Molecular Pathways

Starving the Addiction: New Opportunities for Durable Suppression of AR Signaling in Prostate Cancer

Karen E. Knudsen1,2,3 and Howard I. Scher4,5,6,7

Abstract
Clinical data and models of human disease indicate that androgen receptor (AR) activity is essential for prostate cancer development, growth, and progression. The dependence of prostatic adenocarcinoma on AR signaling at all stages of disease has thereby been exploited in the treatment of disseminated tumors, for which ablation of AR function is the goal of first-line therapy. Although these strategies are initially effective, recurrent tumors arise with restored AR activity, and no durable treatment has yet been identified to combat this stage of disease. New insights into AR regulation and the mechanisms underlying resurgent AR activity have provided fertile ground for the development of novel strategies to more effectively inhibit receptor activity and prolong the transition to therapeutic failure.

Background
Prostate cancer remains the second leading cause of cancer death in the United States and the most frequently diagnosed noncutaneous malignancy. Estimates are that over 192,000 cases will be diagnosed and greater than 27,000 will succumb to the disease in 2009 (1). The goal of treatment for clinically localized disease is cure, typically by surgery or radiation therapy (2). For patients who recur systemically after definitive treatment, or who present with locoregional or metastatic disease, long-term disease control is the primary objective. Typically, this entails a series of hormonal therapies that suppress androgen receptor (AR) signaling, as prostate cancers are exquisitely dependent on AR function for survival and progression. Although AR-directed therapies inhibit tumor growth, disease is rarely eliminated, and resistance to therapy is acquired through restored AR function. Once progression is documented, the course is inevitably fatal. Docetaxel-based chemotherapy is essential for prostate cancer development, growth, and progression. The dependence on AR function for survival and progression. Although AR-directed therapies inhibit tumor growth, disease is rarely eliminated, and resistance to therapy is acquired through restored AR function. Once progression is documented, the course is inevitably fatal. Docetaxel-based chemotherapy can prolong life but is likewise not curative, highlighting the need for more effective treatments (3, 4).

Androgen exerts its biological effects through AR, a ligand-dependent transcription factor and a member of the nuclear receptor superfamily (5). In prostatic adenocarcinoma cells, the most abundant serum androgen, testosterone, is converted into a higher-affinity ligand for AR, dihydrotestosterone (DHT), via the action of 5-alpha reductase (6). Upon ligand binding, AR sheds inhibitory chaperones such as heat-shock proteins, undergoes rapid homodimerization and nuclear translocation, and binds to DNA at specific sequences termed androgen-responsive elements (AREs; ref. 7). Once bound, the receptor recruits cooperative transcriptional cofactors (coactivators) that assist in inducing gene expression (8, 9). This AR-dependent gene expression program results in varied biological outcomes dependent on cell context, including the induction of genes encoding secretory products of the prostate [e.g., prostate-specific antigen, (PSA)], cell survival proteins, and genes that promote cell cycle initiation (10). The striking requirement of prostate cancer cells for AR activity is illustrated in the clinic, wherein therapeutic suppression of AR signaling, as typically achieved through ligand depletion and/or the use of direct AR antagonists, results in decreased PSA production, objective tumor regressions, and palliation of symptoms when present (11). The durability of the effect can range from months to years but, unfortunately, are not permanent, and after a variable period of time tumor regrowth occurs. This is heralded first by rising PSA values (“biochemical failure”), followed by increased tumor size, new metastatic spread, and disease-related symptoms (12).

Recurrent, “castration-resistant” cancers, or CRPC, represent the lethal phenotype of the illness. Considerable effort has been expended to better understand the targets and mechanisms contributing to progression, with the hope that innovative new approaches can be brought forward. Rising PSA levels, however, serve as an indication that AR activity is inappropriately restored in CRPC (13), a hypothesis that has been solidified by a litany of studies investigating mechanisms of therapeutic failure. These mechanisms have been extensively reviewed elsewhere and include (a) AR amplification and/or overexpression; (b) gain-of-function AR mutations (largely occurring in the ligand-binding domain and conferring ligand promiscuity); (c) intracellular androgen production (thus

Authors’ Affiliations: Departments of 1Cancer Biology, 2Urology, and 3Kimmel Cancer Center, Thomas Jefferson University, Philadelphia, Pennsylvania; 4Genitourinary Oncology Service and 5Department of Medicine, Memorial Sloan-Kettering Cancer Center, New York, New York, 6Department of Medicine, Joan and Sanford E. Weill College of Medicine of Cornell University, New York, New York, and 7Genitourinary Oncology Service, Department of Medicine, Sidney Kimmel Center for Prostate and Urologic Cancers, Memorial Sloan-Kettering Cancer Center, New York, New York

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Requests for reprints: Karen E. Knudsen, Thomas Jefferson University/Kimmel Cancer Center, 233 South 10th Street, Bluemle Building, Room 100B, Philadelphia, PA 19107. Phone: 215-503-8574; Fax: 215-923-4498; E-mail: karen.knudsen@jefferson.edu.

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providing tumor-produced ligand to AR); (d) overexpression of AR coactivators (thus sensitizing cells to low-level ligand); and (e) indirect AR activation via growth factors, cytokines, or aberrant AR phosphorylation (Fig. 1; refs. 14-21). Strikingly, circulating tumor cells isolated from patients with CRPC have evidence of AR amplification in 50% of cases, further supporting AR as a major effector of CRPC (22). Inflammation has also been proposed to indirectly negate the inhibitory effects of AR antagonists through molecular cascades that convert AR antagonists into agonists (23). Most recently, it was shown that AR mRNA can undergo alternative splicing events that delete the LBD, thus producing a constitutively active receptor that does not require ligand and is refractory to current AR antagonists (24, 25). These observations strongly suggest that androgen deprivation initiates a selective process for AR reactivation and resultant CRPC development. Recent clinical trials with novel AR antagonists further credentialed the AR pathway as one of therapeutic relevance. This premise applies to both the chemotherapy naïve setting and the postchemotherapy setting, a point when many tumors are considered to be “hormone refractory” and not amenable to further hormonal manipulations. Novel means to durably inhibit AR therefore are urgently needed, and current advances toward this goal are the focus of this review.

**Clinical-Translational Advances**

Major breakthroughs in the development of novel androgen-ablative and AR antagonist strategies have been recently described and have the potential to improve the efficacy of AR targeting and subsequent therapeutic outcome. As will be discussed, these advances were developed based on substantive evidence that the current standard of practice fails to achieve complete androgen ablation and/or sufficient suppression of AR signaling in the prostate. Parallelizing these findings, advances in understanding of AR biology revealed an unexpected need to develop new classes of AR-targeting agents directed against the N-terminal domain (Fig. 2). The potential utility of these new strategies and the likely impact of combination therapy with AR-directed therapies will be discussed below.
Improving Androgen Depletion

The current standard of care for patients with disseminated disease is treatment with gonadotropin releasing hormone (GnRH) agonists (e.g., leuprolide; refs. 3, 26). After an initial increase in testosterone (which may produce or exacerbate symptoms such as urinary obstruction or pain), sustained GnRH agonists desensitize gonadotropin release and suppress testicular androgen synthesis. These regimens are initially effective at suppressing AR activity (as judged by declines in PSA) and initiating tumor regression. Upon progression, ketoconazole, which blocks adrenal androgens, can be used (28). However, the recent discovery that serum androgen depletion selects for intracrine androgen production (e.g., as achieved by induction of enzymes that convert weak adrenal-derived androgens to testosterone) highlights the transient and/or incomplete efficacy of GnRH agonists to achieve “true” androgen depletion (17-19, 29). Exemplifying this, studies evaluating androgen levels in tissue showed that androgen-depletion therapies reduced intratumoral androgens by only 75% at a time when androgen levels in the sera remained in the castrate range; in these tissues, persistent expression of AR and AR target genes (e.g., PSA and TMPRSS2) validated the concept that the residual androgen is sufficient to sustain AR activity. AR gene amplification can further enhance the ability of the receptor to “adapt” to the environment of low testosterone, in part by using low-affinity ligands (30). These observations underscore the emerging view that current androgen-depletion strategies are incomplete, and that residual androgen contributes to sustained AR activity and disease progression.

A new means to further deplete androgens is provided by a selective CYP17 inhibitor that blocks 17α-hydroxylase and C17,20-lyase enzymes in the adrenal steroid synthetic pathway, abiraterone, which has entered clinical trial (31). Abiraterone inhibits both testicular-derived androgen production and tumor-derived androgen synthesis, and preclinical studies showed that abiraterone principally reduces the weight of androgen-dependent/AR-dependent organs (32). Recent clinical data showed that abiraterone can suppress testosterone in noncastrate patients (33). Whether abiraterone can formally suppress intratumoral androgen production remains to be fully tested (34), but recent phase I trials in patients with CRPC showed significant PSA declines, tumor regression, and palliation of symptoms in patients who had not received chemotherapy (35), as well as in...
the postchemotherapy setting, wherein “hormonal” agents are typically not considered.\(^8\) A phase III trial of abiraterone versus placebo plus prednisone with a primary endpoint of survival is ongoing in postchemotherapy-treated patients. Thus, abiraterone could provide a significant advantage toward the goal of durable androgen depletion and suppression of AR activity. Alternatively, VN/124-1 is a CYP17 inhibitor that both reduces androgen production and, among other effects, can also reduce AR expression levels (36). VN/124-1 has entered clinical trial, and it is hoped that the multiple functions of the drug will provide an advance toward the goal of blocking AR activity. Lastly, estrogens [including diethylstilbestrol (DES)] can be used to suppress both testicular and adrenal androgens (37), but formal trials of adequate size and power to address a survival impact have not been conducted.

GnRH antagonists are also available and could be used as an alternative means for androgen suppression. Like GnRH agonists, these agents prevent testicular androgen synthesis, but do not induce the transient testosterone flair associated with GnRH agonists (38, 39). Administration or use of first-generation GnRH antagonists such as abarelix was limited by untoward side effects (27, 40), but recent modifications to the structure of the product resulted in reduced immunostimulatory activity and safer administration (41). One such second-generation product, degarelix, can suppress circulating androgen levels without inducing testosterone flares or allergic reactions (42, 43). To date, however, GnRH antagonists have not been shown to provide superior antitumor effects relative to a combination of GnRH agonist in combination with an antiandrogen. Past clinical trials attempted to further suppress androgen by combining ketoconazole, hydrocortisone, and a 5-alpha reductase inhibitor, but advantages relative to conventional treatment remain uncertain (44). Whether a combination of a GnRH antagonist and abiraterone, shown to further reduce serum androgen levels below those achieved with GnRH agonist therapy alone, and separately intratumoral androgens, will prove superior requires prospective testing.

Collectively, these observations indicate that contemporary understanding of androgen regulation in prostate cancer has allowed for the development of new means to inhibit AR through ligand depletion. Using this knowledge, ongoing studies and clinical trials could provide a firm foundation upon which to formally test the prevailing hypothesis that more complete androgen deprivation (especially in combination with AR antagonists, discussed below) can enhance response rates and cure.

**AR Antagonists**

Direct AR antagonists are frequently utilized in combination with orchiectomy or GnRH agonists/antagonists, in an effort to further inhibit AR signaling (45). Although the precise mechanisms of action remain a subject of controversy, substantive evidence supports the contention that docking of AR antagonists such as bicalutamide into the AR C-terminal LBD results in both passive AR inhibition (via competition for agonists) and an active mechanism of AR inhibition (e.g., prevention of coactivator binding and/or induced corepressor recruitment; refs. 46, 47). Thus, AR antagonists would be expected to act in concert with androgen deprivation to further suppress AR activity. Clinically, the validity of this supposition remains uncertain. Analyses of long-term outcomes in patients receiving “combined androgen blockade” (androgen deprivation plus an AR antagonist) showed varying results with regard to both overall and progression-free survival (11). Indeed, the question of whether AR antagonists (e.g., bicalutamide) or mixed agonists/antagonists (e.g., flutamide) improve outcome has been one of the most extensively studied questions in the field. Taking into account the methodologic differences in trials (including variances in dose, differences in scheduling, and distinctions in the timing of androgen blockade), the beneficial effects on long-term outcomes proved to be modest at best.

Uneven results obtained with current AR antagonists are attributed, at least partially, to the observation that these agents show relatively low affinity for AR as compared to DHT, and therefore are required in vast excess for molecular efficacy (48). Thus, recent studies have endeavored to improve the underlying basis by which AR antagonists suppress receptor activity, and several new agents are in clinical trial (4). The BMS-641988 compound was shown in preclinical studies to have activity in bicalutamide-resistant cells; however, the drug has been discontinued because clinical findings on safety and efficacy do not support benefit/risk sufficient for further development. A full publication of study results is planned. Alternative strategies emerged through the development of MDV3100, a new AR antagonist identified as active in bicalutamide-resistant cancer models that overexpress AR. Preliminary studies suggest that MDV3100 has no agonist effects, and that it prevents both AR nuclear translocation and DNA binding (49). Given these properties, it may be expected that MDV3100 provides a mechanistic advantage over bicalutamide, and therefore enforce sustained suppression of AR activity. Preliminary phase I/II trial data are highly suggestive (presented at the 2009 ASCO meeting), in that MDV3100 treatment correlated with declining serum PSA, reduced circulating tumor cells, and radiographic disease stabilization. Importantly, MDV3100 was active in both pre- and postchemotherapy-treated patients, and a phase III trial in post-chemotherapy treated patients, AFFIRM, will be initiated later this year. Based on these examples, it is clear that basic knowledge of AR function can foster the development of new approaches for targeting the AR LBD. Whether these compounds will result in meaningful clinical outcomes for CRPC should be shortly revealed, as will the impact of these agents on long-term AR and prostate cancer management.

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**The AR N-terminal Domain: A Cause for Concern?**

Androgen-depletion therapies and direct AR antagonists have commonality, in that both approaches rely on an intact AR ligand-binding domain and stand on the premise that agonist binding to the LBD is universally required for AR activation. Recent analyses suggest that this presumption is likely premature. Pioneering work by Dehm, Tindall, and colleagues showed in model systems that AR can be alternatively spliced so that the C-terminal domain is deleted, rendering production of a receptor that is constitutively active (24). This observation is consistent with previous studies that showed that, unlike most other
nuclear receptors, the predominant transcriptional transactivation function of AR resides in the N terminus, and that deletion of the LBD confers ligand-independent activity (5, 50). Constitutively active splice variants have recently been observed in tumor tissue, wherein it was shown that variants lacking C-terminal residues are overproduced in CRPC, and that these receptors are constitutively active (25). Together, these unexpected observations highlight yet another mechanism by which tumors bypass androgen deprivation and/or AR antagonists, as it is predicted that the splice variants would be refractory to both. Accordingly, it is evident that a new class of AR-inhibitory agents must be developed for successful management of tumors expressing truncated AR, wherein even complete androgen ablation would have no effect on receptor activity.

Several options for potentially suppressing the function of C-terminal-deficient ARs may already exist. It has been suggested that either HSP90 inhibitors (e.g., geldanamycin or analogs) or agents that modulate HSP90-HDAC interactions (e.g., genistein) may reduce overall AR levels (51, 52), and could potentially be used to suppress the action of both full-length and truncated AR. Knockdown strategies have also been proposed, as siRNA directed against AR suppresses prostate cancer growth in model systems (53); however, such strategies are hindered by the uncertain feasibility of using siRNAs for cancer therapy. Alternative means to thwart AR function by using expressed peptides have been documented in proof-of-principle studies, wherein “decays” of the AR N terminus were shown to suppress cell growth and survival (54), as has expression of corepressor domains that target AR N-terminal transactivation function (55). Although translating such observations to the clinical setting remains a major challenge, the observation that CRPC tumors can express truncated, androgen-deprivation and AR antagonist-resistant receptors is a cause for concern, and underscores the need for the intensive development of strategies to target the AR N terminus.

Combination Therapies: What Does the Future Hold?

Because prostate cancers utilize a multitude of genetic alterations to restore AR activity and tumor growth under conditions of androgen deprivation and/or combined androgen blockade, the development of successful combinatorial therapies will likely be required to eliminate disease and prevent recurrence. Radiation therapy in combination with androgen depletion can improve response in locally advanced disease (34) and provides benefit for this subset of tumors. Docetaxel can extend survival in patients with CRPC (56); however, the benefit is modest, with an average extension of only 2-3 months. It has been hypothesized that the scheduling of docetaxel in combination with AR-antagonizing strategies may need refinement (57). This poset was initially supported by the observation that cancer cells that survive AR-inhibitory strategies accumulate in the G1 phase of the cell cycle (10), whereas docetaxel acts predominately in later phases (G2/M) to induce cell death. The supposition that concurrent administration of AR-antagonizing strategies with docetaxel may impede the cytotoxic effects of the chemotherapeutic was validated in models of androgen-dependent cancer (58) and is further supported by clinical data that showed an improved response to docetaxel in the presence of androgen (59). These observations provide the impetus for re-examining how AR- ablative therapies might be optimized in combination with antimitotics and emphasize the importance of considering AR biology in the design of combinatorial therapeutic approaches. Furthermore, it should be considered under which conditions AR acts as a survival factor to counteract chemotherapeutic response, and how combination therapy could be optimized to suppress AR-associated survival activity.

Other combinations yet to be rigorously considered include abrogation of growth factor or transcriptional regulatory pathways that contribute to ligand-independent AR activity. Several growth factor and cytokine pathways, including FGF, IGF, EGF, IL-6, and heregulin, were shown in model systems to facilitate AR activity in the presence of no or low androgen and are therefore preliminarily implicated in disease progression (15, 20). Theoretically, antagonists of these pathways could, in some cases, act in concert with androgen ablation and/or AR antagonists to further suppress AR activity, tumor growth, and the development of CRPC. A question to be addressed is whether growth factor pathways are upregulated as a survival mechanism after androgen depletion, so as to determine how potential combinations of AR-directed therapies with growth factor receptor antagonists would be most effective. For all proposed combinations, it will be imperative to also consider the tumor microenvironment, as tumor cells residing near activated stroma and/or neuroendocrine cells that supply growth factors may show a differential response to therapy (60). Lastly, it has been recently shown that AR may require HDACs for transcriptional activation (61), and that HDAC inhibitors may cooperate with AR-directed therapeutics to elicit an enhanced cellular response (62). The combination of HDAC inhibitors with GnRH analogs is the focus of an ongoing InterProstate Cancer SPORE neoadjuvant trial.

Future Directions

The contribution of AR to prostate tumorigenesis and disease progression is incontrovertible. Characterization of CRPC at the molecular level and in model systems has validated the concept that AR activity is regained as part of disease progression. Until recently, the development of innovative new strategies for durable suppression of AR activity has been modest. Although ligand depletion or the use of C-terminal-binding receptor antagonists can induce tumor remission, these strategies do not provide a means by which to sustain suppression of AR activity and do not completely eliminate the tumor. Under prospective study is whether more complete inhibition of AR signaling (e.g., through the combination of a GnRH agonist with an HDAC inhibitor) can completely abrogate AR activity. As the degree of androgen dependence may vary, characterization of the surviving cell population in the neoadjuvant setting may provide important new insights into the mechanisms of resistance and points of therapeutic attack.

At present, new understandings of AR function during disease progression have already led to potential breakthroughs in the development of novel AR antagonists and ligand-depletion strategies. Although it is hoped that these agents will be of clinical benefit, several hurdles remain. First, it should be determined how new agents function under disparate conditions of AR reactivation, and whether patient stratification based on these criteria would be of benefit. For example, if recurrence is associated with AR mutations or splice variants that induce resistance to AR antagonists, it is unlikely that use of these agents
would be of benefit. A notable advance toward this end is the development of mechanisms to characterize CRPC at the molecular level by using circulating tumor cells. This innovation is expected to provide needed new insight into the mechanisms governing castration resistance, and could provide a basis for personalized medicine (49). Second, new strategies to target the AR-N-terminal domain are needed, given the recent observations of recurrence-associated, C-terminal-deficient AR splice variants. Third, mechanisms to destabilize AR or required cofactors would be expected to produce a marked improvement in the durability of response and should be prioritized for development. Fourth, given advances in the understanding of AR-dependent cell cycle control, it should be considered how use of AR-ablative strategies might be more effectively combined with existing antimitotics to improve outcome. Finally, it should be considered that even if achieved, "durable" AR ablation may not provide a cure, as tumor cells are likely to adapt to true AR inhibition through the development of secondary dependencies or signaling pathways, and it remains possible that putative prostate cancer stem cells would be therapy resistant. Thus, it is imperative to identify alternative targets that may act in concert with AR antagonists. Through these collective strategies, it is hoped that the goal of sustained AR management will lead to predicted improvements in clinical response and reduce death from prostate cancer.

Disclosure of Potential Conflicts of Interest

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