

**COMBINATION OF ANTI-CLUSTERIN OLIGONUCLEOTIDE WITH HSP90 INHIBITOR
FOR THE TREATMENT OF PROSTATE CANCER**

5 This application claims priority of U.S. Provisional Application No. 61/453,102, filed March 15, 2011, the contents of which are hereby incorporated by reference.

10 Throughout this application, various publications are referenced, including referenced in parenthesis. Full citations for publications referenced in parenthesis may be found listed in alphabetical order at the end of the specification immediately preceding the claims. The disclosures of all referenced publications in their entirety are hereby incorporated by
15 reference into this application in order to more fully describe the state of the art to which this invention pertains.

Field of the Invention

20 The subject invention relates to combination therapy for treating prostate cancer.

Background of the Invention

Prostate cancer (PCa) is the most common cancer and the third most common cause of cancer related mortality in men in the United
25 States (Jemal et al., 2006). Androgen ablation remains the standard effective therapy for patients with advanced PCa, inhibiting proliferation and inducing apoptosis in tumor cells (Kyprianou et al., 1990). Unfortunately, after short-term remissions, surviving tumor cells recur with castrate resistant prostate cancer (CRPC)
30 and death usually within 3 years in most men (Gleave et al., 1999). CRPC progression results from mechanisms attributed to re-activation of androgen receptor axis (Knudsen et al., 2009), alternative mitogenic growth factor pathways (Miyake et al., 2000; Culig et al., 2004), and stress-induced prosurvival gene (Gleave et al.,
35 al., 1999; Miyake et al., 1999) and cytoprotective chaperone networks (Rocchi et al., 2004; Miyake et al., 2000). To

significantly improve survival in men with PCa, new therapeutic strategies to inhibit the appearance of this phenotype must be developed. It has been observed that numerous proteins are expressed in increased amounts by prostate tumor cells following androgen withdrawal. At least some of these proteins are assumed to be associated with the observed apoptotic cell death which is observed upon androgen withdrawal. (Raffo et al., 1995; Krajewska et al., 1996; McDonnell et al., 1992). The functions of many of the proteins, however, is not completely understood. Clusterin (also known as sulfated glycoprotein-2 (SGP-2) or TRPM-2) is within this latter category.

Clusterin

Clusterin is a cytoprotective chaperone protein that promotes cell survival and confers broad-spectrum resistance to cancer treatments (Chi et al. 2005). In Sensibar et al., Cancer Research 55: 2431-2437, 1995, the authors reported on LNCaP cells transfected with a gene encoding clusterin, and watched to see if expression of this protein altered the effects of tumor necrosis factor α (TNF α), to which LNCaP cells are very sensitive. Treatment of the transfected LNCaP cells with TNF α was shown to result in a transient increase in clusterin levels for a period of a few hours, but these levels had dissipated by the time DNA fragmentation preceding cell death was observed.

As described in U.S. Patent No. 7,534,773, the contents of which are incorporated by reference, enhancement of castration-induced tumor cell death and delay of the progression of androgen-sensitive cancer cells to androgen-independence may be achieved by inhibiting the expression of clusterin by the cells.

Custirsen

Custirsen is a second-generation antisense oligonucleotide that inhibits clusterin expression. Custirsen is designed specifically to bind to a portion of clusterin mRNA, resulting in the inhibition of

the production of clusterin protein. The structure of custirsen is available, for example, in U.S. Patent No. 6,900,187, the contents of which are incorporated herein by reference. A broad range of studies have shown that custirsen potently regulates the expression of clusterin, facilitates apoptosis, and sensitizes cancerous human prostate, breast, ovarian, lung, renal, bladder, and melanoma cells to chemotherapy (Miyake et al. 2005), see also, U.S. Patent Application Publication No. 2008/0119425 A1. In a clinical trial for androgen-dependent prostate cancer, the drugs flutamide and buserelin were used together in combination with custirsen, increasing prostate cancer cell apoptosis (Chi et al. 2004; Chi et al., 2005).

Hsp90

Heat shock protein 90 (Hsp90) is an ATPase-dependent molecular chaperone required for protein folding, maturation and conformational stabilization of many "client" proteins (Young et al., 2000; Kamal et al., 2003). Hsp90 interacts with several proteins involved in CRPC, including growth factor receptors, cell cycle regulators and signaling kinases like Akt, androgen receptor (AR) or Raf-1, (Whitesell et al., 2005; Takayama et al., 2003). Tumor cells express higher Hsp90 levels compared with benign cells (Kamal et al., 2003; Chiosis et al., 2003), and Hsp90 inhibition has emerged as an exciting target in CRPC and other cancers. Many Hsp90 inhibitors were developed targeting its ATP-binding pocket, including natural compounds such as geldanamycin and its analogs, or synthetic compounds. These agents have been shown to inhibit Hsp90 function and induce apoptosis in preclinical studies of colon, breast, PCa and other cancers (Kamal et al., 2003; Solit et al., 2003; Solit et al., 2002).

Combination Therapy

The administration of two drugs to treat a given condition, such as prostate cancer, raises a number of potential problems. *In vivo* interactions between two drugs are complex. The effects of any single drug are related to its absorption, distribution, and

elimination. When two drugs are introduced into the body, each drug can affect the absorption, distribution, and elimination of the other and hence, alter the effects of the other. For instance, one drug may inhibit, activate or induce the production of enzymes involved in a metabolic route of elimination of the other drug (Guidance for Industry. *In vivo* drug metabolism/drug interaction studies - study design, data analysis, and recommendations for dosing and labeling). Thus, when two drugs are administered to treat the same condition, it is unpredictable whether each will complement, have no effect on, or interfere with, the therapeutic activity of the other in a human subject.

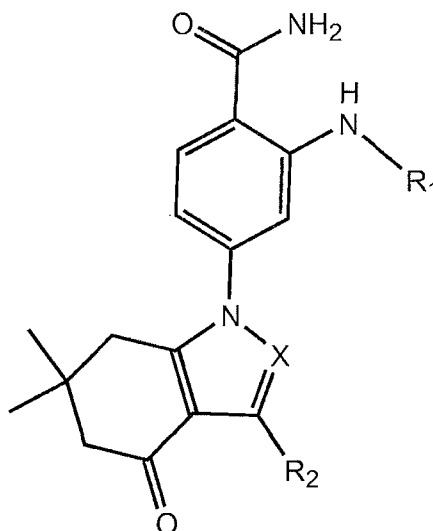
Not only may the interaction between two drugs affect the intended therapeutic activity of each drug, but the interaction may increase the levels of toxic metabolites (Guidance for Industry. *In vivo* drug metabolism/drug interaction studies - study design, data analysis, and recommendations for dosing and labeling). The interaction may also heighten or lessen the side effects of each drug. Hence, upon administration of two drugs to treat a disease, it is unpredictable what change will occur in the profile of each drug.

Additionally, it is difficult to accurately predict when the effects of the interaction between the two drugs will become manifest. For example, metabolic interactions between drugs may become apparent upon the initial administration of the second drug, after the two have reached a steady-state concentration or upon discontinuation of one of the drugs (Guidance for Industry. *In vivo* drug metabolism/drug interaction studies - study design, data analysis, and recommendations for dosing and labeling).

Thus, the success of one drug or each drug alone in an *in vitro* model, an animal model, or in humans, may not correlate into efficacy when both drugs are administered to humans.

Summary of the Invention

The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin
 5 expression and ii) a Heat Shock Protein 90 (Hsp90) inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

10 R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from
 15 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ - $C_6)$ alkylamino, nitro, halo $(C_1$ - $C_6)$ alkyl, halo $(C_1$ - $C_6)$ alkoxy, or carboxamide, wherein

when R_1 is a C_1 - C_{14} alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R_4 , carbonyl, ethenyl,
 20 ethynyl or a moiety selected from N, O, S, SO_2 , or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R_4 is
 25 (i) heteroaryl,

- (ii) aryl,
 (iii) saturated or unsaturated C₃-C₁₀
 cycloalkyl, or
 (iv) saturated or unsaturated C₂-C₁₀
 heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or
 unsaturated cycloalkyl, or saturated or
 unsaturated heterocycloalkyl,
 independently, is optionally substituted
 with at least one group, which
 independently is hydroxy, halo, amino,
 cyano, carboxy, carboxamido, nitro, oxo,
 -S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-
 aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂,
 -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-
 C₆)alkoxy, or mono- or di-(C₁-
 C₁₀)alkylamino; and R₄ is optionally fused
 to a C₆-C₁₀ aryl group, C₅-C₈ saturated
 cyclic group, or a C₄-C₁₀ heterocycloalkyl
 group; and

R₁ is optionally substituted at any available position
 with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-
 C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo,
 amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-
 C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-
 aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀
 alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-
 C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
 wherein

Z is OR₆ or -N(R₆)₂, wherein

each R₆ is independently -H or C₁-C₆ alkyl, or
 N(R₆)₂ represents pyrrolidinyl, piperidinyl,
 piperazinyl, azepanyl, 1,3- or 1,4-
 diazepanyl, or morpholinyl, each of which

is optionally substituted with hydroxy, amino, aminoalkyl, C₁-C₆ alkyl, mono- or di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or halogen, and

R₀ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀ alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl; and

R₅ is

(1) heteroaryl,

(2) aryl,

(3) saturated or unsaturated C₅-C₁₀ cycloalkyl, or

(4) saturated or unsaturated C₅-C₁₀ heterocycloalkyl,

and

the R₅ groups are optionally substituted at least one group which is independently hydroxy, oxo, halo, amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl, -SO-(C₁-C₆) alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆) alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino;

R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

R₃ is H, halogen, or CH₃,

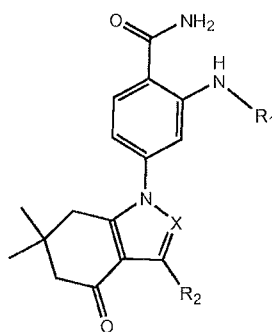
or a prodrug thereof, each in an amount that when in combination with the other is effective to treat the mammalian subject.

Hsp90 inhibitors having structure above are described in U.S. Patent No. 7,928,135, the entire contents of which are hereby incorporated herein by reference.

The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor, which inhibitor is other than Hsp90i-1, each in an amount that when in combination with the other is effective to treat the mammalian subject.

The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50 nmol/L, or a prodrug thereof, each in an amount that when in combination with the other is effective to treat the mammalian subject.

The present invention provides a pharmaceutical composition comprising an amount of an oligonucleotide which reduces clusterin expression, and a Hsp90 inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ -

C₆)alkylamino, nitro, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, or carboxamide, wherein

when R₁ is a C₁-C₁₄ alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R₄, carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO₂, or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R₄ is

(i) heteroaryl,

(ii) aryl,

(iii) saturated or unsaturated C₃-C₁₀ cycloalkyl, or

(iv) saturated or unsaturated C₂-C₁₀ heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or unsaturated cycloalkyl, or saturated or unsaturated heterocycloalkyl,

independently, is optionally substituted with at least one group, which independently is hydroxy, halo, amino, cyano, carboxy, carboxamido, nitro, oxo,

-S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino; and R₄ is optionally fused

to a C₆-C₁₀ aryl group, C₅-C₈ saturated cyclic group, or a C₄-C₁₀ heterocycloalkyl group; and

R₁ is optionally substituted at any available position with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo,

amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀ alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
 5 wherein

Z is OR₀ or -N(R₆)₂, wherein

each R₆ is independently -H or C₁-C₆ alkyl, or
 N(R₆)₂ represents pyrrolidinyl, piperidinyl,
 10 piperazinyl, azepanyl, 1,3- or 1,4-
 diazepanyl, or morpholinyl, each of which
 is optionally substituted with hydroxy,
 amino, aminoalkyl, C₁-C₆ alkyl, mono- or
 di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or
 15 halogen, and

R₀ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀
 alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl;
 and

R₅ is

- 20 (1) heteroaryl,
- (2) aryl,
- (3) saturated or unsaturated C₅-C₁₀
 cycloalkyl, or
- (4) saturated or unsaturated C₅-C₁₀
 25 heterocycloalkyl,

and

the R₅ groups are optionally substituted at
 least one group which is independently
 hydroxy, oxo, halo, amino, cyano, nitro, -SH,
 30 -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl,
 -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
 (C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or
 mono- or di-(C₁-C₁₀)alkylamino;

R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or
 35 halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

R₃ is H, halogen, or CH₃,

or a prodrug thereof, for use in treating a mammalian subject
5 affected by prostate cancer.

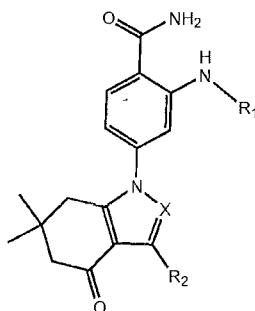
Hsp90 inhibitors having structure above are described in U.S.
Patent No. 7,928,135, the entire contents of which are hereby
incorporated herein by reference.

10

The present invention provides a pharmaceutical composition
comprising an amount of an oligonucleotide which reduces clusterin
expression, and a Hsp90 inhibitor, which inhibitor is other than
Hsp90i-1, for use in treating a mammalian subject affected by
15 prostate cancer.

The present invention provides a pharmaceutical composition
comprising an amount of an oligonucleotide which reduces clusterin
expression, and a Hsp90 inhibitor which binds to Hsp90 α and Hsp90 β
20 with a K_a of less than 50 nmol/L, or a prodrug thereof, for use in
treating a mammalian subject affected by prostate cancer.

The present invention provides an oligonucleotide which reduces
clusterin expression for use in combination with a Hsp90 inhibitor
25 having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R₁ is H, C₁-C₁₄ alkyl, C₁-C₁₀ haloalkyl, C₃-C₇ cycloalkyl, heterocycloalkyl, C₁-C₆ acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, hydroxy, amino, mono- or di-(C₁-C₆)alkylamino, nitro, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, or carboxamide, wherein

when R₁ is a C₁-C₁₄ alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R₄, carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO₂, or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R₄ is

- (i) heteroaryl,
- (ii) aryl,
- (iii) saturated or unsaturated C₃-C₁₀ cycloalkyl, or
- (iv) saturated or unsaturated C₂-C₁₀ heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or unsaturated cycloalkyl, or saturated or unsaturated heterocycloalkyl, independently, is optionally substituted with at least one group, which independently is hydroxy, halo, amino, cyano, carboxy, carboxamido, nitro, oxo, -S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino; and R₄ is optionally fused

to a C₆-C₁₀ aryl group, C₅-C₈ saturated cyclic group, or a C₄-C₁₀ heterocycloalkyl group; and

R₁ is optionally substituted at any available position with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo, amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀ alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅, wherein

Z is OR₆ or -N(R₆)₂, wherein

each R₆ is independently -H or C₁-C₆ alkyl, or N(R₆)₂ represents pyrrolidinyl, piperidinyl, piperazinyl, azepanyl, 1,3- or 1,4-diazepanyl, or morpholinyl, each of which is optionally substituted with hydroxy, amino, aminoalkyl, C₁-C₆ alkyl, mono- or di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or halogen, and

R₆ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀ alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl; and

R₅ is

- (1) heteroaryl,
- (2) aryl,
- (3) saturated or unsaturated C₅-C₁₀ cycloalkyl, or
- (4) saturated or unsaturated C₅-C₁₀ heterocycloalkyl,

and

the R₅ groups are optionally substituted at least one group which is independently hydroxy, oxo, halo, amino, cyano, nitro, -SH,

-S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl,
-SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or
mono- or di-(C₁-C₁₀)alkylamino;

5 R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀alkyl, or
 halo(C₁-C₆)alkyl; and
 X is N or CR₃, wherein
 R₃ is H, halogen, or CH₃,

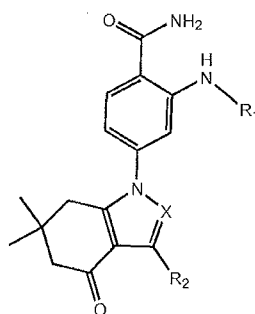
10 or a prodrug thereof, in treating a mammalian subject affected by
 prostate cancer.

Hsp90 inhibitors having structure above are described in U.S.
Patent No. 7,928,135, the entire contents of which are hereby
15 incorporated herein by reference.

The present invention provides an oligonucleotide which reduces
clusterin expression for use in combination with a Hsp90 inhibitor,
which inhibitor is other than Hsp90i-1, in treating a mammalian
20 subject affected by prostate cancer.

The present invention provides an oligonucleotide which reduces
clusterin expression for use in combination with a Hsp90 inhibitor
which binds to Hsp90 α and Hsp90 β with a K_a of less than 50 nmol/L,
25 or a prodrug thereof, in treating a mammalian subject affected by
 prostate cancer.

The present invention provides a composition for treating a
mammalian subject affected by prostate cancer comprising i) an
30 oligonucleotide which reduces clusterin expression and ii) a Hsp90
 inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

5 R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ -
10 $C_6)$ alkylamino, nitro, halo(C_1 - C_6)alkyl, halo(C_1 - C_6)alkoxy, or carboxamide, wherein

when R_1 is a C_1 - C_{14} alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R_4 , carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO_2 , or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R_4 is
20 (i) heteroaryl,
(ii) aryl,
(iii) saturated or unsaturated C_3 - C_{10} cycloalkyl, or
(iv) saturated or unsaturated C_2 - C_{10}
25 heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or unsaturated cycloalkyl, or saturated or unsaturated heterocycloalkyl,

independently, is optionally substituted with at least one group, which independently is hydroxy, halo, amino, cyano, carboxy, carboxamido, nitro, oxo, 5 -S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino; and R₄ is optionally fused to a C₆-C₁₀ aryl group, C₅-C₉ saturated cyclic group, or a C₄-C₁₀ heterocycloalkyl group; and

R₁ is optionally substituted at any available position with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo, amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂(C₁-C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀ alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅, 20 wherein

Z is OR₀ or -N(R₆)₂, wherein

each R₆ is independently -H or C₁-C₆ alkyl, or N(R₆)₂ represents pyrrolidinyl, piperidinyl, 25 piperazinyl, azepanyl, 1,3- or 1,4-diazepanyl, or morpholinyl, each of which is optionally substituted with hydroxy, amino, aminoalkyl, C₁-C₆ alkyl, mono- or di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or 30 halogen, and

R₀ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀ alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl; and

R₅ is

(1) heteroaryl,

(2) aryl,

(3) saturated or unsaturated C₅-C₁₀
cycloalkyl, or

(4) saturated or unsaturated C₅-C₁₀
heterocycloalkyl,

and

the R₅ groups are optionally substituted at
least one group which is independently
hydroxy, oxo, halo, amino, cyano, nitro, -SH,
-S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl,
-SO-(C₁-C₆) alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
(C₁-C₆) alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or
mono- or di-(C₁-C₁₀)alkylamino;

R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or
halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

R₃ is H, halogen, or CH₃,

or a prodrug thereof, each in an amount that when in combination
with the other is effective to treat the mammalian subject.

The present invention provides a composition for treating a
mammalian subject affected by prostate cancer comprising i) an
oligonucleotide which reduces clusterin expression and ii) a Hsp90
inhibitor, which inhibitor is other than Hsp90i-1, each in an
amount that when in combination with the other is effective to
treat the mammalian subject.

The present invention provides a composition for treating a
mammalian subject affected by prostate cancer comprising i) an
oligonucleotide which reduces clusterin expression and ii) a Hsp90
inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50
nmol/L, or a prodrug thereof, each in an amount that when in
combination with the other is effective to treat the mammalian
subject.

Brief Description of the Drawings

Figure 1. Hsp90i-1 and Hsp90i-2 induce HSPs and clusterin (CLU) expression in prostate cancer (PCa) cells in vitro. PC-3 and LNCaP cells were treated with 1 μ M Hsp90i-2 (A) or 1 μ M Hsp90i-1 (C) for the indicated time points. In parallel, PC-3 and LNCaP cells were treated for 48h with Hsp90i-2 for the indicated doses (B). Protein extracts were analyzed for CLU, Hsp70, Akt and vinculin. Tumor cells were treated for 24h with 1 μ M Hsp90i-2 or 1 μ M Hsp90i-1 (D). mRNA extracts were analyzed by real-time PCR for CLU, Hsp90 and Hsp70. ***, $p < 0.001$.

Figure 2. Hsp90i-2 induces HSP and CLU expression in PCa xenografts. Mice were treated for 6 weeks with 50mg/kg Hsp90i-2-PRO (the prodrug of Hsp90i-2) or vehicle (Control). A, tumors were collected and CLU and Hsp70 were evaluated by immunohistochemical analysis. B, total proteins were extracted from the xenograft tumors and CLU expression was analyzed by western blotting. The relative levels were normalized with GAPDH and estimated in densitometric units. ***, $p < 0.001$.

Figure 3. CLU induction following Hsp90 inhibitor treatment is cytoprotective via an increase of HSF-1 activity. A, LNCaP cells were treated with indicated concentrations of Hsp90i-1 or Hsp90i-2 for 48h. B, LNCaP cells were transiently transfected with indicated concentrations of CLU-plasmid for 48h. Total amount of plasmid DNA transfected was normalized to 2 μ g per well by the addition of an

empty vector. C (Top), LNCaP cells were transfected with 20nM CLU siRNA or control siScr, followed of Hsp90i-1 or Hsp90i-2 treatment (1 μ M) for 48h. C (bottom), LNCaP cells were treated twice with 300nM custirsen or control ScrB ASO. D, LNCaP and PC-3 cells were treated twice with 300nM custirsen or control ScrB ASO, followed by 1 μ M of Hsp90i-1 or Hsp90i-2 for 48h. Cells were harvested, and HSE-luciferase activity or western blotting analyses were performed. Means of at least three independent experiments done in triplicate. ***, $p < 0.001$; *, $p < 0.05$; ns, not significant.

Figure 4. Increased potency of CLU knockdown and Hsp90

inhibitor combination treatment in PCa cells. A, LNCaP cells were treated twice with 300nM custirsen or control ScrB ASO, followed by the indicated concentration of Hsp90i-1 or Hsp90i-2 for 48h. Cell growth was determined by crystal violet and compared with control. B, dose dependent effects and combination index (CI) values calculated by CalcuSyn software were assessed in LNCap cells treated for 48h with custirsen alone, Hsp90i-2 alone or combined treatment at indicated concentration with constant ratio design between both drugs. The CI for ED₅₀ and ED₇₅ was 0.4 and 0.75, respectively, indicative of a combination effect of this combined treatment. C and D, LNCaP cells were treated twice with 300nM custirsen or control ScrB, followed by 1 μ M Hsp90i-1 or Hsp90i-2 for 48h. Cells were harvested, and western blotting analyses were performed (C). The proportion of cells in subG1, G0-G1, S, G2-M was

determined by propidium iodide staining and caspase-3 activity was determined on the cell lysates and the results are expressed in arbitrary units and corrected for protein content (D). All experiments were repeated at least thrice. \$\$\$, $p < 0.001$; ***, $p < 0.001$; **, $p < 0.01$; *, $p < 0.05$.

Figure 5. Increased potency of custirsen + Hsp90i-1 combination in PC-3 xenograft model. Mice were treated IP with 25mg/kg Hsp90i-1 and 15mg/kg custirsen starting when tumors reached 300mm as described in Example 6. A, The mean tumor volume of mice custirsen + Hsp90i-1 was compared with control ScrB ASO + Hsp90i-1 $\pm$ SEM (n=7). **, $p < 0.01$. B, in Kaplan-Meier curve, cancer-specific survival was compared between mice treated with custirsen + Hsp90i-1 and control ScrB ASO + Hsp90i-1 over a 72-d period. *, $p < 0.05$. C, tumors were collected after 72-d and CLU, Ki67 and TUNEL were evaluated by immunohistochemical analysis (original magnification: x200).

Figure 6. Increased potency of custirsen + Hsp90i-2-PRO combination in LNCaP xenograft model. Mice were treated with 25mg/kg Hsp90i-2-PRO and 15mg/kg custirsen starting when serum PSA values relapsed to pre-castration levels. The mean tumor volume (A) and the serum PSA level (B) were compared between the 4 groups $\pm$ SEM (n=10). ***, $p < 0.001$. C, PSA doubling time and velocity were calculated as described in Example 6. *, $p < 0.05$. D, in Kaplan-Meier curve, cancer-specific survival was compared between the 4

groups over a 62-d period. ***, $p < 0.001$. Progression-free survival was defined as time for the first tumor volume doubling.

5 **Figure 7. Increased potency of custirsen + Hsp90i-2-PRO combination treatment apoptosis levels in CRPC LNCaP tumors.** A, tumors were collected after 57 days and CLU, Ki67, AR, AKT and TUNEL were evaluated by immunohistochemical analysis (original
10 magnification: x200). B, total proteins were extracted from the xenograft tumors and CLU, AR, Akt and PSA were analyzed by western blotting. The relative levels were normalized with vinculin and estimated in densitometric units $\pm$ SEM.

15

Figure 8. Clusterin protects tumor cells to Hsp90 inhibitors via a regulation of HSF-1. A, PC-3 cells were transfected to overexpress CLU compared to wt-PC-3 and treated with indicated concentrations of Hsp90i-
20 2 for 48h. Cell growth was determined by crystal violet and compared with control. **, $p \leq 0.01$. B, tumor cells were treated with 20nM HSF-1 siRNA vs control Scr siRNA and treated with μ M Hsp90i-2 for 48h. Total proteins were extracted and, western
25 blotting and caspase 3/7 activity were performed. C, PC-3 cells were treated with 20nM CLU siRNA vs control Scr siRNA and treated with 1 μ M Hsp90i-1 for 24h. HSF-1 localization was assessed by immunofluorescence staining.

30

Figure 9. LNCaP and PC-3 cells were treated twice with 300nM custirsen or control ScrB ASO, followed by 1 μ M of

Hsp90i-1 or Hsp90i-2 for 48h. Cells were harvested, and HSE-luciferase activity or western blotting analyses were performed.

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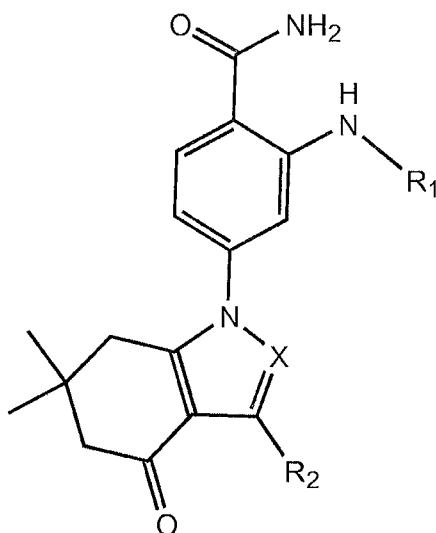
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Detailed Description of the Invention

The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin
 5 expression and ii) a Heat Shock Protein 90 (Hsp90) inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl,
 10 heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl,
 wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl,
 and heteroaryl group is optionally substituted with from
 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6
 15 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ -
 $C_6)$ alkylamino, nitro, halo $(C_1$ - $C_6)$ alkyl, halo $(C_1$ - $C_6)$ alkoxy,
 or carboxamide, wherein

when R_1 is a C_1 - C_{14} alkyl group, up to five of the
 carbon atoms in the alkyl group are optionally
 replaced independently by R_4 , carbonyl, ethenyl,
 20 ethynyl or a moiety selected from N, O, S, SO_2 , or SO,
 with the proviso that two O atoms, two S atoms, or an
 O and S atom are not immediately adjacent each other,
 wherein

R_4 is
 25 (i) heteroaryl,

- (ii) aryl,
 (iii) saturated or unsaturated C₃-C₁₀
 cycloalkyl, or
 (iv) saturated or unsaturated C₂-C₁₀
 heterocycloalkyl,

5 wherein
 each aryl, heteroaryl, saturated or
 unsaturated cycloalkyl, or saturated or
 unsaturated heterocycloalkyl,
 10 independently, is optionally substituted
 with at least one group, which
 independently is hydroxy, halo, amino,
 cyano, carboxy, carboxamido, nitro, oxo,
 -S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-
 15 aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂,
 -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-
 C₆)alkoxy, or mono- or di-(C₁-
 C₁₀)alkylamino; and R₄ is optionally fused
 to a C₈-C₁₀ aryl group, C₅-C₈ saturated
 20 cyclic group, or a C₄-C₁₀ heterocycloalkyl
 group; and

R₁ is optionally substituted at any available position
 with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-
 C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo,
 25 amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-
 C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-
 aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀
 alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-
 C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
 30 wherein

Z is OR₆ or -N(R₆)₂, wherein
 each R₆ is independently -H or C₁-C₆ alkyl, or
 N(R₆)₂ represents pyrrolidinyl, piperidinyl,
 piperazinyl, azepanyl, 1,3- or 1,4-
 35 diazepanyl, or morpholinyl, each of which

is optionally substituted with hydroxy, amino, aminoalkyl, C₁-C₆ alkyl, mono- or di(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or halogen, and

R₆ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀ alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl; and

R₅ is

(1) heteroaryl,

(2) aryl,

(3) saturated or unsaturated C₅-C₁₀ cycloalkyl, or

(4) saturated or unsaturated C₅-C₁₀ heterocycloalkyl,

and

the R₅ groups are optionally substituted at least one group which is independently hydroxy, oxo, halo, amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl, -SO-(C₁-C₆) alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆) alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino;

R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

R₃ is H, halogen, or CH₃,

or a prodrug thereof, each in an amount that when in combination with the other is effective to treat the mammalian subject.

The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor, which inhibitor is other than

Hsp90i-1, each in an amount that when in combination with the other is effective to treat the mammalian subject.

5 The present invention provides a method for treating a mammalian subject affected by prostate cancer comprising administering to the mammalian subject i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50 nmol/L, or a prodrug thereof, each in an amount that when in combination with the other is effective
10 to treat the mammalian subject.

In some embodiments, the Hsp90 inhibitor binds to Hsp90 α and/or Hsp90 β with a K_a of less than about 70, 60, 50, 40, 35, 30, 25, 20, 15, 10, or 5 nmol/L.

15 In some embodiments, the cancer is androgen-independent prostate cancer.

In some embodiments, the amount of the oligonucleotide and the
20 amount of the Hsp90 inhibitor when taken together is more effective to treat the subject than when each agent is administered alone.

In some embodiments, the amount of the oligonucleotide in combination with the amount of the Hsp90 inhibitor is less than is
25 clinically effective when administered alone.

In some embodiments, the amount of the Hsp90 inhibitor in combination with the amount of the oligonucleotide is less than is clinically effective when administered alone.

30 In some embodiments, the amount of the oligonucleotide and the amount of the Hsp90 inhibitor when taken together is effective to reduce a clinical symptom of prostate cancer in the subject.

35 In some embodiments, the mammalian subject is human.

In some embodiments, the oligonucleotide is an antisense oligonucleotide.

- 5 In some embodiments, the antisense oligonucleotide spans either the translation initiation site or the termination site of clusterin-encoding mRNA.

10 In some embodiments, the antisense oligonucleotide comprises nucleotides in the sequence set forth in SEQ ID NO: to 11.

In some embodiments, the antisense oligonucleotide comprises nucleotides in the sequence set forth in SEQ ID NO: 3.

- 15 In some embodiments, the antisense oligonucleotide is modified to enhance *in vivo* stability relative to an unmodified oligonucleotide of the same sequence.

20 In some embodiments, the oligonucleotide is custirsen.

In some embodiments, the amount of custirsen is less than 640mg.

In some embodiments, the amount of custirsen is less than 480mg.

- 25 In some embodiments, the amount of custirsen is administered intravenously once in a seven day period.

30 In some embodiments, the amount of the Hsp90 inhibitor is less than 50mg/kg.

In some embodiments, the amount of the Hsp90 inhibitor is 25mg/kg or less.

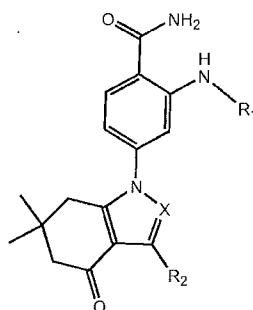
35 In some embodiments, the Hsp90 inhibitor is Hsp90i-2.

In some embodiments, a prodrug of the Hsp90 inhibitor is administered to the mammalian subject which prodrug is Hsp90i-2-PRO.

In some embodiments, a prodrug of the Hsp90 inhibitor is administered to the mammalian subject which prodrug is Hsp90i-2-PRO2.

In some embodiments, the combination of the oligonucleotide and the Hsp90 inhibitor is effective to inhibit the proliferation of prostate cancer cells.

The present invention provides a pharmaceutical composition comprising an amount of an oligonucleotide which reduces clusterin expression, and a Hsp90 inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ - $C_6)$ alkylamino, nitro, halo $(C_1$ - $C_6)$ alkyl, halo $(C_1$ - $C_6)$ alkoxy, or carboxamide, wherein

when R_1 is a C_1 - C_{14} alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R_4 , carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO_2 , or SO, with the proviso that two O atoms, two S atoms, or an

O and S atom are not immediately adjacent each other,
wherein

R₄ is

- (i) heteroaryl,
(ii) aryl,
(iii) saturated or unsaturated C₃-C₁₀
cycloalkyl, or
(iv) saturated or unsaturated C₂-C₁₀
heterocycloalkyl,

10

wherein

each aryl, heteroaryl, saturated or
unsaturated cycloalkyl, or saturated or
unsaturated heterocycloalkyl,
independently, is optionally substituted
with at least one group, which
independently is hydroxy, halo, amino,
cyano, carboxy, carboxamido, nitro, oxo,
-S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-
aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂,
-SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-
C₆)alkoxy, or mono- or di-(C₁-
C₁₀)alkylamino; and R₄ is optionally fused
to a C₆-C₁₀ aryl group, C₅-C₈ saturated
cyclic group, or a C₄-C₁₀ heterocycloalkyl
group; and

25

R₁ is optionally substituted at any available position
with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-
C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo,
amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-
C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-
aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀
alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-
C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
wherein

35

Z is OR₆ or -N(R₆)₂, wherein

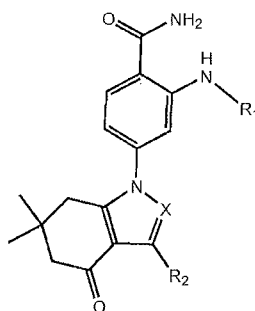
- each R_6 is independently -H or C_1-C_6 alkyl, or
 $N(R_6)_2$ represents pyrrolidinyl, piperidinyl,
piperazinyl, azepanyl, 1,3- or 1,4-
diazepanyl, or morpholinyl, each of which
is optionally substituted with hydroxy,
amino, aminoalkyl, C_1-C_6 alkyl, mono- or
di(C_1-C_6)alkylamino, C_1-C_6 alkoxy, or
halogen, and
 R_0 is -H, $-C_1-C_{10}$ alkyl, $-C_2-C_{10}$ alkenyl, $-C_2-C_{10}$
alkynyl, aryl, heteroaryl, or $-C_1-C_6$ acyl;
and
 R_5 is
(1) heteroaryl,
(2) aryl,
(3) saturated or unsaturated C_5-C_{10}
cycloalkyl, or
(4) saturated or unsaturated C_5-C_{10}
heterocycloalkyl,
and
the R_5 groups are optionally substituted at
least one group which is independently
hydroxy, oxo, halo, amino, cyano, nitro, -SH,
-S-(C_1-C_6)alkyl, -SO₂-(C_1-C_6)alkyl, -SO₂-aryl,
-SO-(C_1-C_6) alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
(C_1-C_6) alkyl, -SO₂NH-aryl, (C_1-C_6)alkoxy, or
mono- or di-(C_1-C_{10})alkylamino;
 R_2 is H, Cl, halogen, CF₃, CHF₂, CH₃, C_1-C_{10} alkyl, or
halo(C_1-C_6)alkyl; and
X is N or CR₃, wherein
 R_3 is H, halogen, or CH₃,

or a prodrug thereof, for use in treating a mammalian subject
affected by prostate cancer.

The present invention provides a pharmaceutical composition comprising an amount of an oligonucleotide which reduces clusterin expression, and a Hsp90 inhibitor, which inhibitor is other than Hsp90i-1, for use in treating a mammalian subject affected by prostate cancer.

The present invention provides a pharmaceutical composition comprising an amount of an oligonucleotide which reduces clusterin expression, and a Hsp90 inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50 nmol/L, or a prodrug thereof, for use in treating a mammalian subject affected by prostate cancer.

The present invention provides an oligonucleotide which reduces clusterin expression for use in combination with a Hsp90 inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ - $C_6)$ alkylamino, nitro, halo(C_1 - C_6)alkyl, halo(C_1 - C_6)alkoxy, or carboxamide, wherein

when R_1 is a C_1 - C_{14} alkyl group, up to five of the carbon atoms in the alkyl group are optionally

replaced independently by R_4 , carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO_2 , or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R_4 is

- (i) heteroaryl,
- (ii) aryl,
- (iii) saturated or unsaturated C_3-C_{10} cycloalkyl, or
- (iv) saturated or unsaturated C_2-C_{10} heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or unsaturated cycloalkyl, or saturated or unsaturated heterocycloalkyl, independently, is optionally substituted with at least one group, which independently is hydroxy, halo, amino, cyano, carboxy, carboxamido, nitro, oxo, $-S-(C_1-C_6)$ alkyl, $-SO_2(C_1-C_6)$ alkyl, $-SO_2$ -aryl, $-SO-(C_1-C_6)$ alkyl, $-SO$ -aryl, $-SO_2NH_2$, $-SO_2NH-(C_1-C_6)$ alkyl, $-SO_2NH$ -aryl, (C_1-C_6) alkoxy, or mono- or di- (C_1-C_{10}) alkylamino; and R_4 is optionally fused to a C_6-C_{10} aryl group, C_5-C_8 saturated cyclic group, or a C_4-C_{10} heterocycloalkyl group; and

R_1 is optionally substituted at any available position with C_1-C_{10} alkyl, C_1-C_{10} haloalkyl, C_2-C_{10} alkenyl, C_2-C_{10} alkynyl, hydroxy, carboxy, carboxamido, oxo, halo, amino, cyano, nitro, $-SH$, $-S-(C_1-C_6)$ alkyl, $-SO_2-(C_1-C_6)$ alkyl, $-SO_2NH_2$, $-SO_2NH(C_1-C_6)$ alkyl, $-SO_2NH$ -aryl, $-SO_2$ -aryl, $-SO-(C_1-C_6)$ alkyl, $-SO_2$ -aryl, C_1-C_6 alkoxy, C_2-C_{10} alkenyloxy, C_2-C_{10} alkynyloxy, mono- or di- $(C_1-$

C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
wherein

Z is OR₆ or -N(R₆)₂, wherein

5 each R₆ is independently -H or C₁-C₆ alkyl, or
N(R₆)₂ represents pyrrolidinyl, piperidinyl,
piperazinyl, azepanyl, 1,3- or 1,4-
diazepanyl, or morpholinyl, each of which
is optionally substituted with hydroxy,
amino, aminoalkyl, C₁-C₆ alkyl, mono- or
10 di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or
halogen, and

R₆ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀
alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl;
and

15 R₅ is

- (1) heteroaryl,
- (2) aryl,
- (3) saturated or unsaturated C₅-C₁₀
cycloalkyl, or
- 20 (4) saturated or unsaturated C₅-C₁₀
heterocycloalkyl,

and

the R₅ groups are optionally substituted at
least one group which is independently
25 hydroxy, oxo, halo, amino, cyano, nitro, -SH,
-S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl,
-SO-(C₁-C₆) alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
(C₁-C₆) alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or
mono- or di-(C₁-C₁₀)alkylamino;

30 R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or
halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

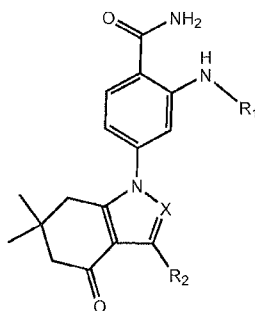
R₃ is H, halogen, or CH₃,

or a prodrug thereof, in treating a mammalian subject affected by prostate cancer.

The present invention provides an oligonucleotide which reduces clusterin expression for use in combination with a Hsp90 inhibitor, which inhibitor is other than Hsp90i-1, in treating a mammalian subject affected by prostate cancer.

The present invention provides an oligonucleotide which reduces clusterin expression for use in combination with a Hsp90 inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50 nmol/L, or a prodrug thereof, in treating a mammalian subject affected by prostate cancer.

The present invention provides a composition for treating a mammalian subject affected by prostate cancer comprising i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor having the structure:



or a pharmaceutically acceptable salt thereof, wherein

R_1 is H, C_1 - C_{14} alkyl, C_1 - C_{10} haloalkyl, C_3 - C_7 cycloalkyl, heterocycloalkyl, C_1 - C_6 acyl, aryl, or heteroaryl, wherein each alkyl, cycloalkyl, heterocycloalkyl, aryl, and heteroaryl group is optionally substituted with from 1-4 groups that are independently C_1 - C_6 alkyl, C_1 - C_6 alkoxy, halogen, hydroxy, amino, mono- or di- $(C_1$ -

C₆)alkylamino, nitro, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, or carboxamide, wherein

when R₁ is a C₁-C₁₄ alkyl group, up to five of the carbon atoms in the alkyl group are optionally replaced independently by R₄, carbonyl, ethenyl, ethynyl or a moiety selected from N, O, S, SO₂, or SO, with the proviso that two O atoms, two S atoms, or an O and S atom are not immediately adjacent each other, wherein

R₄ is

(i) heteroaryl,
 (ii) aryl,
 (iii) saturated or unsaturated C₃-C₁₀ cycloalkyl, or

(iv) saturated or unsaturated C₂-C₁₀ heterocycloalkyl,

wherein

each aryl, heteroaryl, saturated or unsaturated cycloalkyl, or saturated or unsaturated heterocycloalkyl,

independently, is optionally substituted with at least one group, which independently is hydroxy, halo, amino, cyano, carboxy, carboxamido, nitro, oxo,

-S-(C₁-C₆) alkyl, -SO₂(C₁-C₆) alkyl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-(C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or mono- or di-(C₁-C₁₀)alkylamino; and R₄ is optionally fused

to a C₆-C₁₀ aryl group, C₅-C₉ saturated cyclic group, or a C₄-C₁₀ heterocycloalkyl group; and

R₁ is optionally substituted at any available position with C₁-C₁₀ alkyl, C₁-C₁₀ haloalkyl, C₂-C₁₀ alkenyl, C₂-C₁₀ alkynyl, hydroxy, carboxy, carboxamido, oxo, halo,

amino, cyano, nitro, -SH, -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂NH-aryl, -SO₂-aryl, -SO-(C₁-C₆)alkyl, -SO₂-aryl, C₁-C₆ alkoxy, C₂-C₁₀ alkenyloxy, C₂-C₁₀ alkynyloxy, mono- or di-(C₁-C₁₀)alkylamino, -C₁-C₁₀ alkyl-Z, -OC₁-C₁₀ alkyl-Z, or R₅,
 5 wherein

Z is OR₀ or -N(R₆)₂, wherein

each R₆ is independently -H or C₁-C₆ alkyl, or
 N(R₆)₂ represents pyrrolidinyl, piperidinyl,
 10 piperazinyl, azepanyl, 1,3- or 1,4-diazepanyl, or morpholinyl, each of which
 is optionally substituted with hydroxy,
 amino, aminoalkyl, C₁-C₆ alkyl, mono- or
 di-(C₁-C₆)alkylamino, C₁-C₆ alkoxy, or
 15 halogen, and

R₀ is -H, -C₁-C₁₀ alkyl, -C₂-C₁₀ alkenyl, -C₂-C₁₀
 alkynyl, aryl, heteroaryl, or -C₁-C₆ acyl;
 and

R₅ is

- 20 (1) heteroaryl,
 - (2) aryl,
 - (3) saturated or unsaturated C₅-C₁₀
cycloalkyl, or
 - (4) saturated or unsaturated C₅-C₁₀
 25 heterocycloalkyl,
- and

the R₅ groups are optionally substituted at
 least one group which is independently
 hydroxy, oxo, halo, amino, cyano, nitro, -SH,
 30 -S-(C₁-C₆)alkyl, -SO₂-(C₁-C₆)alkyl, -SO₂-aryl,
 -SO-(C₁-C₆)alkyl, -SO-aryl, -SO₂NH₂, -SO₂NH-
 (C₁-C₆)alkyl, -SO₂NH-aryl, (C₁-C₆)alkoxy, or
 mono- or di-(C₁-C₁₀)alkylamino;

R₂ is H, Cl, halogen, CF₃, CHF₂, CH₃, C₁-C₁₀ alkyl, or
 35 halo(C₁-C₆)alkyl; and

X is N or CR₃, wherein

R₃ is H, halogen, or CH₃,

or a prodrug thereof, each in an amount that when in combination
5 with the other is effective to treat the mammalian subject.

The present invention provides a composition for treating a
mammalian subject affected by prostate cancer comprising i) an
oligonucleotide which reduces clusterin expression and ii) a Hsp90
10 inhibitor, which inhibitor is other than Hsp90i-1, each in an
amount that when in combination with the other is effective to
treat the mammalian subject.

The present invention provides a composition for treating a
15 mammalian subject affected by prostate cancer comprising i) an
oligonucleotide which reduces clusterin expression and ii) a Hsp90
inhibitor which binds to Hsp90 α and Hsp90 β with a K_a of less than 50
nmol/L, or a prodrug thereof, each in an amount that when in
combination with the other is effective to treat the mammalian
20 subject.

In some embodiments, the combination of the oligonucleotide and the
Hsp90 inhibitor is effective to inhibit the proliferation of
prostate cancer cells.

25

In some embodiments, Hsp90 inhibitor-mediated induction of
clusterin expression is attenuated by custirsen, wherein the
combination of the Hsp90 inhibitor and custirsen delays the
progression of CRPC. In some embodiments, the combination of the
30 Hsp90 inhibitor and custirsen inhibits tumor growth in the
mammalian subject. In some embodiments, the combination of the
Hsp90 inhibitor and custirsen prolongs the survival of the
mammalian subject.

An aspect of the invention provides pharmaceutical composition comprising an amount of an oligonucleotide which reduces clusterin expression, and a Hsp90 inhibitor for use in treating a mammalian subject affected by prostate cancer.

5

An aspect of the invention provides oligonucleotide which reduces clusterin expression for use in combination with a Hsp90 inhibitor in treating a mammalian subject affected by prostate cancer.

10 An aspect of the invention provides a composition for treating a mammalian subject affected by prostate cancer comprising i) an oligonucleotide which reduces clusterin expression and ii) a Hsp90 inhibitor each in an amount that when in combination with the other is effective to treat the mammalian subject.

15

Aspects of the invention involve the increased potency of a combination treatment comprising an oligonucleotide that targets clusterin expression and an Hsp90 inhibitor compared to oligonucleotide or Hsp90 inhibitor monotherapy. In some embodiments
20 of the invention, the combination of an oligonucleotide that targets clusterin expression and an HSP90 inhibitor increases prostate cancer cell apoptosis and/or decreases prostate cancer cell proliferation compared to oligonucleotide or Hsp90 inhibitor monotherapy. In some embodiments, the combination of an
25 oligonucleotide that targets clusterin expression and an Hsp90 inhibitor decreases the protein expression and/or a function of HSF-1 compared to oligonucleotide or Hsp90 inhibitor monotherapy.

Aspects of the invention provide targeted strategies employing an
30 oligonucleotide which reduces clusterin expression in combination with Hsp90 inhibitors to improve patient outcome in castration-resistant prostate cancer.

The present invention relates to a method for treating a mammalian
35 subject affected by prostate cancer comprising i) an

oligonucleotide which reduces clusterin expression and ii) a Heat Shock Protein 90 (Hsp90) inhibitor, each in an amount that when in combination with the other is effective to treat the mammalian subject.

5

In some embodiments, the Hsp90 inhibitor is Hsp90i-1.

Each embodiment disclosed herein is contemplated as being applicable to each of the other disclosed embodiments. Thus, all combinations of the various elements described herein are within the scope of the invention.

It is understood that where a parameter range is provided, all integers within that range, and tenths thereof, are also provided by the invention. For example, "0.2-5 mg/kg/day" includes 0.2 mg/kg/day, 0.3 mg/kg/day, 0.4 mg/kg/day, 0.5 mg/kg/day, 0.6 mg/kg/day etc. up to 5.0 mg/kg/day.

Terms

As used herein, and unless stated otherwise, each of the following terms shall have the definition set forth below.

As used herein, "about" in the context of a numerical value or range means $\pm 10\%$ of the numerical value or range recited or claimed.

As used in the specification and claims of this application, the term "clusterin" refers to a glycoprotein present in mammals, including humans, and denominated as such in the humans. The sequences of numerous clusterin species are known. For example, the sequence of human clusterin is described by Wong et al., Eur. J. Biochem. 221 (3), 917-925 (1994), and in NCBI sequence accession number NM_001831 (SEQ ID NO: 43). In this human sequence, the coding sequence spans bases 48 to 1397.

35

As used herein, "oligonucleotide which reduces clusterin expression" is an oligonucleotide with a sequence which is effective to reduce clusterin expression in a cell. The oligonucleotide which reduces clusterin expression may be, for example, an antisense oligonucleotide or an RNA interference inducing molecule.

As used herein, "antisense oligonucleotide" refers to a non-RNAi oligonucleotide that reduces clusterin expression and that has a sequence complementary to clusterin mRNA. Antisense oligonucleotides may be antisense oligodeoxynucleotides (ODN). Exemplary sequences which can be employed as antisense molecules in the invention are disclosed in PCT Patent Publication WO 00/49937, U.S. Patent Publication US-2002-0128220-A1, and U.S. Pat. No. 6,383,808, all of which are incorporated herein by reference. Specific antisense sequences are set forth in the present application as SEQ ID NOs: 1 to 18, and may be found in Table 1.

Table 1. Sequence Identification Numbers for Antisense Oligonucleotides

SEQ ID NO:	Sequence
1	GCACAGCAGG AGAATCTTCA T
2	TGGAGTCTTT GCACGCCTCG G
3	CAGCAGCAGA GTCTTCATCA T
4	ATTGTCTGAG ACCGTCTGGT C
5	CCTTCAGCTT TGTCTCTGAT T
6	AGCAGGGAGT CGATGCGGTC A
7	ATCAAGCTGC GGACGATGCG G
8	GCAGGCAGCC CGTGGAGTTG T
9	TTCAGCTGCT CCAGCAAGGA G
10	AATTTAGGGT TCTTCCTGGA G
11	GCTGGGCGGA GTTGGGGGCC T
12	GGTGTAGACG CCGCACG
13	GCAGCGCAGC CCCTGG
14	GCAGCAGCCG CAGCCCGGCT CC

15	AGCCGCAGCC CGGCTCCT
16	CAGCAGCCGC AGCCCGGCTC
17	CAGCAGCCGC AGCCCGGCTC
18	AGCAGCCGCA GCCCGGCTCC

The ODNs employed may be modified to increase the stability of the ODN *in vivo*. For example, the ODNs may be employed as phosphorothioate derivatives (replacement of a non-bridging phosphoryl oxygen atom with a sulfur atom) which have increased resistance to nuclease digestion. MOE (2'-O-(2-methoxyethyl)) modification (ISIS backbone) is also effective. The construction of such modified ODNs is described in detail in U.S. Patent No. 6,900,187 B2, the contents of which are incorporated by reference. In some embodiments, the ODN is custirsén.

As used herein, "custirsén" refers to an antisense oligonucleotide that reduces clusterin expression having the sequence CAGCAGCAGAGTCTTCATCAT (SEQ ID NO: 3), wherein the anti-clusterin oligonucleotide has a phosphorothioate backbone throughout, has sugar moieties of nucleotides 1-4 and 18-21 bearing 2'-O-methoxyethyl modifications, has nucleotides 5-17 which are 2'-deoxynucleotides, and has 5-methylcytosines at nucleotides 1, 4, and 19. Custirsén is also known as TV-1011, OGX-011, ISIS 112989 and Custirsén Sodium.

As used herein, "RNA inducing molecule" refers to a molecule capable of inducing RNA interference or "RNAi" of clusterin expression. RNAi involves mRNA degradation, but many of the biochemical mechanisms underlying this interference are unknown. The use of RNAi has been described in Fire et al., 1998, Carthew et al., 2001, and Elbashir et al., 2001, the contents of which are incorporated herein by reference.

Isolated RNA molecules can mediate RNAi. That is, the isolated RNA molecules of the present invention mediate degradation or block expression of mRNA that is the transcriptional product of the gene,

which is also referred to as a target gene. For convenience, such mRNA may also be referred to herein as mRNA to be degraded. The terms RNA, RNA molecule(s), RNA segment(s) and RNA fragment(s) may be used interchangeably to refer to RNA that mediates RNA interference. These terms include double-stranded RNA, small interfering RNA (siRNA), hairpin RNA, single-stranded RNA, isolated RNA (partially purified RNA, essentially pure RNA, synthetic RNA, recombinantly produced RNA), as well as altered RNA that differs from naturally occurring RNA by the addition, deletion, substitution and/or alteration of one or more nucleotides. Such alterations can include addition of non-nucleotide material, such as to the end(s) of the RNA or internally (at one or more nucleotides of the RNA). Nucleotides in the RNA molecules of the present invention can also comprise nonstandard nucleotides, including non-naturally occurring nucleotides or deoxyribonucleotides. Collectively, all such altered RNAi molecules are referred to as analogs or analogs of naturally-occurring RNA. RNA of the present invention need only be sufficiently similar to natural RNA that it has the ability to mediate RNAi.

As used herein the phrase "mediate RNAi" refers to and indicates the ability to distinguish which mRNA molecules are to be affected by the RNAi machinery or process. RNA that mediates RNAi interacts with the RNAi machinery such that it directs the machinery to degrade particular mRNAs or to otherwise reduce the expression of the target protein. In one embodiment, the present invention relates to RNA molecules that direct cleavage of specific mRNA to which their sequence corresponds. It is not necessary that there be perfect correspondence of the sequences, but the correspondence must be sufficient to enable the RNA to direct RNAi inhibition by cleavage or blocking expression of the target mRNA.

As noted above, the RNA molecules of the present invention in general comprise an RNA portion and some additional portion, for example a deoxyribonucleotide portion. The total number of

nucleotides in the RNA molecule is suitably less than in order to be effective mediators of RNAi. In preferred RNA molecules, the number of nucleotides is 16 to 29, more preferably 18 to 23, and most preferably 21-23. Suitable sequences are set forth in the present application as SEQ ID NOs:19 to 42 (Table 2).

Table 2. Sequence Identification Numbers for RNA Interference Inducing Molecules

SEQ ID NO:	Sequence
19	CCAGAGCUCG CCCUUCUACT T
20	GUAGAAGGGC GAGCUCUGGT T
21	GAUGCUC AAC ACCUCCUCCT T
22	GGAGGAGGUG UUGAGCAUCT T
23	UAAUUCAACA AAACUGUTT
24	GACAGUUUUA UUGAAUUAGT T
25	UAAUUCAACA AAACUGUTT
26	ACAGUUUUGU UGAUUUATT
27	AUGAUGAAGA CUCUGCUGCT T
28	GCAGCAGAGU CUUCAUCAUT T
29	UGAAUGAAGG GACUAACCUG TT
30	CAGGUUAGUC CCUUCAUUA TT
31	CAGAAUAGA CAAAGUGGGG TT
32	CCCCACUUUG UCUUUUCUG TT
33	ACAGAGACUA AGGGACCAGA TT
34	ACAGAGACUA AGGGACCAGA TT
35	CCAGAGCUCG CCCUUCUACT T
36	GUAGAAGGGC GAGCUCUGGT T
37	GUCCCGCAUC GUCCGCAGCT T
38	GCUGCGGACG AUGCGGGACT T
39	CUAAUCAAU AAAACUGUCT T
40	GACAGUUUUA UUGAAUUAGT T
41	AUGAUGAAGA CUCUGCUGC
42	GCAGCAGAGU CUUCAUCAU

The siRNA molecules of the invention are used in therapy to treat patients, including human patients, that have cancers or other diseases of a type where a therapeutic benefit is obtained by the inhibition of expression of the targeted protein. siRNA molecules of the invention are administered to patients by one or more daily injections (intravenous, subcutaneous or intrathecal) or by continuous intravenous or intrathecal administration for one or more treatment cycles to reach plasma and tissue concentrations suitable for the regulation of the targeted mRNA and protein.

10

As used herein, a "mammalian subject affected by prostate cancer" means a mammalian subject who has been affirmatively diagnosed to have prostate cancer.

15

As used herein, "androgen-independent prostate cancer" encompasses cells and tumors containing cells that are not androgen-dependent (not androgen sensitive); often such cells progress from being androgen-dependent to being androgen-independent. In some embodiments, androgen independent prostate cancer has progressed since the administration of hormone ablation therapy and/or hormone blockade therapy. In some embodiments, there is increased AR expression in the androgen-independent prostate cancer compared to prostate cancer that is not androgen-independent.

20

25

As used herein, "castration-resistant prostate cancer" encompasses any androgen-independent prostate cancer that is resistant to hormone ablation therapy and/or hormone blockade therapy. In some embodiments, castration-resistant prostate cancer has progressed since the administration of hormone ablation or hormone blockade therapy. In some embodiments, there is increased AR expression in the castration-resistant prostate cancer compared to prostate cancer that is not castration resistant.

30

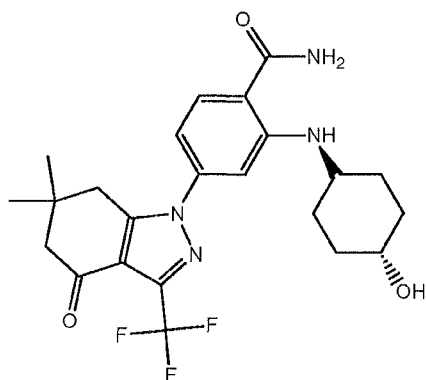
As used herein, "Hsp90 inhibitor" refers to an agent that perturbs or reduces a function of Hsp90, including inhibiting a Hsp90-protein

35

interaction, Hsp90 signaling, or Hsp90 protein expression. Hsp90 inhibitors include but are not limited to Hsp90-specific monoclonal antibodies, oligonucleotides that target Hsp90 expression (such as Hsp90 targeting antisense oligonucleotides or RNA inducing molecules), peptide agents specific for Hsp90, and small molecule inhibitors specific for Hsp90. Non-limiting examples of Hsp90 inhibitors are Hsp90i-1, Hsp90i-2, Hsp90i-2-PRO and Hsp90i-2-PRO2.

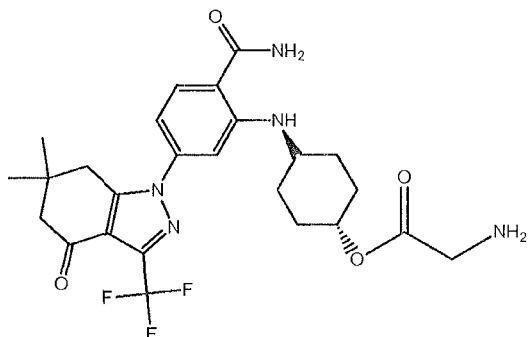
Hsp90i-1 is a Hsp90 inhibitor. Hsp90i-1 is also known as 17-allylamino-17-demethoxygeldanamycin (17-AAG), Telatinib, Tanespimycin, NSC-330507, CNF-101, KOS-953, GLD-36, and CP 127374. The CAS Registry Number of Hsp90i-1 is 75747-14-7. The Hsp90i-1 used for the experiments described herein is also referred to as 17-AAG and was obtained from the National Institutes of Health (Bethesda, Maryland, USA). 17-AAG has been discussed in Egorin et al., 1998, and Koga et al., 2009, and is also available for purchase from Invivogen (San Diego, California, USA).

Hsp90i-2 is a Hsp90 inhibitor. Hsp90i-2 is also known as PF-04928473, and SNX-2112, and (4-(6,6-Dimethyl-4-oxo-3-trifluoromethyl-4,5,6,7-tetrahydro-indazol-1-yl)-2-(4-hydroxy-cyclohexylamino)-benzamide). The CAS Registry No. for Hsp90i-2 is 908112-43-6. Hsp90i-2 has the following structure:



Hsp90i-2 is discussed in Lamoureux et al., 2011, the entire contents of which are incorporated herein by reference.

Hsp90i-2-PRO is a Hsp90 inhibitor. Hsp90i-2-PRO is the prodrug of Hsp90i-2. The CAS Registry No. for Hsp90i-2-PRO is 908115-27-5. Hsp90i-2-PRO is also known as SNX-5422 and PF-04929113. Hsp90i-2-PRO has the following structure:



Hsp90i-2-PRO is discussed in Lamoureux et al., 2011, the entire contents of which are incorporated herein by reference.

Hsp90i-2-PRO2 is another prodrug of Hsp90i-2. Hsp90i-2-PRO2 is discussed in Chandarlapaty et al., 2008, the entire contents of which are incorporated herein by reference. Hsp90i-2-PRO2 is also known as SNX-5542.

Methods of synthesis for Hsp90i-2, Hsp90i-2-PRO and Hsp90i-2-PRO2 are described in Huang et al., *J. Med Chem.* 52:4288-4305 (2009), and U.S. Patent No. 7,928,135, the entire contents of which are incorporated herein by reference. Alternatively, Hsp90i-2, Hsp90i-2-PRO and Hsp90i-2-PRO2 are available from Pfizer Inc. (New York, New York, USA) and Serenex Inc. (Durham, North Carolina, USA).

Those having ordinary skill in the art of organic synthesis will appreciate that modifications to the general procedures shown in the synthesis schemes of this application can be made to yield structurally diverse compounds. For example, where aryl rings are present, all positional isomers are contemplated and may be synthesized using standard aromatic substitution chemistry. The number and types of substituents may also vary around the aryl rings. Furthermore, where alkyl groups are present, the chain length may be

modified using methods well known to those of ordinary skill in the art. Where ester formation is contemplated, lactones may be used wherein the lactone ring is opened by reaction with a nucleophile, such as an ether-containing moiety described hereinabove. Suitable organic transformations are described in March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure (Wiley-Interscience; 6th edition, 2007), the content of which is hereby incorporated by reference.

- 10 Compounds of the subject invention can be converted to prodrugs to optimize absorption and bioavailability. Formation of a prodrug include, but is not limited to, reaction of a free hydroxyl group with a carboxylic acid to form an ester, reaction of a free hydroxyl group with an phosphorus oxychloride followed by hydrolysis to form a
15 phosphate, or reaction of a free hydroxyl group with an amino acid to form an amino acid ester, the process of which has been described previously by Chandran in WO 2005/046575. The substituents are chosen and resulting analogs are evaluated according to principles well known in the art of medicinal and pharmaceutical chemistry, such as
20 quantification of structure-activity relationships, optimization of biological activity and ADMET (absorption, distribution, metabolism, excretion, and toxicity) properties.

- Except where otherwise specified, when the structure of a compound
25 of this invention includes an asymmetric carbon atom, it is understood that the compound occurs as a racemate, racemic mixture, and isolated single enantiomer. All such isomeric forms of these compounds are expressly included in this invention. Except where otherwise specified, each stereogenic carbon may be of the R or S
30 configuration. It is to be understood accordingly that the isomers arising from such asymmetry (e.g., all enantiomers and diastereomers) are included within the scope of this invention, unless indicated otherwise. Such isomers can be obtained in substantially pure form by classical separation techniques and by
35 stereochemically controlled synthesis, such as those described in

"Enantiomers, Racemates and Resolutions" by J. Jacques, A. Collet and S. Wilen, Pub. John Wiley & Sons, NY, 1981. For example, the resolution may be carried out by preparative chromatography on a chiral column.

5

The subject invention is also intended to include all isotopes of atoms occurring on the compounds disclosed herein. Isotopes include those atoms having the same atomic number but different mass numbers. By way of general example and without limitation, isotopes of hydrogen include tritium and deuterium. Isotopes of carbon include C-13 and C-14.

It will be noted that any notation of a carbon in structures throughout this application, when used without further notation, are intended to represent all isotopes of carbon, such as ^{12}C , ^{13}C , or ^{14}C . Furthermore, any compounds containing ^{13}C or ^{14}C may specifically have the structure of any of the compounds disclosed herein.

It will also be noted that any notation of a hydrogen in structures throughout this application, when used without further notation, are intended to represent all isotopes of hydrogen, such as ^1H , ^2H , or ^3H . Furthermore, any compounds containing ^2H or ^3H may specifically have the structure of any of the compounds disclosed herein.

Isotopically-labeled compounds can generally be prepared by conventional techniques known to those skilled in the art using appropriate isotopically-labeled reagents in place of the non-labeled reagents employed.

As used herein, "alkyl" includes both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms and may be unsubstituted or substituted. Thus, $\text{C}_1\text{-C}_n$ as in " $\text{C}_1\text{-C}_n$ alkyl" is defined to include groups having 1, 2, ...,

n-1 or n carbons in a linear or branched arrangement. For example, C₁-C₆, as in "C₁-C₆ alkyl" is defined to include groups having 1, 2, 3, 4, 5, or 6 carbons in a linear or branched arrangement, and specifically includes methyl, ethyl, n-propyl, isopropyl, n-butyl, 5 t-butyl, pentyl, hexyl, and octyl.

As used herein, "aryl" is intended to mean any stable monocyclic, bicyclic or polycyclic carbon ring of up to 10 atoms in each ring, wherein at least one ring is aromatic, and may be unsubstituted or 10 substituted. Examples of such aryl elements include phenyl, p-toluenyl (4-methylphenyl), naphthyl, tetrahydro-naphthyl, indanyl, phenanthryl, anthryl or acenaphthyl. In cases where the aryl substituent is bicyclic and one ring is non-aromatic, it is understood that attachment is via the aromatic ring.

15 The term "heteroaryl", as used herein, represents a stable monocyclic, bicyclic or polycyclic ring of up to 10 atoms in each ring, wherein at least one ring is aromatic and contains from 1 to 4 heteroatoms selected from the group consisting of O, N and S. 20 Bicyclic aromatic heteroaryl groups include but are not limited to phenyl, pyridine, pyrimidine or pyridazine rings that are (a) fused to a 6-membered aromatic (unsaturated) heterocyclic ring having one nitrogen atom; (b) fused to a 5- or 6-membered aromatic (unsaturated) heterocyclic ring having two nitrogen atoms; (c) 25 fused to a 5-membered aromatic (unsaturated) heterocyclic ring having one nitrogen atom together with either one oxygen or one sulfur atom; or (d) fused to a 5-membered aromatic (unsaturated) heterocyclic ring having one heteroatom selected from O, N or S. Heteroaryl groups within the scope of this definition include but 30 are not limited to: benzoimidazolyl, benzofuranyl, benzofurazanyl, benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl, carbolinyl, cinnolinyl, furanyl, indolinyl, indolyl, indolaziny, indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl, naphthpyridinyl, oxadiazolyl, oxazolyl, 35 oxazoline, isoxazoline, oxetanyl, pyranyl, pyrazinyl, pyrazolyl,

pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl,
pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrazolyl,
tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl,
azetidiny, aziridinyl, 1,4-dioxanyl, hexahydroazepinyl,
5 dihydrobenzoimidazolyl, dihydrobenzofuranyl,
dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl,
dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl,
dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl,
10 dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl,
dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,
dihydrothienyl, dihydrotriazolyl, dihydroazetidiny,
methylenedioxybenzoyl, tetrahydrofuranyl, tetrahydrothienyl,
acridinyl, carbazolyl, cinnolinyl, quinoxalinyl, pyrrazolyl,
15 indolyl, benzotriazolyl, benzothiazolyl, benzoxazolyl, isoxazolyl,
isothiazolyl, furanyl, thienyl, benzothienyl, benzofuranyl,
quinolinyl, isoquinolinyl, oxazolyl, isoxazolyl, indolyl,
pyrazinyl, pyridazinyl, pyridinyl, pyrimidinyl, pyrrolyl, tetra-
hydroquinoline. In cases where the heteroaryl substituent is
20 bicyclic and one ring is non-aromatic or contains no heteroatoms,
it is understood that attachment is via the aromatic ring or via
the heteroatom containing ring, respectively. If the heteroaryl
contains nitrogen atoms, it is understood that the corresponding N-
oxides thereof are also encompassed by this definition.

25

The alkyl, aryl, and heteroaryl substituents may be substituted or unsubstituted, unless specifically defined otherwise.

The compounds of the instant invention may be in a salt form. As used
30 herein, a "salt" is a salt of the instant compounds which has been
modified by making acid or base salts of the compounds. In some
embodiments, the salt is pharmaceutically acceptable. Examples of
pharmaceutically acceptable salts include, but are not limited to,
mineral or organic acid salts of basic residues such as amines;
35 alkali or organic salts of acidic residues such as phenols. The salts

can be made using an organic or inorganic acid. Such acid salts are chlorides, bromides, sulfates, nitrates, phosphates, sulfonates, formates, tartrates, maleates, malates, citrates, benzoates, salicylates, ascorbates, and the like. Phenolate salts are the alkaline earth metal salts, sodium, potassium or lithium. The term "pharmaceutically acceptable salt" in this respect, refers to the relatively non-toxic, inorganic and organic acid or base addition salts of compounds of the present invention. These salts can be prepared *in situ* during the final isolation and purification of the compounds of the invention, or by separately reacting a purified compound of the invention in its free base or free acid form with a suitable organic or inorganic acid or base, and isolating the salt thus formed. Representative salts include the hydrobromide, hydrochloride, sulfate, bisulfate, phosphate, nitrate, acetate, valerate, oleate, palmitate, stearate, laurate, benzoate, lactate, phosphate, tosylate, citrate, maleate, fumarate, succinate, tartrate, naphthylate, mesylate, glucoheptonate, lactobionate, and laurylsulphonate salts and the like. (See, e.g., Berge et al. (1977) "Pharmaceutical Salts", J. Pharm. Sci. 66:1-19).

The compositions of this invention may be administered in various forms, including those detailed herein. The treatment with the compound may be a component of a combination therapy or an adjunct therapy, i.e. the subject or patient in need of the drug is treated or given another drug for the disease in conjunction with one or more of the instant compounds. This combination therapy can be sequential therapy where the patient is treated first with one drug and then the other or the two drugs are given simultaneously. These can be administered independently by the same route or by two or more different routes of administration depending on the dosage forms employed.

As used herein, a "pharmaceutically acceptable carrier" is a pharmaceutically acceptable solvent, suspending agent or vehicle, for delivering the instant compounds to the animal or human. The carrier

may be liquid or solid and is selected with the planned manner of administration in mind. Liposomes are also a pharmaceutically acceptable carrier.

- 5 The dosage of the compounds administered in treatment will vary depending upon factors such as the pharmacodynamic characteristics of a specific chemotherapeutic agent and its mode and route of administration; the age, sex, metabolic rate, absorptive efficiency, health and weight of the recipient; the nature and extent of the
10 symptoms; the kind of concurrent treatment being administered; the frequency of treatment with; and the desired therapeutic effect.

A dosage unit of the compounds may comprise a single compound or mixtures thereof with additional anticancer agents. The compounds can
15 be administered in oral dosage forms as tablets, capsules, pills, powders, granules, elixirs, tinctures, suspensions, syrups, and emulsions. The compounds may also be administered in intravenous (bolus or infusion), intraperitoneal, subcutaneous, or intramuscular form, or introduced directly, e.g. by injection, topical application,
20 or other methods, all using dosage forms well known to those of ordinary skill in the pharmaceutical arts.

The compounds can be administered in admixture with suitable pharmaceutical diluents, extenders, excipients, or carriers
25 (collectively referred to herein as a pharmaceutically acceptable carrier) suitably selected with respect to the intended form of administration and as consistent with conventional pharmaceutical practices. The unit will be in a form suitable for oral, rectal, topical, intravenous or direct injection or parenteral
30 administration. The compounds can be administered alone or mixed with a pharmaceutically acceptable carrier. This carrier can be a solid or liquid, and the type of carrier is generally chosen based on the type of administration being used. The active agent can be co-administered in the form of a tablet or capsule, liposome, as an agglomerated
35 powder or in a liquid form. Examples of suitable solid carriers

include lactose, sucrose, gelatin and agar. Capsule or tablets can be easily formulated and can be made easy to swallow or chew; other solid forms include granules, and bulk powders. Tablets may contain suitable binders, lubricants, diluents, disintegrating agents, coloring agents, flavoring agents, flow-inducing agents, and melting agents. Examples of suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent granules. Such liquid dosage forms may contain, for example, suitable solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents. Oral dosage forms optionally contain flavorants and coloring agents. Parenteral and intravenous forms may also include minerals and other materials to make them compatible with the type of injection or delivery system chosen.

Techniques and compositions for making dosage forms useful in the present invention are described in the following references: 7 Modern Pharmaceutics, Chapters 9 and 10 (Banker & Rhodes, Editors, 1979); Pharmaceutical Dosage Forms: Tablets (Lieberman et al., 1981); Ansel, Introduction to Pharmaceutical Dosage Forms 2nd Edition (1976); Remington's Pharmaceutical Sciences, 17th ed. (Mack Publishing Company, Easton, Pa., 1985); Advances in Pharmaceutical Sciences (David Ganderton, Trevor Jones, Eds., 1992); Advances in Pharmaceutical Sciences Vol 7. (David Ganderton, Trevor Jones, James McGinity, Eds., 1995); Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms (Drugs and the Pharmaceutical Sciences, Series 36 (James McGinity, Ed., 1989); Pharmaceutical Particulate Carriers: Therapeutic Applications: Drugs and the Pharmaceutical Sciences, Vol 61 (Alain Rolland, Ed., 1993); Drug Delivery to the Gastrointestinal Tract (Ellis Horwood Books in the Biological Sciences. Series in Pharmaceutical Technology; J. G. Hardy, S. S. Davis, Clive G. Wilson,

Eds.); Modern Pharmaceutics Drugs and the Pharmaceutical Sciences, Vol 40 (Gilbert S. Banker, Christopher T. Rhodes, Eds.). All of the aforementioned publications are incorporated by reference herein.

5 Tablets may contain suitable binders, lubricants, disintegrating agents, coloring agents, flavoring agents, flow-inducing agents, and melting agents. For instance, for oral administration in the dosage unit form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic, pharmaceutically acceptable, inert
10 carrier such as lactose, gelatin, agar, starch, sucrose, glucose, methyl cellulose, magnesium stearate, dicalcium phosphate, calcium sulfate, mannitol, sorbitol and the like. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth, or
15 sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes, and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride, and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite,
20 xanthan gum, and the like.

The compounds can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be
25 formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines. The compounds may be administered as components of tissue-targeted emulsions.

The compounds may also be coupled to soluble polymers as targetable
30 drug carriers or as a prodrug. Such polymers include polyvinylpyrrolidone, pyran copolymer, polyhydroxylpropylmethacrylamide-phenol, polyhydroxyethylaspartamidephenol, or polyethyleneoxide-polylysine substituted with palmitoyl residues. Furthermore, the compounds may be coupled to a
35 class of biodegradable polymers useful in achieving controlled

release of a drug, for example, polylactic acid, polyglycolic acid, copolymers of polylactic and polyglycolic acid, polyepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihydropyrans, polycyanoacylates, and crosslinked or amphipathic
5 block copolymers of hydrogels.

Gelatin capsules may contain the active ingredient compounds and powdered carriers, such as lactose, starch, cellulose derivatives, magnesium stearate, stearic acid, and the like. Similar diluents can
10 be used to make compressed tablets. Both tablets and capsules can be manufactured as immediate release products or as sustained release products to provide for continuous release of medication over a period of hours. Compressed tablets can be sugar coated or film coated to mask any unpleasant taste and protect the tablet from the
15 atmosphere, or enteric coated for selective disintegration in the gastrointestinal tract.

For oral administration in liquid dosage form, the oral drug components are combined with any oral, non-toxic, pharmaceutically
20 acceptable inert carrier such as ethanol, glycerol, water, and the like. Examples of suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions
25 reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent granules. Such liquid dosage forms may contain, for example, suitable solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents.

30 Liquid dosage forms for oral administration can contain coloring and flavoring to increase patient acceptance. In general, water, a suitable oil, saline, aqueous dextrose (glucose), and related sugar solutions and glycols such as propylene glycol or polyethylene
35 glycols are suitable carriers for parenteral solutions. Solutions for

parenteral administration preferably contain a water soluble salt of the active ingredient, suitable stabilizing agents, and if necessary, buffer substances. Antioxidizing agents such as sodium bisulfite, sodium sulfite, or ascorbic acid, either alone or combined, are suitable stabilizing agents. Also used are citric acid and its salts and sodium EDTA. In addition, parenteral solutions can contain preservatives, such as benzalkonium chloride, methyl- or propylparaben, and chlorobutanol. Suitable pharmaceutical carriers are described in Remington's Pharmaceutical Sciences, Mack Publishing Company, a standard reference text in this field.

The compounds of the instant invention may also be administered in intranasal form via use of suitable intranasal vehicles, or via transdermal routes, using those forms of transdermal skin patches well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery system, the dosage administration will generally be continuous rather than intermittent throughout the dosage regimen.

Parenteral and intravenous forms may also include minerals and other materials to make them compatible with the type of injection or delivery system chosen.

The inhibition of clusterin expression may be transient, and may occur in combination with a Hsp90 inhibitor. In humans with prostate cancer, this means that inhibition of expression should be effective starting within a day or two of Hsp90 inhibition or administration of an Hsp90 inhibitor, and extending for about 3 to 6 months thereafter. This may require multiple doses to accomplish. It will be appreciated, however, that the period of time may be more prolonged, starting Hsp90 inhibition and extending for substantial time afterwards without departing from the scope of the invention.

Aspects of the invention can be applied to the treatment of androgen-independent prostate cancer, or to prevent prostate cancer from becoming androgen-independent.

- 5 Aspects of the invention can be applied to the treatment of castration-resistant prostate cancer, or to prevent prostate cancer from becoming castration-resistant.

10 "Combination" means either at the same time and frequency, or more usually, at different times and frequencies as an oligonucleotide targeting clusterin expression, as part of a single treatment plan. Aspects of the invention include the administration of the oligonucleotide before, after, and/or during the administration of a Hsp90 inhibitor. A Hsp90 inhibitor may therefore be used, in
15 combination with an oligonucleotide according to the invention, but yet be administered at different times, different dosages, and at a different frequency, than the oligonucleotide.

As used herein, an "amount" or "dose" of an oligonucleotide measured
20 in milligrams refers to the milligrams of oligonucleotide present in a drug product, regardless of the form of the drug product.

As used herein, "effective" when referring to an amount of oligonucleotide which reduces clusterin expression, a Hsp90
25 inhibitor, or any combination thereof refers to the quantity of oligonucleotide, Hsp90 inhibitor, or any combination thereof that is sufficient to yield a desired therapeutic response without undue adverse side effects (such as toxicity, irritation, or allergic response) commensurate with a reasonable benefit/risk ratio when
30 used in the manner of this invention.

As used herein, "treating" encompasses, e.g., inhibition, regression, or stasis of the progression of prostate cancer. Treating also encompasses the prevention or amelioration of any
35 symptom or symptoms of prostate cancer.

As used herein, "inhibition" of disease progression or disease complication in a subject means preventing or reducing the disease progression and/or disease complication in the subject.

5

As used herein, a "symptom" associated with prostate cancer includes any clinical or laboratory manifestation associated with prostate cancer, and is not limited to what the subject can feel or observe.

10

As used herein, "pharmaceutically acceptable carrier" refers to a carrier or excipient that is suitable for use with humans and/or animals without undue adverse side effects (such as toxicity, irritation, and allergic response) commensurate with a reasonable benefit/risk ratio. It can be a pharmaceutically acceptable solvent, suspending agent or vehicle, for delivering the instant compounds and/or combinations to the subject.

The following abbreviations are used herein

20	PCa	prostate cancer
	CRPC	castrate resistant prostate cancer
	HSP	heat shock proteins
	CLU	clusterin
	PSA	prostate specific antigen
25	17-AAG	17-allylamino-17-demethoxygeldanamycin
	Aso	antisense oligonucleotide

Dosage Units

Administration of an oligonucleotide that targets clusterin expression can be carried out using the various mechanisms known in the art, including naked administration and administration in pharmaceutically acceptable lipid carriers. For example, lipid carriers for antisense delivery are disclosed in U.S. Patent Nos. 5,855,911 and 5,417,978, which are incorporated herein by reference.

35 In general, the oligonucleotide is administered by intravenous

(i.v.), intraperitoneal (i.p.), subcutaneous (s.c.), or oral routes, or direct local tumor injection. In preferred embodiments, an oligonucleotide targeting clusterin expression is administered by i.v. injection. In some embodiments, the amount of oligonucleotide
5 administered is 640mg.

The amount of antisense oligonucleotide administered is one effective to inhibit the expression of clusterin in prostate cells. It will be appreciated that this amount will vary both with the
10 effectiveness of the antisense oligonucleotide employed, and with the nature of any carrier used.

The amount of antisense oligonucleotide targeting clusterin expression administered may be from 40 to 640 mg, or 300-640 mg.
15 Administration of the antisense oligonucleotide may be once in a seven day period, 3 times a week, or more specifically on days 1, 3 and 5, or 3, 5 and 7 of a seven day period. In some embodiments administration of the antisense oligonucleotide is less frequent than once in a seven day period. Dosages may be calculated by
20 patient weight, and therefore a dose range of about 1-20 mg/kg, or about 2-10 mg/kg, or about 3-7 mg/kg, or about 3-4 mg/kg could be used. This dosage is repeated at intervals as needed. One clinical concept is dosing once per week with 3 loading doses during week one of treatment. The amount of antisense oligonucleotide administered
25 is one that has been demonstrated to be effective in human patients to inhibit the expression of clusterin in cancer cells.

In some embodiments of the invention, the amount of oligonucleotide targeting the expression of clusterin required for treatment of
30 prostate cancer is less in combination with a Hsp90 inhibitor, than would be required with oligonucleotide monotherapy.

Custirsen may be formulated at a concentration of 20 mg/mL as an isotonic, phosphate-buffered saline solution for IV administration

and can be supplied as an 8 mL solution containing 160 mg custirsén sodium in a single vial.

5 Custirsén may be added to 250 mL 0.9% sodium chloride (normal saline). The dose may be administered using either a peripheral or central indwelling catheter intravenously as an infusion over 2 hours. Additionally, an infusion pump may be used.

10 Administration of an Hsp90 inhibitor may be oral, nasal, pulmonary, parenteral, i.v., i.p., intra-articular, transdermal, intradermal, s.c., topical, intramuscular, rectal, intrathecal, intraocular, and buccal. One of skill in the art will recognize that higher doses may be required for oral administration than for i.v. injection.

15 The dose of Hsp90 inhibitor may be 60mg/kg, 55mg/kg, 45mg/kg, 40mg/kg, 35mg/kg, 25mg/kg, 20mg/kg, 15mg/kg, 10mg/kg, 5mg/kg or less.

20 A dosage unit of the oligonucleotide which reduces clusterin expression and an Hsp90 inhibitor may comprise one of each singly or mixtures thereof. A combination of an oligonucleotide which reduces clusterin expression and an Hsp90 inhibitor can be administered in oral dosage forms as tablets, capsules, pills, powders, granules, elixirs, tinctures, suspensions, syrups, and emulsions. An oligonucleotide which reduces clusterin expression and/or Hsp90
25 inhibitor may also be administered in intravenous (bolus or infusion), intraperitoneal, subcutaneous, or intramuscular form, or introduced directly, e.g. by injection or other methods, into or onto a prostate cancer lesion, all using dosage forms well known to those of ordinary skill in the pharmaceutical arts.

30 An oligonucleotide which reduces clusterin expression and/or Hsp90 inhibitor can be administered in admixture with suitable pharmaceutical diluents, extenders, excipients, or carriers (collectively referred to herein as a pharmaceutically acceptable
35 carrier) suitably selected with respect to the intended form of

administration and as consistent with conventional pharmaceutical practices. The unit will be in a form suitable for oral, rectal, topical, intravenous or direct injection or parenteral administration. The oligonucleotide and/or Hsp90 inhibitor can be administered alone or mixed with a pharmaceutically acceptable carrier. This carrier can be a solid or liquid, and the type of carrier is generally chosen based on the type of administration being used. Capsule or tablets can be easily formulated and can be made easy to swallow or chew; other solid forms include granules, and bulk powders. Tablets may contain suitable binders, lubricants, diluents, disintegrating agents, coloring agents, flavoring agents, flow-inducing agents, and melting agents. Examples of suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent granules. Such liquid dosage forms may contain, for example, suitable solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents. Oral dosage forms optionally contain flavorants and coloring agents. Parenteral and intravenous forms may also include minerals and other materials to make them compatible with the type of injection or delivery system chosen.

An oligonucleotide which reduces clusterin expression and/or Hsp90 inhibitor can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines. The compounds may be administered as components of tissue-targeted emulsions.

For oral administration in liquid dosage form, an Hsp90 inhibitor may be combined with any oral, non-toxic, pharmaceutically acceptable

inert carrier such as ethanol, glycerol, water, and the like. Examples of suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent granules. Such liquid dosage forms may contain, for example, suitable solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents.

In some embodiments of the invention, the amount of Hsp90 inhibitor required for treatment of prostate cancer is less in combination with an oligonucleotide targeting the expression of clusterin, than would be required with Hsp90 monotherapy.

A dosage unit may comprise a single compound or mixtures of compounds. A dosage unit can be prepared for oral or injection dosage forms.

According to an aspect of the invention, there is provided an oligonucleotide which reduces clusterin expression-containing pharmaceutical composition packaged in dosage unit form, wherein the amount of the oligonucleotide in each dosage unit is 640mg or less. Said pharmaceutical composition may include an Hsp90 inhibitor, and may be in an injectable solution or suspension, which may further contain sodium ions.

According to another aspect of the invention, there is provided the use of an oligonucleotide targeting clusterin expression and a Hsp90 inhibitor in the manufacture of a medicament for the treatment of cancer, where the medicament is formulated to deliver a dosage of 640mg or less of oligonucleotide to a patient. The medicament may contain sodium ions, and/or be in the form of an injectable solution.

General techniques and compositions for making dosage forms useful in the present invention are described in the following references: 7
Modern Pharmaceutics, Chapters 9 and 10 (Banker & Rhodes, Editors, 1979); Pharmaceutical Dosage Forms: Tablets (Lieberman et al., 1981);
5 Ansel, Introduction to Pharmaceutical Dosage Forms 2nd Edition (1976); Remington's Pharmaceutical Sciences, 17th ed. (Mack Publishing Company, Easton, Pa., 1985); Advances in Pharmaceutical Sciences (David Ganderton, Trevor Jones, Eds., 1992); Advances in
10 Pharmaceutical Sciences Vol 7. (David Ganderton, Trevor Jones, James McGinity, Eds., 1995); Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms (Drugs and the Pharmaceutical Sciences, Series 36 (James McGinity, Ed., 1989); Pharmaceutical Particulate Carriers: Therapeutic Applications: Drugs and the Pharmaceutical Sciences, Vol
15 61 (Alain Rolland, Ed., 1993); Drug Delivery to the Gastrointestinal Tract (Ellis Horwood Books in the Biological Sciences. Series in Pharmaceutical Technology; J. G. Hardy, S. S. Davis, Clive G. Wilson, Eds.); Modern Pharmaceutics Drugs and the Pharmaceutical Sciences, Vol. 40 (Gilbert S. Banker, Christopher T. Rhodes, Eds.). These references in their entirety are hereby incorporated by reference
20 into this application.

This invention will be better understood by reference to the Experimental Details which follow, but those skilled in the art will readily appreciate that the specific experiments detailed are only
25 illustrative of the invention as described more fully in the claims which follow thereafter.

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Experimental Details

Example 1. Hsp90 inhibitors induce expression of HSPs in prostate cancer (PCa) cells *in vitro* and *in vivo*.

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Dose- and time-dependent effects of Hsp90i-1 or Hsp90i-2 on the expression of CLU, Hsp90, Hsp70 and Akt protein and mRNA levels was evaluated in LNCaP and PC-3 cells. Both Hsp90i-1 and Hsp90i-2 increased Hsp70 and CLU protein levels up to 3 fold in a dose- and time-dependent manner (Fig. 1A, B and C). Hsp90 inhibition induced a dose- and a time dependent decline of Akt expression as previously reported (Lamoureux et al., 2011). mRNA levels of CLU, Hsp70 and Hsp90 also increased after Hsp90 inhibitor treatment (Fig. 1D).

15

Next the effects of Hsp90i-2 treatment on CLU expression were assessed *in vivo* in CRPC LNCaP xenografts using immunohistochemistry and western blot (Fig. 2). CLU expression increased 4-fold after treatment with Hsp90i-2-PRO (***, $p < 0.001$) compared with vehicle treated tumor (Fig. 2A, B). Similarly, Hsp70, which is considered a pharmacodynamic measure of Hsp90 inhibition (Solit et al., 2003; Eccles et al., 2008), increased 2.3-fold after treatment with Hsp90i-2-PRO (***, $p < 0.001$) (Fig. 2A).

20

Example 2. Treatment-induced feed forward loop involving CLU and HSF-1 activity.

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Since HSF-1 is the pre-dominant regulator of the heat shock response (Banerji et al., 2008; Workman et al., 2007), the effect of Hsp90 inhibition on HSF-1-activity and expression of HSPs was evaluated. Hsp90i-1 or Hsp90i-2 significantly induced CLU (Fig. 1) as well as HSF-1 activity in a dose-dependent manner (***, $p \leq 0.001$; Fig. 3A). CLU overexpression protected PC-3 tumor cells from Hsp90i-2-induced apoptosis (**, $p \leq 0.01$; Fig. 8A). Moreover, HSF-1 knockdown using siRNA decreases CLU expression, sensitizing tumor cells to apoptosis-induced by Hsp90i-2 (Fig. 8B), confirming that the protective effect of CLU is mediated by HSF-1. Surprisingly,

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overexpression of CLU also increased HSF-1 activity (***, $p \leq 0.001$, Fig. 3B), while CLU knockdown using siRNA or custirsen significantly decreased HSF-1 activity (*, $p \leq 0.05$; ***, $p \leq 0.001$; Fig. 3C), identifying novel feed-forward regulation of HSF-1 by CLU. Indeed, silencing of CLU inhibited HSF-1 transcriptional activity-induced by Hsp90i-1 or Hsp90i-2 (Fig. 3C), as well as HSF-1 regulated genes such as Hsp27 and Hsp70 (Fig. 3D). This effect can be explained by the ability of CLU knockdown to sequester HSF-1 in the cytoplasm (Fig. 8B).

10

Example 3. Increased potency of combination treatment comprising custirsen and Hsp90 inhibitor in increasing apoptosis in prostate tumor cell lines compared to monotherapy.

Since Hsp90 inhibitors induce up-regulation of CLU and functions as a mediator in treatment resistance (Zoubeidi et al. 2010; Gleave et al., 2005; Zellweger et al., 2003), it was next evaluated if CLU knockdown combined with Hsp90 inhibition increased treatment effectiveness. LNCaP cells were treated with custirsen and subsequently treated with indicated concentrations of Hsp90i-1 or Hsp90i-2. The combination had significantly enhanced Hsp90i-1 or Hsp90i-2 effectiveness, reducing cell viability by 20% at 100nM and 1000nM (*, $p < 0.05$) compared with cells treated with control ScrB ASO and Hsp90 inhibitors (Fig. 4A). To determine whether this effect was additive or a combination effect, the dose-dependent effects

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with constant ratio design and the combination index (CI) values was performed and calculated according to the Chou and Talalay median effect principal (Chou et al., 1984). Figure 4B shows the dose response curve (combination treatment, custirsen or HSP90i-2 monotherapy) and the combination index plots, indicating that custirsen and HSP90i-2 had enhanced combined potency on tumor cell growth compared to custirsen or Hsp90i-2 inhibitor monotherapy.

30

Moreover, OGX-011 potentiates the effect of Hsp90 inhibitor to induce apoptosis (Fig. 4C and D). Flow cytometric analysis shows

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that apoptotic rates (subG1 fraction) increased significantly ($p < 0.001$) when custirsen is combined with Hsp90i-1 (53%) or Hsp90i-2 (65.4%), compared to control ScrB ASO (4.2%), custirsen alone (17.4%), control ScrB ASO+Hsp90i-1 (18.3%) or control ScrB ASO+Hsp90i-2 (24.8%; Fig. 4C). Moreover, the combination custirsen with Hsp90i-1 or Hsp90i-2 induced more caspase-dependent apoptosis compared to Hsp90 inhibitor- or custirsen monotherapy, as shown by cleaved PARP and caspase-3 expression (Fig 4C). The significant increase of caspase-3 activity confirms that custirsen sensitizes cells to Hsp90 inhibition with increased apoptotic rates (Fig. 4D). Reduced cell viability from combined CLU plus Hsp90 inhibition results, in part, from decreases in p-Akt levels in both PC-3 and LNCaP cells, as well as AR (and PSA) expression in LNCaP cells (Fig. 4C).

15

Example 4. Potent combination therapy of custirsen and Hsp90i-1 in PC-3 xenografts *in vivo*.

The effects of combining custirsen with Hsp90i-1 was evaluated in PC-3 tumors *in vivo*. Male nude mice bearing PC-3 xenograft were randomly selected for treatment (custirsen + Hsp90i-1 vs control ScrB + Hsp90i-1; $n=7$). Custirsen + Hsp90i-1 had significantly enhanced the antitumor effects compared to of ScrB + Hsp90i-1 *in vivo*, reducing the mean tumor volume from 2935.3 mm³ to 1176.9 mm³ after 68 days (**; $p \leq 0.01$), compared to control ScrB (Fig. 5A). Cancer specific survival was significantly prolonged with combined custirsen + Hsp90i-1 compared with controls (71.4% vs 14.3% at day 72, respectively; *, $p \leq 0.05$; Fig. 5B. Immunohistochemical analysis reveals decreased CLU, Ki67, and Akt, expression after treatment with custirsen + Hsp90i-1 compared to other groups (Fig 5C). Additionally, custirsen + Hsp90i-1 treated tumors had higher apoptosis as shown by increased TUNEL staining compared with other groups (Fig 5C).

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Example 5. Potent combination therapy of custirsen and Hsp90i-2-PRO in LNCaP CRPC xenografts *in vivo*.

Next the effects of combined treatment with custirsen and Hsp90i-2-PRO was assessed in castrate resistant LNCaP tumors. Mice bearing LNCaP tumors were castrated when PSA values exceeded 50ng/ml. Once PSA levels relapsed above pre-castration levels mice were randomly assigned to vehicle control, Hsp90i-2-PRO alone, Hsp90i-2-PRO + control ScrB, or Hsp90i-2-PRO + custirsen (n=10 in each group). Mice treated with Hsp90i-2-PRO + custirsen had significant delays in tumor growth compared with all other groups (Fig 6A) (at 10 days, respectively 265.3 mm³, and 892.7 mm³ for control, 646.4mm³ for Hsp90i-2-PRO alone and 551.56 mm³ for Hsp90i-2-PRO+control ScrB). By 7 weeks post treatment, all mice in the control had been euthanized; tumor volume in the Hsp90i-2-PRO + custirsen group was 1517.4 mm³ compared to 2483.6mm³ for HSP90i-2-PRO alone and 2176.4 mm³ for Hsp90i-2-PRO+control ScrB; ***, p<0.001; Fig. 6A).

Serum PSA levels were also significantly lower (~4-fold) in the mice receiving custirsen + Hsp90i-2-PRO compared with other groups (***, p<0.001; Fig 6B). The combination custirsen + Hsp90i-2-PRO group had a mean PSA level of 120ng/ml after 42 days compared to 418.7ng/ml in vehicle group, 527ng/ml in Hsp90i-2-PRO alone, or 480.3ng/ml in scrB + Hsp90i-2-PRO groups. The combination custirsen+ Hsp90i-2-PRO group had a significantly increased PSA doubling time (33.6 weeks; *, p<0.05) and decreased PSA velocity (13.78ng/mL/week; *, p<0.05) compared with other groups (PSA doubling time: ~2.4weeks; velocity: ~85ng/mL/week; Fig. 6C).

Overall survival was also significantly longer in mice treated with combined custirsen + Hsp90i-2-PRO (Fig 6D). By day 57, all mice died or were euthanized due to high tumor burden in control, Hsp90i-2-PRO alone, or control ScrB + Hsp90i-2-PRO groups compared with the combined custirsen + Hsp90i-2-PRO group, where all mice were still alive (p<0.001) after 62 days. These data demonstrate that targeting CLU using custirsen in combination with HSP90i-2-PRO

inhibits tumor growth and prolongs survival in human CRPC xenograft model significantly more than monotherapy.

Consistent with in vitro findings, immunohistochemical analysis revealed decreased CLU, Ki67, Akt, and AR expression after treatment with combined custirsen + Hsp90i-2-PRO compared with other groups (Fig 7A). The immunostaining results were corroborated by western blotting (Fig 7B). Additionally, tumors treated with combination custirsen + Hsp90i-2-PRO had higher apoptosis rates compared with other groups as shown by increased TUNEL staining (Fig 7A). These data suggest that decreases in tumor progression custirsen + Hsp90i-2-PRO treated tumors result from both reduced proliferation rates as well as increased apoptosis rates.

Example 6. Materials and Methods for Examples 1-5.

Tumor cell lines and reagents:

The human PCa cell line PC-3 was purchased from the American Type Culture Collection (2008, ATCC-authentication by isoenzymes analysis) and maintained in DMEM (Invitrogen-Life Technologies, Inc.) supplemented with 5% fetal bovine serum and 2mmol/L Lglutamine. LNCaP cells were kindly provided by Dr. Leland W.K. Chung (1992, MDACC, Houston Tx) and tested and authenticated by whole-genome and whole-transcriptome sequencing on Illumina Genome Analyzer IIx platform in July 2009. LNCaP cells were maintained RPMI 1640 (Invitrogen Life Technologies, Inc.) supplemented with 5% fetal bovine serum and 2mmol/L L-glutamine. All cell lines were cultured in a humidified 5% CO₂/air atmosphere at 37°C. All cell lines were passaged for less than 3 months after resurrection. Western blotting and/or real time PCR was performed for AR and PSA each time when LNCaP cells were resurrected.

Therapeutic agents:

Hsp90 inhibitor, HSP90i-2 (4-(6,6-Dimethyl-4-oxo-3-trifluoromethyl-4,5,6,7-tetrahydro-indazol-1-yl)-2-(4-hydroxy-cyclohexylamino)-benzamide) and its prodrug HSP90i-2-PRO were used respectively for

- in vitro and *in vivo* studies. These compounds are novel synthetic small molecular weight inhibitors that bind the N-terminal adenosine triphosphate binding site of Hsp90 and HSP90i-2-PRO is orally bioavailable. For the *in vitro* studies, HSP90i-2 was dissolved in dimethyl sulfoxide (DMSO) at 10mM stock solutions and stored at -20°C. For the *in vivo* studies, HSP90i-2-PRO was dissolved in PBS 1% carboxymethylcellulose and 0.5% Tween 80 (Invitrogen-Life Technologies, Inc.) at 15mg/ml and stored at 4°C.
- HSP90i-1 (17-allylamino-17-demethoxygeldanamycin (17-AAG)) was used for *in vitro* and *in vivo* studies. For the studies, 17-AAG was dissolved in dimethyl sulfoxide (DMSO) at 10mM stock solutions and stored at -20°C.
- Clusterin siRNA and Antisense Oligonucleotides*
- siRNAs were purchased from Dharmacon Research, Inc. (Lafayette, CO) using the siRNA sequence corresponding to the human CLU initiation site in exon 2 and a scramble control as previously described (Sowery et al., 2008). Second-generation antisense (custirsen) and scrambled (ScrB) oligonucleotides with a 2'-O-(2-methoxy)ethyl modification were supplied by OncoGenex Pharmaceuticals (Vancouver, British Columbia, Canada). Custirsen sequence (5'-CAGCAGCAGAGTCTTCATCAT-3'), SEQ ID NO:3 corresponds to the initiation site in exon II of human CLU. The ScrB control sequence was 5'-CAGCGCTGACAACAGTTTCAT-3' (SEQ ID NO: 44). Prostate cells were treated with siRNA or oligonucleotides using protocols described previously (Sowery et al., 2008).

Cell proliferation and apoptosis assays:

- Prostate cells lines were plated in appropriate media (DMEM or RPMI) with 5% FBS and treated with Hsp90i-2-PRO or Hsp90i-1 at indicated concentration and time and cell growth was measured using the crystal violet assay as described previously (Leung et al., 2000). Detection and quantitation of apoptotic cells were done by

flow-cytometry (described below) and western blotting analysis. Each assay was repeated in triplicate.

The combination index (CI) was evaluated using CalcuSyn dose effect analysis software (Biosoft, Cambridge, UK). This method, based on the multiple drug effect equation of Chou-Talalay (Chou et al., 1984), is suitable for calculating combined drug activity over a wide range of growth inhibition: CI =1, additivity; CI >1, antagonism; CI <1, combination effect. CI was calculated at ED₅₀ and ED₇₅.

Caspase-3 activity was assessed 3 days after treatment using the kit CaspACE Assay System, Fluorometric (Promega, Madison, WI, USA). Fifty µg of total cell lysate were incubated with caspase-3 substrate AC-DEVD-AMC at room temperature for 4h and caspase-3 activity was quantified in a fluorometer with excitation at 360nm and emission 460nm.

Cell cycle analysis:

Prostate cancer cell lines were incubated in the absence or the presence of 1µM Hsp90i-2 or Hsp90i-1 for 72h, trypsinized, washed twice and incubated in PBS containing 0.12% Triton X-100, 0.12mM EDTA and 100µg/ml ribonuclease A; 50µg/ml propidium iodide was then added to each sample for 20min at 4°C. Cell cycle distribution was analyzed by flow cytometry (Beckman Coulter Epics Elite, Beckman, Inc., Miami, FL), based on 2N and 4N DNA content. Each assay was done in triplicate.

Western blotting analysis:

Samples containing equal amounts of protein (depending on the antibody, 5-50µg) from lysates of cultured tumor prostate cell lines underwent electrophoresis on SDS-polyacrylamide gel and were transferred to nitrocellulose filters. The filters were blocked in Odyssey Blocking Buffer (LI-COR Biosciences) at room temperature for 1h and blots were probed overnight at 4°C with primary

antibodies to detect proteins of interests. After incubation, the filters were washed 3 times with washing buffer (PBS containing 0.1% Tween) for 5min. Filters were then incubated for 1h with 1:5,000 diluted Alexa Fluor secondary antibodies (Invitrogen) at room temperature. Specific proteins were detected using ODYSSEY IR imaging system (LI-COR Biosciences) after washing.

Quantitative Reverse Transcription-PCR:

Total RNA was extracted from cultured cells after 48h of treatment using TRIzol reagent (Invitrogen Life Technologies, Inc.). Two µg of total RNA was reversed transcribed using the Transcriptor First Strand cDNA Synthesis Kit (Roche Applied Science). Real-time monitoring of PCR amplification of complementary DNA (cDNA) was performed using DNA primers (supplemental table S1) on ABI PRISM 7900 HT Sequence Detection System (applied Biosystems) with SYBR PCR Master Mix (Applied Biosystems). Target gene expression was normalized to GAPDH levels in respective samples as an internal standard, and the comparative cycle threshold (Ct) method was used to calculated relative quantification of target mRNAs. Each assay was performed in triplicate.

Luciferase assay:

LNCaP and C4-2 cells (2.5×10^5) were plated on six-plates and transfected using lipofectin (6µL per well; Invitrogen Life Technologies, Inc.). The total amount HSE plasmids DNA used were normalized to 1µg per well by the addition of a control plasmid. One µM HSP90i-2 or Hsp90i-1 was added 4h after the transfection and for 48h. HSE-luciferase activity was measured using Dual-Luciferase Reporter Assay System (Promega) with the aid of a microplate luminometer (EG&G Berthold). All experiments were carried out in triplicate wells and repeated 3 times using different preparations of plasmids.

Immunofluorescence:

Tumor cells were grown on coverslips and treated with different concentration of Hsp90i-2 or Hsp90i-1 for 48h. After treatment, cells were fixed in ice-cold methanol completed with 3% acetone for 10min at -20°C. Cells were the washed thrice with PBS and incubated with 0.2% Triton/PBS for 10min, followed by washing and 30min blocking in 3% nonfat milk before the addition of antibody overnight to detect HSF-1 (1:250). Antigens were visualized using anti-mouse antibody coupled with FITC (1:500; 30 min). Photomicrographs were taken at 20X magnification using Zeiss Axioplan II fluorescence microscope, followed by analysis with imaging software (Northern Eclipse, Empix Imaging, Inc.).

Animal Treatment:

Male athymic nude mice (Harlan Sprague-Dawley, Inc.) were injected s.c. with 2×10^6 LNCaP cells (suspended in 0.1mL Matrigel; BD Biosciences). The mice were castrated once tumors reach between 300 and 500mm³ or the PSA level increased above 50ng/mL. Once tumors progressed to castrate resistance, mice were randomly assigned to vehicle, Hsp90i-2-PRO alone, Hsp90i-2-PRO + ScrB ASO or Hsp90i-2-PRO + custirsen. Hsp90i-2-PRO (Prodrug, 25mg/kg; formulation in 0.5% CMC+0.5% Tween-80) is orally administered three times per week and custirsen or ScrB ASO (15mg/kg) was injected intra-peritoneally once daily for the first week and then three times per week. Each experimental group consisted of 10 mice. Tumor volume was measured twice weekly (length x width x depth x 0.5432). Serum PSA was determined weekly by enzymatic immunoassay (Abbott IMX, Montreal, Quebec, Canada). PSA doubling time (PSAdt) and velocity were calculated by the log-slope method ($PSAt = PSA_{initial} \times e^{mt}$). Data points were expressed as average tumor volume $\pm$ SEM or average PSA concentration $\pm$ SEM.

To establish PC-3 tumors, 2×10^6 PC-3 cells were inoculated s.c. in the flank region of 6-8 week-old male athymic mice (Harlan Sprague-Dawley, Inc.). When tumors reached 100mm³, usually 3-4 weeks after

injection, mice were randomly selected for treatment with Hsp90i-1 (25mg/kg) + control ScrB ASO (15mg/kg) or Hsp90i-1 + custirsen (15mg/kg). Hsp90i-1 was injected i.p. three times per week, and custirsen or ScrB were injected i.p. once/day for the first week and then three times per week. For each experimental group consisted of 7 mice. Tumor volume was measured twice weekly. Data points were expressed as average tumor volume $\pm$ SEM.

When tumor volume reached $\geq 10\%$ of body weight, mice were sacrificed and tumors harvested for evaluation of protein expression by western blotting analyses and immunohistochemistry. All animal procedures were performed according to the guidelines of the Canadian Council on Animal Care and appropriate institutional certification.

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Immunohistochemistry:

Immunohistochemical stains were performed on formalin-fixed and paraffin-embedded 4 μ m sections of tumor samples using adequate primary antibody, and the Ventana autostainer Discover XT (Ventana Medical System) with enzyme labeled biotin streptavidin system and solvent resistant 3,3'-diaminobenzidine Map kit. All comparisons of staining intensities were made at 200x magnifications.

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Statistical analysis:

All *in vitro* data were assessed using the Student t test and Mann-Whitney test. Tumor volumes of mice were compared using Kruskal-Wallis test. Overall survival was analyzed using Kaplan-Meier curves and statistical significance between the groups was assessed with the log-rank test (Graphpad Prism). Levels of statistical significance were set at $P < 0.05$.

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Antibodies used for western blotting:

PARP (1/1000) Caspase 3 (1/1000), Akt (1/1000), p-Akt (1/500), are from cell signaling. Cyclin D1 (1/1000), HSP90 (1/1000), HSP70 (1/1000), clusterin (1/1000), AR (1/1000), PSA (1/1000)

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HSF-1 (1/1000) are from Santa Cruz. HSP27 (1/5000) is from Assays Designs.

Table 3: Primers used for quantitative real-time PCT

Sequence name	Sequence 5' to 3' forward	Sequence 5' to 3' reverse
Clusterin	GAGCAGCTGAACGAGCAGTTT (SEQ ID NO: 45)	CTTCGCCTTGCGTGAGGT (SEQ ID NO: 46)
Hsp70	TGCCCTATCCAGATCCTGCTA (SEQ ID NO: 47)	GAGCCATCAGACTGAGGAGTGA (SEQ ID NO: 48)
Hsp90	TTCAGGCCCTTCCCGAAT (SEQ ID NO: 49)	TCACTCCTTCCTTGGCAACAT (SEQ ID NO: 50)

Discussion

Prostate cancer responds initially to anti-androgen therapies, however, progression of castration resistant disease frequently occurs. Small molecule inhibitors of Hsp90 show promise in the treatment of castration-resistant prostate cancer (CRPC) and other cancers, however these inhibitors trigger a heat shock response that attenuates drug effectiveness. In prostate cancer, treatment resistance emerges early due to compensatory mechanisms involving activation of heat shock factor 1 (HSF-1). Once released from Hsp90, HSF-1 translocates to the nucleus, binds to heat shock elements (HSE) of Hsp genes and increases Hsp transcription activity (Whitesell et al., 2005). Therefore, Hsp90 inhibition induces a heat shock response with increased expression of several Hsps including Hsp90, Hsp70, Hsp27 and clusterin (CLU), which enhance tumor cell survival and treatment resistance. The up-regulation of these molecular chaperones has been reported to play a role in cellular recovery from stress by restoring protein homeostasis, promoting thermotolerance, cell survival, and treatment resistance (Takayama et al., 2003; Zoubeidi et al., 2010). The data herein show that preventing CLU induction in this response would enhance Hsp90 inhibitor-induced CRPC cell death *in vitro* and *in vivo*. As disclosed herein, CRPC was treated with Hsp90 inhibitor HSP90i-2-PRO or HSP90i-1 in the absence or presence of custirsen, an antisense drug that targets CLU. Treatment with either Hsp90 inhibitor alone increased nuclear translocation and transcriptional activity of the heat shock factor HSF-1, which stimulated dose- and time-dependent increases in heat shock protein expression, including especially CLU expression. Treatment-induced increases in CLU were blocked by custirsen, such that the combination of custirsen and either Hsp90 inhibitor had enhanced inhibition activity on CRPC cell growth and apoptosis compared to custirsen or Hsp90 inhibitor monotherapy. Accompanying these effects was a decrease in HSF-1 transcriptional activity as well as expression of HSPs, Akt, PSA and androgen receptor. *In vivo* evaluation of the Hsp90 inhibitors with custirsen in xenograft models of human CRPC demonstrated that custirsen markedly potentiated anti-tumor efficacy,

leading to an 80% inhibition of tumor growth with prolonged survival compared to Hsp90 inhibitor monotherapy. Together, the findings herein indicate that Hsp90 inhibitor-induced activation of the heat shock response and CLU is attenuated by custirsen, with combination
5 therapy having increased potency on delaying CRPC progression.

Development of treatment resistance is a common feature of most malignancies and the underlying basis for most cancer deaths. Treatment resistance evolves, in part, from selective pressures of
10 treatment that collectively increase the apoptotic rheostat of cancer cells. Survival proteins up-regulated after treatment stress include anti-apoptotic members of the bcl-2 protein family, survivin, and molecular chaperones like CLU and other HSPs (Zellweger et al. 2003).

15 Molecular chaperones help cells cope with stress-induced protein aggregation, and play prominent roles in cell signaling and transcriptional regulatory networks. Chaperones act as genetic buffers stabilizing the phenotype of various cells and organisms at
20 times of environmental stress, and enhance Darwinian fitness of cells during cancer progression and treatment resistance (Whitesell et al., 2005). Heat shock chaperones are key components of the heat shock response, a highly conserved stress-activated protective mechanism also associated with oncogenic transformation and thermo-
25 tolerance (Dai et al., 2007). Chaperones are particularly important in regulating misfolded protein and endoplasmic reticular (ER) stress responses, an emerging area of interest in treatment stress and resistance. A growing enthusiasm for therapeutic modulation of this proteostasis network highlights Hsp's and CLU as rational
30 targets because of their multifunctional roles in signaling and transcriptional networks associated with cancer progression and treatment resistance. Cancer cells express higher levels of molecular chaperones and pirate the protective functions of HSF1 to support their transformation (Dai et al., 2007). Indeed, inhibitors
35 of Hsp90, Hsp70, Hsp27 or CLU have all been reported to induce

cancer cell death and sensitize chemotherapy (Lamoureux et al., 2011; Guo et al., 2005).

Increased expression of clusterin (CLU) has been associated with
5 chemoresistance, radioresistance, and hormone resistance (Zellweger
et al., 2003; July et al., 2004). CLU is a stress-induced
cytoprotective chaperone that inhibits protein aggregation in a
manner analogous to small HSPs, and its promoter contains a 14-bp
element recognized by the transcription factor HSF-1 (Humphreys et
10 al., 1999). In human PCa, CLU levels are low in Gleason grade 3
untreated hormone-naive tissues, but increase with higher Gleason
score (Steinberg et al., 1997) and within weeks after androgen
deprivation (July et al., 2002). CLU expression correlates with
loss of the tumor suppressor gene Nkx3.1 during the initial stages
15 of prostate tumorigenesis in Nkx3.1 knockout mice (Song et al.,
2009). Experimental and clinical studies associate CLU with
development of treatment resistance, where CLU suppresses
treatment-induced cell death in response to androgen withdrawal,
chemotherapy or radiation (Miyake et al., 2000a; July et al., 2002;
20 Miyake et al., 2000b; Miyake et al., 2000c). Over-expression of CLU
in human prostate LNCaP cells accelerates progression after
hormone- or chemo-therapy (Miyake et al., 2000a; Miyake et al.,
2000c), identifying CLU as an anti-apoptotic gene up-regulated by
treatment stress that confers therapeutic resistance. Custirsen is
25 a second-generation phosphorothioate antisense oligonucleotide
currently in late stage clinical development that potently inhibits
CLU expression and enhances the efficacy of anticancer therapies in
various human cancers including PCa (Zoubeidi et al., 2010, Gleave
et al., 2005). While targeting CLU enhances the cytotoxic effects
30 of chemotherapy and delays tumor growth in various human cancers
including PCa (Miyake et al., 2005), a role for CLU has not been
characterized in the context of Hsp90 inhibitor treatment and
resistance. As shown herein, Hsp90 inhibition induces a heat shock
response with increased HSF-1 activity and CLU expression, which
35 functions to inhibit treatment-induced apoptosis and enhance

emergence of treatment resistance. Knockdown of CLU using custirsen potentiates the effect of Hsp90 inhibitors in CRPC.

Aspects of the present invention relate to the unexpected discovery
5 that an oligonucleotide targeting clusterin expression such as
custirsen, together with a Hsp90 inhibitor is a potent combination
for treatment of prostate cancer. The discovery that an anti-
clusterin therapy combined with Hsp90 is so potent is particularly
surprising because Hsp90 is known to increase the expression of
10 multiple cytoprotective proteins.

Several Hsp90 inhibitors including HSP90i-2 have potent anti-tumor
activity in various preclinical models (Lamoureux et al., 2001;
Chandarlapaty et al., 2008; Okawa et al., 2009) and are in clinical
15 trials (Lamoureux et al., 2011; Sydor et al., 2006). Consistent
with prior reports (Lamoureux et al., 2011; Cervantes-Gomez et al.,
2009), the data herein show that Hsp90 inhibitors induce a stress
response with activation of the transcription factor HSF-1 and
subsequent increased levels of Hsp90 itself, Hsp70 and CLU. This
20 heat shock response likely enhance emergence of treatment
resistance, as inhibition of transcription using Actinomycin D
attenuates HSP90i-1-mediated Hsp70 and Hsp27 expression and
potentiates the effect of HSP90i-1 *in vitro* (Cervantes-Gomez et
al., 2009). Additionally, inhibition of the stress response by
25 silencing HSF-1 also increases the activity of Hsp90 inhibitors
(Bagatell et al., 2000). The experiments disclosed herein evaluated
the role of CLU in this heat shock response since CLU is
dramatically induced by Hsp90 inhibitor treatment and CLU
inhibitors are in late stage clinical development.

30 CLU is associated with many varied patho-physiological processes
including reproduction, lipid transport, complement regulation and
apoptosis (Zoubeidi et al. 2010; Rosenberg et al., 1995). CLU
expression is rapidly upregulated in various tissues undergoing
35 apoptosis, including normal and malignant prostate and breast

tissues following hormone withdrawal (Kyprianou et al., 1990; Kyprianou et al., 1991). Previous studies have also linked CLU expression with induction and progression of many cancers, including CRPC (Zoubeidi et al., 2010). Furthermore, CLU up-regulation following androgen ablation in xenograft tumor models accelerates progression to castrate resistance and renders cells resistant to other apoptotic stimuli, including taxane chemotherapy (Miyake et al., 2000; Miyake et al., 2001). Consistent with these accumulated findings (Miyake et al., 2001), inhibition of CLU using custirsen synergistically enhances conventional as well as molecular targeted therapies in PCa preclinical models (Sowery et al., 2008). Indeed, custirsen is now in Phase III trials as Phase II studies reported >90% inhibition of CLU in human prostate cancer tissues (Chi et al., 2005), and 7 months prolonged survival when OGX-011 is combined with docetaxel in CRPC (Chi et al., 2008; Chi et al., 2010).

The data herein show that Hsp90 inhibitors increase CLU levels both *in vitro* and *in vivo*, while clusterin inhibits HSP90i-2 or HSP90i-1 induced CLU. As expected (Cervantes-Gomez et al., 2009; Bagatell et al., 2000), HSP90i-2 or HSP90i-1 induces HSF-1 transcriptional activity leading to up-regulation of HSPs expression. Surprisingly, the experiments described herein found that CLU silencing abrogates, while CLU overexpression enhances, Hsp90 inhibitor-induced HSF-1 transcription activity, identifying a role for CLU in the regulation of HSF-1 and the heat shock response itself. CLU knockdown blocks the translocation to HSF-1 to the nucleus following treatment with Hsp90 inhibitors. This effect of CLU on HSF-1 activity is biologically relevant since CLU overexpression protects, while CLU silencing enhances, cytotoxicity of Hsp90 inhibitors. Consistent with these *in vitro* results, synergistic effects were also observed *in vivo* in PC-3 and LNCaP models when custirsen was combined with Hsp90 inhibitors. Combination custirsen plus Hsp90 inhibitor significantly delay CRPC tumor growth and prolonged survival in PC-3 and LNCaP models. Increased apoptotic

rates with combined Hsp90 and CLU inhibition suggests that delayed tumor progression resulted from enhanced treatment-induced apoptosis. Systemic administration of an oligonucleotide which reduces clusterin expression plus a Hsp90 inhibitor decreases tumor growth compared with control ScrB ASO plus an Hsp90 inhibitor in PC-3 model and LNCaP castration-resistant prostate cancer, respectively. This inhibition of tumor progression is accompanied with a prolongation of survival in both prostate cancer models. Detection of increased apoptosis after combined clusterin plus Hsp90 inhibition by detection of TUNEL using immunohistochemistry suggests that delayed tumor progression after combined therapy results from enhanced Hsp90 inhibitor-induced apoptosis. Collectively, these results highlight, for the first time, a biologically relevant feed-forward regulation loop of CLU on HSF-1 and the heat shock response.

The effect of an oligonucleotide which reduces clusterin expression in combination with an Hsp90 inhibitor on PSA level was examined in the LNCaP castration-resistant prostate cancer model as disclosed herein above. As shown herein, targeting CLU using siRNA or the antisense drug, custirsen, suppressed treatment-induced CLU induction and enhanced Hsp90 inhibitor-induced cell death in prostate cancer cells. Serum PSA level is an established and useful biomarker regulated by androgen receptor (AR) in the presence of androgens (Magklara et al., 2002), and a valuable tool in the follow-up of patients to assess the efficacy of chemotherapy. In addition to the effects of CLU inhibition on the heat shock response, observations in the castrate-sensitive, AR-positive LNCaP model highlight another possible benefit of combined CLU and Hsp90 suppression involving AR activity. Hsp90 inhibition is known to destabilize and degrade the AR with decreased PSA expression (Solit et al., 2002; Georget et al., 2002). *In vivo*, serum PSA levels as well as PSA doubling time and velocity, were significantly reduced with combination OGX-011 therapy compared with PF-04929113 monotherapy. Serum PSA level is an established and useful AR-

regulated biomarker (Kim et al., 2004) and a valuable tool in assessing efficacy of chemotherapy. Interestingly, at the low doses of Hsp90 inhibitor used in this in vivo study, no effect on serum PSA level was apparent. Lower PSA levels with combination therapy correlated with lower AR levels. This correlation between CLU inhibition and lower AR levels may involve the regulation loop of CLU on HSF-1 and the role of HSF-1 in regulating expression of other AR chaperones (eg. Hsp27, Hsp70, Hsp90, FKBP5.2) and we are actively exploring the molecular basis in ongoing experiments. While CLU is known to be transcriptionally activated by HSF-1 (Zoubeidi et al., 2010), the data herein also show that CLU exerts a feed forward loop that in turn activates HSF-1. CLU knockdown decreases HSF-1 transcriptional activity and abrogates its nuclear translocation, which subsequently leads to decreased Hsp27, Hsp70 and Hsp90 expression, similar to that observed after HSF-1 knockdown (Rossi et al., 2006). Consequently, AR stability is reduced because of lowered chaperone levels.

In addition to increased potency of anti-tumor activity, combination therapy may also allow dose reduction strategies to reduce toxicity. For example, HSP90i-1 induced hepatotoxicity as monotherapy at 60mg/kg/day (Glaze et al., 2005), while HSP90i-2-PRO caused body weight loss at 50mg/kg/day. In a previous study, 50mg/kg HSP90i-2-PRO as monotherapy inhibited LNCaP CRPC tumor progression (Lamoureux et al., 2011). At sub-therapeutic doses of 25 mg/kg/day used in the present study, HSP90i-2-PRO monotherapy showed marginal, non-significant decreases in tumor volume and no effect on serum PSA levels; however, significant delays in tumor progression were seen at this lower dose when HSP90i-2-PRO was combined with custirsen, with no toxicity observed.

The data disclosed herein help define how stress induced by Hsp90 inhibitors regulates CLU by induction of HSF-1 activity and, in turn, how CLU regulates HSF-1 activity, cell survival, and treatment resistance. As demonstrated herein, for the first time,

that CLU inhibition abrogates the heat shock response induced Hsp90 inhibitors. These observations are clinically relevant since CLU inhibitors are in phase III clinical trials, and provide a framework for building new drug combinations based on mechanism-based interventions to overcome drug resistance. The present invention relates to the development of targeted strategies employing custirsen in combination with Hsp90 inhibitors to improve patient outcome in CRPC.

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What is claimed is:

1. Use of
 - i) an antisense or RNAi oligonucleotide that is complementary to the sequence of clusterin of a mammalian subject, and that reduces clusterin expression; and
 - ii) a Heat Shock Protein 90 (Hsp90) inhibitor comprising 4-(6,6-Dimethyl-4-oxo-3-trifluoromethyl-4,5,6,7-tetrahydro-indazol-1-yl)-2-(4-hydroxy-cyclohexylamino)-benzamide, or a pharmaceutically acceptable salt thereof, or a prodrug that is metabolized to release 4-(6,6-Dimethyl-4-oxo-3-trifluoromethyl-4,5,6,7-tetrahydro-indazol-1-yl)-2-(4-hydroxy-cyclohexylamino)-benzamidein the manufacture of a medicament for treating prostate cancer in the mammalian subject.
2. The use of claim 1, wherein the cancer is androgen-independent prostate cancer.
3. The use of claim 1 or 2, wherein the mammalian subject is human.
4. The use of any one of claims 1-4, wherein the oligonucleotide is an antisense oligonucleotide.
5. The use of claim 5, wherein the antisense oligonucleotide comprises one of Seq ID NOs: 1-11.
6. The use of claim 5, wherein the antisense oligonucleotide comprises SEQ ID NO: 3.
7. The use of claim 6, wherein the antisense oligonucleotide is modified to enhance *in vivo* stability relative to an unmodified oligonucleotide of the same sequence.
8. The use of claim 7, wherein the oligonucleotide is custirsen.

9. The use of any one of claims 1-4, wherein the oligonucleotide is an RNAi oligonucleotide.
10. The use of claim 9, wherein the RNAi oligonucleotide comprises one of SEQ ID NOs: 19-42.
11. The use of any one of claims 1-10, wherein the combination comprises Hsp90i-2-PRO.
12. The use of any one of claims 1-11, wherein the combination of the oligonucleotide and the Hsp90 inhibitor is effective to inhibit the proliferation of prostate cancer cells.