-3.69 (m, 3H), 3.65-3.61 (dd, $J = 11.2$, 5.6, 1H), 3.40 (d, $J = 5.2$, 2H); $^{13}$C NMR (100 MHz, DMSO-de): $\delta$ 157.5, 157.0, 151.2, 139.3, 137.9, 137.4, 128.7, 128.7, 127.8, 127.5, 125.9, 120.4, 114.2, 114.2, 69.9, 69.5, 68.8, 68.5, 63.6, 62.7, 46.7.

EXAMPLE 15
IN VITRO ACTIVITY OF SELECT COMPOUNDS

LNCaP cells were transiently cotransfected with PSA (6.1 kb)-luciferase (0.25 µg/well) in 24-well plates for 24 h prior to pre-treatment with compounds for 1 hour before the addition of synthetic androgen, R1881 (1 nM) to induce PSA production or vehicle. The total amount of plasmid DNA transfected was normalized to 0.75 µg/well by the addition of the empty vector. After 48 h of incubation with R1881, the cells were harvested, and relative luciferase activity was determined. Test
compounds were added to the cells at various concentrations and activity for each treatment was normalized to the predicted maximal activity induction (in the absence of test compounds, vehicle only). Plotting of sigmoidal curves (Boltzmann Function) and IC50 calculations were done using OriginPro 8.1 Software (Northampton, MA, USA).

Furthermore, toxicity was assessed by both microscopic examination and reduction of protein levels. Solubility was assessed both macroscopically (cloudy media) and microscopically (formation of granules or crystals).

TABLE 3 shows the compounds tested using the above-described assays and their respective activities.

<table>
<thead>
<tr>
<th>COMPOUND</th>
<th>IC50</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="image1.png" alt="Chemical Structure 1" /></td>
<td>0.8 µM</td>
</tr>
<tr>
<td><img src="image2.png" alt="Chemical Structure 4" /></td>
<td>0.9 µM</td>
</tr>
<tr>
<td><img src="image3.png" alt="Chemical Structure 5" /></td>
<td>0.8 µM</td>
</tr>
</tbody>
</table>
In vivo dose response of compounds of the invention is determined according to the following procedure: Male athymic SCID-NOD mice, 6- to 8-weeks old, are inoculated subcutaneously with LNCaP cells \((1 \times 10^6)\) suspended in 75 \(\mu\)l of RPMI 1640 (5% FBS) and 75 \(\mu\)l of Matrigel (Becton Dickinson Labware) in the flank region via a 27-gauge needle under isofluorane anesthesia. Mice bearing LNCaP subcutaneous tumors are castrated when tumor volumes are approximately 100 mm\(^3\). Seven days after castration, mice are injected intravenously by tail vein every other day for a total of 7 doses with compounds of the invention in 15% DMSO and 25.5% PEG.

<table>
<thead>
<tr>
<th>COMPOUND</th>
<th>IC50</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="image" alt="Compound 8" /></td>
<td>1.5 (\mu)M</td>
</tr>
<tr>
<td><img src="image" alt="Compound 9" /></td>
<td>&lt; 0.2 (\mu)M</td>
</tr>
<tr>
<td><img src="image" alt="Compound 10" /></td>
<td>1.4 (\mu)M</td>
</tr>
</tbody>
</table>
The experiment is complete 2 days after the last injection. Tumours are measured with calipers and their volumes calculated by the formula L x W x H x 0.5236. Tumor volume as a function of compound dose is plotted.

Dose response of comparative compounds are also determined according to the above procedure.

Although various embodiments of the invention are disclosed herein, many adaptations and modifications may be made within the scope of the invention in accordance with the common general knowledge of those skilled in this art. Such modifications 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 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 the examples and drawings.
CLAIMS

1. A compound having a structure of Formula I:

or a pharmaceutically acceptable salt or tautomer thereof, wherein:

- at least one $Z^1$ is independently C-Q;
- at least one $Z^2$ is independently C-T, CCH$_3$, CF, CCl, CBr, Cl, COH, CG$^1$, COG$^1$, CNH$_2$, CNHG$^1$, CN(G$^1$)$_2$, COS0$_3$H, COP0$_3$H$_2$, CSG$^1$, CSOG$^1$, or CSO$_2$G$^1$;
- $Z^3$, $Z^4$ and each remaining $Z^1$ and $Z^2$ are, at each occurrence, independently C-T, N, CH, CCH$_3$, CF, CC1, CBr, Cl, COH, CG$^1$, COG$^1$, CNH$_2$, CNHG$^1$, CN(G$^1$)$_2$, COSO$_3$H, COPO$_3$H$_2$, CSG$^1$, CSOG$^1$, or CSO$_2$G$^1$;
- Q is
- J is G$^1$, O, CH$_2$, CHG$^1$, C(G$^1$)$_2$, S, NH, NG$^1$, SO, S0$_2$, or NR;
- M is H, F, Cl, Br, CH$_2$OH, CH$_2$OD, CH$_2$OG$^1$, CH$_2$F, CH$_2$Cl, CHCl$_2$, CCl$_3$, CH$_2$Br, CHBr$_2$, CBr$_3$ or C≡CH;
- L is H or A-D;
- A is O, S, NH, NG$^1$, N$^+$H$_2$ or N$^+$HG$^1$;
- D is, at each occurrence, independently H, G$^1$, R,

or a moiety from TABLE 1;

- each of q, r and t may independently be 0, 1, 2, 3, 4, 5, 6 or 7;
- n is 0, 1, 2, 3, 4, 5, 6, 7 or 8.
T is, at each occurrence, independently; J is, at each occurrence, independently; G, O, CHG, C(G)2, S, NH, NG, SO, S02, or NR;

M2 is, at each occurrence, independently H, CH3, F, Cl, Br, CH2F, CH2Cl, CHCl2, CCl3, CH2Br, CHBr2, CBr3, CH2OH, CH2OD, CH2OJ", G1, CH2OG1, CH2OR, CH2OG1OG1, GOG11, GOG11OG1'1, CH2SG, CH2NH2, CH2NHG1, CH2NG2, or C=CH;

L2 is, at each occurrence, independently H or A2-D2;

A2 is, at each occurrence, independently O, S, SO, S02, NH, NG1, N+H2, orN +HG1;

D2 is, at each occurrence, independently H, G1, R,

or a moiety selected from TABLE 1;

each of u, y and j are, at each occurrence, independently 0, 1, 2, 3, 4, 5, 6 or 7;

m is, at each occurrence, independently 0, 1, 2, 3, 4, 5, 6, 7 or 8;

J" and J"" are, at each occurrence, independently a moiety selected from TABLE 1;

G1, G1" and G1"" are, at each occurrence, independently a linear or branched, aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C10 alkyl, wherein the optional substituents for the C1-C10 alkyl are oxo, CH2C02R', OJ"", COOH, R1, OH, OR1, F, Cl, Br, I, NH2, NHR, N(R')2, CN, SH, SR1, SO3H, SO3R1, SO2R1, OSO3R1, OR2, CO2R1, CONHR1, CONHR2, CON(R')2, NHR3, OP03H, CON2R2, NR3R2 or N02;

each R' is independently H, linear or branched, aromatic cyclic or non-aromatic cyclic, substituted or unsubstituted, saturated or unsaturated C1-C10 alkyl or a metal counter ion, wherein the metal counter ion is Li, Na, K, Mg or Ca;

each R1 is independently unsubstituted C1-C10 alkyl; and
each R and R² are independently Ci-Cio acyl.

2. The compound of claim 1, wherein at least one Z² is C-T.

3. The compound of claim 1, wherein the compound has a structure of Formula II:

![Image of structure II]

4. The compound of claim 1, wherein the compound has a structure of Formula III:

![Image of structure III]

wherein:

R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ are each independently hydrogen, halo, or linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cio alkyl.

5. The compound of claim 1, wherein the compound has a structure of Formula IV:

![Image of structure IV]
6. The compound of claim 1, wherein the compound has a structure of Formula V:

![Structure V](image)

7. The compound of claim 1, wherein the compound has a structure of Formula Va, Vb, Vc or Vd:

![Structure Va](image)

![Structure Vb](image)

![Structure Vc](image) or

![Structure Vd](image)

8. The compound of any one of claims 1 to 7, wherein the compound has one of the following structures:
wherein:

$R^{11}$ is hydrogen or linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cio alkyl;

$Y$ is Cl or OH;

$q$ is 0, 1, 2, 3, 4, 5, 6 or 7; and

$m$ is 0, 1, 2, 3, 4, 5, 6, 7 or 8.

9. The compound of any one of claims 1 to 5, wherein $J$ is $G^1$, O, CH$_2$, CHG$^1$, CG$^A$, NH, SO, or NR.

10. The compound of any one of claims 1 to 5, wherein $J$ is O.
11. The compound of any one of claims 1 to 7, 9 or 10, wherein M is Cl, Br, CH₂OH, CH₂OD, CH₂OG¹, CH₂F, CH₂Cl, CHCl₂, CCl₃, CH₂Br, CHBr₂, CBr₃, or C≡CH.

12. The compound of any one of claims 1 to 7, 9 or 10, wherein M is CH₂OH, CH₂F, C≡CH, CH₂OCH₂C≡CH, CH₂OCH₃, CH₂0-2-propyl, CH₂0-i-butyl, or CH₂OD, wherein D is

13. The compound of any one of claims 1 to 7, 9 or 10, wherein M is CH₂OH.

14. The compound of any one of claims 1 to 7, 9 or 10, wherein M is H.

15. The compound of any one of claims 1 to 7 or 9-14, wherein L is H.

16. The compound of any one of claims 1 to 7 or 9-14, wherein L is A-D.

17. The compound of any one of claims 1 to 7 or 9-14, wherein A is O.

18. The compound of any one of claims 16 or 17, wherein D is H, R, or a moiety from TABLE 1; and each of q, r and t is independently 0, 1, 2, 3, 4, 5, 6 or 7.

19. The compound of any one of claims 16 or 17, wherein D is H.

20. The compound of any one of claims 16 or 17, wherein D is R.

21. The compound of any one of claims 16 or 17, wherein D is a moiety selected from TABLE 1.

22. The compound of claim 21, wherein the moiety from TABLE 1

23. The compound of any one of claims 1 to 22, wherein n is 0.
24. The compound of any one of claims 1 to 22, wherein \( n \) is 1, 2, 3, 4, or 5.

25. The compound of any one of claims 1 to 22, wherein \( n \) is 1.

26. The compound of any one of claims 1 to 5, wherein \( J^2 \) is \( G^1, O, \text{CH}_2, \text{CHG}^1, \text{CG}^\wedge, \text{NH}, \text{SO}, \text{or NR} \).

27. The compound of any one of claims 1 to 5, wherein \( J^2 \) is \( O \).

28. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( H, \text{CH}_2\text{F}, \text{CH}_2\text{Cl}, \text{CH}_2\text{Br}, \text{CH}_2\text{OH}, \text{CH}_2\text{O}^\wedge, \text{CF}_2\text{O}^\wedge, \text{or C≡CH} \).

29. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( \text{CH}_2\text{F} \).

30. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( \text{CH}_2\text{Cl} \).

31. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( \text{CH}_2\text{Br} \).

32. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( \text{CH}_2\text{OH} \).

33. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( H \).

34. The compound of any one of claims 1 to 7 or 9-27, wherein \( M^2 \) is \( \text{C≡CH} \).

35. The compound of any one of claims 1 to 7 or 9-34, wherein \( L^2 \) is \( H \).

36. The compound of any one of claims 1 to 7 or 9-34, wherein \( L^2 \) is \( A^2\cdot D^2 \).

37. The compound of claim 36, wherein \( A^2 \) is \( O \).

38. The compound of any one of claims 36 or 37, wherein \( D^2 \) is \( H, \text{RO}_u\text{OH}^\wedge, \text{OG}^1, \text{OR}_j\text{OR}, \text{or a moiety from TABLE 1}; \text{and each of u, y and j is independently 0, 1, 2, 3, 4, 5, 6 or 7.} \)
39. The compound of any one of claims 36 or 37, wherein \( D_2 \) is \( H \).
40. The compound of any one of claims 36 or 37, wherein \( D_2 \) is \( R \).
41. The compound of any one of claims 36 or 37, wherein \( D_2 \) is a moiety from TABLE 1.
42. The compound of claim 41, wherein the moiety from TABLE 1

\[
\begin{align*}
\text{O} & \quad \text{NH}_2 \\
\text{O} & \quad \text{NH}_3^+ \\
\end{align*}
\]

is or.
43. The compound of any one of claims 1 to 42, wherein \( m \) is 0.
44. The compound of any one of claims 1 to 42, wherein \( m \) is 1, 2, 3, 4, or 5.
45. The compound of any one of claims 1 to 42, wherein \( m \) is 1.
46. The compound of any one of claims 1 to 7, wherein \( M \) is \( \text{C}_3\text{H}_4\text{OH} \) and \( L \) is \( \text{OH} \).
47. The compound of any one of claims 1 to 7, wherein \( M_2 \) is \( \text{CH}_2\text{C}_1 \) and \( L \) is \( \text{OH} \).
48. The compound of any one of claims 1 to 7, wherein \( M \) is \( \text{CH}_2\text{OH} \), \( M_2 \) is \( \text{CH}_2\text{C}_1 \), \( L \) is \( \text{OH} \) and \( L_2 \) is \( \text{OH} \).
49. The compound of claim 48, wherein \( n \) and \( m \) are each 1.
50. The compound of any one of claims 4 to 49, wherein at least one of \( R_3 \), \( R_4 \), \( R_5 \), \( R_6 \), \( R_7 \), \( R_8 \), \( R_9 \) or \( R_{10} \) is hydrogen.
51. The compound of any one of claims 4 to 49, wherein each of \( R_3 \), \( R_4 \), \( R_5 \), \( R_6 \), \( R_7 \), \( R_8 \), \( R_9 \) and \( R_{10} \) are hydrogen.
52. The compound of any one of claims 4 to 49, wherein each of \( R_3 \), \( R_4 \), \( R_5 \) and \( R_6 \) are hydrogen.
53. The compound of any one of claims 4 to 49, wherein at least one of \( R_3 \), \( R_4 \), \( R_5 \), \( R_6 \), \( R_7 \), \( R_8 \), \( R_9 \) or \( R_{10} \) is methyl.
54. The compound of any one of claims 4 to 49, wherein each of \( R_3 \), \( R_4 \), \( R_5 \), \( R_6 \), \( R_7 \), \( R_8 \), \( R_9 \) and \( R_{10} \) are methyl.
55. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵ and R⁶ are methyl.
56. The compound of any one of claims 4 to 49, wherein at least one of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ or R¹⁰ is fluoro.
57. The compound of any one of claims 4 to 49, wherein each R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ are fluoro.
58. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵, and R⁶ are fluoro.
59. The compound of any one of claims 4 to 49, wherein at least one of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ or R¹⁰ is chloro.
60. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ are chloro.
61. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵ and R⁶ are chloro.
62. The compound of any one of claims 4 to 49, wherein at least one of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ or R¹⁰ is bromo.
63. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ are bromo.
64. The compound of any one of claims 4 to 49, wherein each of R³, R⁴, R⁵ and R⁶ are bromo.
65. The compound of any one of claims 1 to 5, wherein Q is
n is 0, 1, 2, 3, 4, 5, 6, 7 or 8;
each of q, r, and t is independently 0, 1, 2, 3, 4, 5, 6 or 7;

m is 0, 1, 2, 3, 4, 5, 6, 7 or 8;

each of u, j and y is independently 0, 1, 2, 3, 4, 5, 6 or 7;

each $G^1$ is independently linear or branched, substituted or unsubstituted, saturated or unsaturated $Ci-Cio$ alkyl, wherein the optional substituents are selected from oxo, $OJ''$, COOH, OH, F, Cl, Br, I, NH$_2$, CN, SH, S0$_3$H, CONH$_2$, OP0$_3$H$_3$ and N0$_2$.

66. The compound of any one of claims 1 to 5, wherein Q

![Chemical structures](image-url)
n is 0, 1, 2, 3, 4, 5, 6, 7 or 8;
each of q, r, and t is independently 0, 1, 2, 3, 4, 5, 6 or 7;
m is 0, 1, 2, 3, 4, 5, 6, 7 or 8;
each of u, j and y is independently 0, 1, 2, 3, 4, 5, 6 or 7;
each G₁ is independently linear or branched, substituted or unsubstituted, saturated or unsaturated Ci-Cίo alkyl, wherein the optional substituents are selected from oxo, OJ"", COOH, OH, F, Cl, Br, I, NH₂, CN, SH, SO₃H, CONH₂, OP0₃H₃, and N0₂.

67. The compound of any one of claims 1 to 5, wherein Q is
n is 0, 1, 2, 3, 4, 5, 6, 7 or 8;
m is 0, 1, 2, 3, 4, 5, 6, 7 or 8; and
each G is independently linear or branched, substituted or unsubstituted,
saturated or unsaturated Ci-Ci alkyl, wherein the optional substituents are selected
from oxo, OJ", COOH, OH, F, Cl, Br, I, NH₂, CN, SH, S0₃H, CONH₂, OP0₃H₃, and
N0₂.

68. The compound of any one of claims 1 to 5, wherein Q is
n is 0, 1, 2, 3, 4, 5, 6, 7 or 8;
q is 0, 1, 2, 3, 4, 5, 6 or 7;
m is 0, 1, 2, 3, 4, 5, 6 or 8;
u is 0, 1, 2, 3, 4, 5, 6 or 7; and
each Gᵢ is independently linear or branched, substituted or unsubstituted, saturated or unsaturated C₁-C₁₀ alkyl, wherein the optional substituents are selected from oxo, OJ‴, COOH, OH, F, Cl, Br, I, NH₂, CN, SH, S₀₃H, CONH₂, OP₀₃H₃, and NO₂.

69. The compound of any one of claims 1 to 5, wherein q is
70. The compound of any one of claims 1 to 5, wherein Q
71. The compound of any one of claims 1 to 5, wherein T is
n, m and q are each independently 0, 1, 2, 3, 4, 5, 6, 7 or 8.

72. The compound of any one of claims 1 to 5, wherein T is
73. The compound of any one of claims 1 to 5, wherein Q is
74. The compound of any one of claims 1 to 5, wherein Q is

and T is

75. The compound of any one of claims 1 to 3, wherein Z\textsuperscript{3}, Z\textsuperscript{4} and each remaining Z\textsuperscript{1} and Z\textsuperscript{2} is independently CCH\textsubscript{3}; CH; CF, CCl or CBr.
76. The compound of any one of claims 1 to 3, wherein \( Z^3, Z^4 \) and each remaining \( Z^1 \) and \( Z^2 \) is CH.

77. The compound of any one of claims 1 to 76, wherein one or more of the OH groups of the compound is substituted to replace the H with a moiety from TABLE 1.

78. The compound of claim 77, wherein the moiety from TABLE 1 is \( \text{O} \text{NH}_2 \) or \( \text{O} \text{NH}_3^+ \).

79. A compound having one of the following structures:
80. Use of the compound of any one of claims 1 to 79, for modulating androgen receptor (AR) activity.

81. The use of claim 80, wherein modulating androgen receptor (AR) activity is in a mammalian cell.
82. The use of any one of claims 80 or 81, wherein 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 muscular atrophy, and age-related macular degeneration.

83. The use of claim 82, wherein the indication is prostate cancer.

84. The use of claim 83, wherein the prostate cancer is castration resistant prostate cancer.

85. The use of claim 83, wherein the prostate cancer is androgen-dependent prostate cancer.

86. The use of claim 82, wherein the spinal and bulbar muscular atrophy is Kennedy's disease.

87. A method of modulating androgen receptor (AR) activity, the method comprising administering a compound, or pharmaceutically acceptable salt thereof, of any one of claims 1 to 79 to a subject in need thereof.

88. The method of claim 87, wherein modulating androgen receptor (AR) activity is for the treatment of one or more of the following: 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.

89. The method of claim 88, wherein the prostate cancer is castration resistant prostate cancer.

90. The method of claim 88, wherein the prostate cancer is androgen-dependent prostate cancer.

91. The method of claim 88, wherein the spinal and bulbar muscular atrophy is Kennedy's disease.

92. A pharmaceutical composition comprising a compound of any one of claims 1 to 79 and a pharmaceutically acceptable carrier.
93. A pharmaceutical composition comprising a compound of any one of claims 1 to 79, an additional therapeutic agent and a pharmaceutically acceptable carrier.

94. The pharmaceutical composition of claim 93, 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.

95. The pharmaceutical composition of claim 93, wherein the additional therapeutic agent is MDV3100, TOK 001, ARN-509; abiraterone, bicalutamide, nilutamide, flutamide, cyproterone acetate, docetaxel, Bevacizumab (Avastin), OSU-HDAC42, VITAXIN, sunitinib, ZD-4054, VN/124-1, Cabazitaxel (XRP-6258), flpiimumab, OGX 427, OGX 011, finasteride, dutasteride, turosteride, bexlosteride, izonsteride, FCE 28260, SKF105,111 or a related compound thereof.

96. Use of the pharmaceutical composition of any one of claims 92 to 95 for modulating androgen receptor (AR) activity.

97. The use of claim 96, wherein modulating androgen receptor (AR) activity is in a mammalian cell.

98. The use of any one of claims 96 or 97, wherein 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 muscular atrophy, and age-related macular degeneration.

99. The use of claim 98, wherein the indication is prostate cancer.

100. The use of claim 99, wherein the prostate cancer is castration resistant prostate cancer.

101. The use of claim 99, wherein the prostate cancer is androgen-dependent prostate cancer.
102. The use of claim 98, wherein the spinal and bulbar muscular atrophy is Kennedy's disease.

103. A method of modulating androgen receptor (AR) activity, the method comprising administering the pharmaceutical composition of any one of claims 92 to 95 to a subject in need thereof.

104. The method of claim 103 wherein modulating androgen receptor (AR) activity is for the treatment of one or more of the following: 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.

105. The method of claim 104, wherein the spinal and bulbar muscular atrophy is Kennedy's disease.

106. The use of claim 104, wherein the indication is prostate cancer.

107. The use of claim 106, wherein the prostate cancer is castration resistant prostate cancer.

108. The use of claim 106, wherein the prostate cancer is androgen-dependent prostate cancer.
INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) ... Box 1450, Alexandria, Virginia 22313-1450
Facsimile No. 571-273-3201 -7774
Form PCT/ISA /210 (second sheet) (July 2009)

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC(8): A01N 31/14; A61K 31/075; C07C 43/295 (2012.01)
USPC: 514/721; 568/630 (text search)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
USPC: 514/224.5, 325, 185 (text search) Find search terms below

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
PubWEST (PGPB,USPT,USOC,EPAB,JPAB), Google Scholar

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category* Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No.
Y WO 2010/000866 A1 (SADAR et al.) 07 January 2010 (07.01.2010) pg 4, para 4 - pg 5, para 2; pg 16, para 3 - pg 22, para 1; Table 1; Table 2 1-10, 26-27, 46-49, 65-76 and 79

Further documents are listed in the continuation of Box C.

* Special categories of cited documents:
“A” document defining the general state of the art which is not considered to be of particular relevance
“E” earlier application or patent but published on or after the international filing date
“L” 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)
“O” document referring to an oral disclosure, use, exhibition or other means
“P” document published prior to the international filing date but later than the priority date claimed
“T” 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
“X” 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
“Y” 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

Date of the actual completion of the international search
03 July 2012 (03.07.2012)

Date of mailing of the international search report
18 JUL 2012

Name and mailing address of the ISA/US
Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, Virginia 22313-1450
Facsimile No. 571-273-3201

Authorized officer: Lee W. Young
PCT Helpdesk: 571-272-4300
PCT OSP: 571-272-7774

Form PCT/ISA /210 (second sheet) (July 2009)
INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 12/33959

<table>
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<th>Box No. II</th>
<th>Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)</th>
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<tr>
<td><strong>This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:</strong></td>
<td></td>
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<tr>
<td><strong>1.</strong></td>
<td>Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:</td>
</tr>
<tr>
<td><strong>2.</strong></td>
<td>Claims 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:</td>
</tr>
<tr>
<td><strong>3.</strong></td>
<td>Claims Nos.: 11-25, 28-45, 50-64, 77-78 and 80-108 because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).</td>
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<th>Box No. III</th>
<th>Observations where unity of invention is lacking (Continuation of item 3 of first sheet)</th>
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<td><strong>This International Searching Authority found multiple inventions in this international application, as follows:</strong></td>
<td></td>
</tr>
<tr>
<td><strong>1.</strong></td>
<td>As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.</td>
</tr>
<tr>
<td><strong>2.</strong></td>
<td>As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.</td>
</tr>
<tr>
<td><strong>3.</strong></td>
<td>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 claims Nos.:</td>
</tr>
<tr>
<td><strong>4.</strong></td>
<td>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 claims Nos.:</td>
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**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.

Form PCT/ISA/210 (continuation of first sheet (2)) (July 2009)