Preclinical and Coclinical Studies in Prostate Cancer

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Men who develop metastatic castration-resistant prostate cancer (mCRPC) will invariably succumb to their disease. Thus there remains a pressing need for preclinical testing of new drugs and drug combinations for late-stage prostate cancer (PCa). Insights from the mCRPC genomic landscape have revealed that, in addition to sustained androgen receptor (AR) signaling, there are other actionable molecular alterations and distinct molecular subclasses of PCa; however, the rate at which this knowledge translates into patient care via current preclinical testing is painfully slow and inefficient. Here, we will highlight the issues involved and discuss a new translational platform, “the co-clinical trial project,” to expedite current preclinical studies and optimize clinical trial and experimental drug testing. With this platform, in vivo preclinical and early clinical studies are closely aligned, enabling in vivo testing of drugs using genetically engineered mouse models (GEMMs) in defined genetic contexts to personalize individual therapies. We will discuss the principles and essential components of this novel paradigm, representative success stories and future therapeutic options for mCRPC that should be explored.

Prostate cancer (PCa) is the third leading cause of cancer-related deaths in U.S. men. Although localized PCa is highly curable, metastatic castration-resistant PCa (mCRPC) is still a fatal disease (Wu et al. 2014). Preclinical studies implicating dysregulated androgen receptor (AR) signaling as the predominant driver of mCRPC have propelled the development of novel agents targeting the AR signaling (Chen et al. 2004). Despite the effectiveness of recently approved AR-targeted therapies, such as abiraterone and enzalutamide, de novo or acquired drug resistance invariably emerge and pose the largest obstacle to implementing curable thera-

py (Watson et al. 2015). Thus, there remains a pressing need for preclinical testing of new drugs and novel drug combinations that are tailored to target potent driver mutations and prevent resistance in mCRPC.

Integrative genomic analyses have shown that PCa displays significant inter- and inpatient heterogeneity, not only in the context of the mutational landscape but also in epigenetic deregulation and aberrant transcriptome of non-coding elements (Boutros et al. 2015; Cooper et al. 2015; Gundem et al. 2015). It is now widely understood that defining the genetic makeup of individual patient tumors is likely to have a sub-

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stantial impact on the effectiveness of their treatment, a concept referred to as precision medicine (Collins and Varmus 2015). The recent development of powerful methods for characterizing patients, particularly high-throughput “omics” analysis, and large-scale biologic databases are now enabling researchers to categorize and stratify patients according to genetic alterations, and in doing so, to distinguish different subsets of human PCa, how these genetic make-ups dictate tumor aggressiveness, and likely response to the standard-of-care (SOC) and/or experimental therapies, ultimately leading to precise and personalized therapies. Indeed, stratification of patients with defined genetic defects has recently shown great promise for targeted therapies in PCa (Mateo et al. 2015). However, this opportunity is not without its own challenges, particularly concerning the design of clinical trials. With increased PCa diversity and complexity, one major hurdle is the stark decrease in the size of patient populations available for enrollment in appropriate clinical trials. This, in turn, makes it difficult to obtain robust statistical analyses, and prolongs the time that it takes new and/or combinatorial therapies to be evaluated.

In recognition of these challenges, we are now compelled to rethink the platforms that are required to rapidly analyze in model systems current and novel therapeutic options for patients to ensure rapid implementation of personalized approaches. Preclinical studies using accurate genetically engineered mouse models (GEMMs) of specific cancer subtypes have proven extremely valuable; indeed, they have already guided and optimized the treatment of several forms of human cancer (Van Dyke and Jacks 2002; Abate-Shen and Pandolfi 2013). However, the traditional “GEMMs-to-human” or “pre-clinical-to-clinical” approach remains painfully slow because data have to be accrued in the mouse model of interest first, and then translated to be able to inform the clinical trials that follow. Importantly, current preclinical methods to predict successful outcomes are extremely limited because only about 5% of new cancer drugs are approved after preclinical studies (Sharpless and Depinho 2006). Given that we

are able to model cancer in mice at an unprecedented level of accuracy and that a large amount of new experimental drugs are currently in early-phase clinical studies, we have implemented a coclinical trial project to align preclinical efforts in GEMMs with clinical trial efforts in humans to optimize and refine phase I/II clinical studies in real time and expedite effective drug approval for personalized therapies.

In this review, we discuss the principles and essential components of the coclinical trial project, with a focus on how the implementation of such a platform can accelerate the development of personalized medicine for PCa. We also summarize what we have learned thus far by using this platform and future therapeutic options for mCRPC that have emerged, and should be explored, on the basis of these efforts.

THE COCLINICAL TRIAL PROJECT FOR THE DEVELOPMENT OF PERSONALIZED THERAPIES

The Approach

The coclinical trial project rests on the notion that (1) cancer develops in many distinct genetic subtypes, (2) this diversity impacts how these tumors respond to or resist various forms of therapy, and (3) that preclinical testing of new drugs and drug combinations in faithful GEMMs of human cancer is a powerful and predictive platform for clinical trial optimization. The prospect of implementing this approach has been significantly improved by the advance of powerful “omics” methods for characterizing patients and our ability to model cancer in mice at an unprecedented level of accuracy. The keys to this approach are “synchronicity” and “integration.” Specifically, by enrolling as many GEMMs representative of the genetic diversity of a given tumor type as in which human trial is performed, and treating GEMMs and human patients in exactly the same way, the relevant clinical, biological, and pharmacologic data (i.e., sensitivity vs. resistance to specific regimens, imaging analysis, microarray and proteomic profiling, somatic mutation, nucleotide polymorphisms, etc.) from both sources are an-

alyzed in parallel, shared, and integrated in real time. The synchronicity and the integration of the approach in turn credential and validate, in real time, what is found in human and mice, thereby facilitating the design of subsequent clinical studies through more timely and informed clinical decisions (Fig. 1).

The coclinical trial project can significantly lessen the gap between preclinical study and patient care via (1) the rapid stratification of patient sensitivity and resistance to a specific treatment on the basis of molecular and genetic criteria, (2) characterization of mechanisms dictating de novo or acquired tumor resistance to therapy, and (3) the use of this platform as a testing ground for new drug combinations (Fig. 1). Instead of attempting to provide a detailed review for the coclinical trial project, here we will focus on the essential components rele-

vant to PCa and refer the reader to previous articles that have addressed specific topics and/or cancer types in depth (Nardella et al. 2011; Chen et al. 2012; Lunardi et al. 2013; Kwong et al. 2015; Whitley et al. 2016).

GEMMs as *In Vivo* Models to Inform Patient Stratification

Mice rarely develop spontaneous PCa; however, they can be genetically or environmentally modified to develop PCa, and are a compelling model system for the *in vivo* testing of new drugs and drug combinations. Implant of cancer cell lines (xenografts) or patient-derived xenografts (PDXs) into heterologous sites of immunodeficient mice represent additional ways to study human cancer in mouse models. These models can also be used “coclinically.” However, cell

Figure 1. The coclinical trial platform. (A) In the traditional clinical trial, little attention is paid to the molecular characteristics of the disease. When a minor fraction of patients are responsive, the overall response of the population might mask the responders. (B) In the coclinical trial project, relevant genetically engineered mouse models (GEMMs) and human patients are treated with the same drug and clinical protocol. Integrated analyses of data accrued in GEMMs and patients serve to stratify responsiveness and resistance on the basis of molecular and genetic criteria. Mechanisms underlying acquired resistance are also rapidly identified, and drug combinations to overcome such resistance are tested in GEMMs for their effectiveness.

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lines or PDXs do harbor multiple concurrent genetic alterations at once, which makes it difficult to determine the impact of specific mutations in modulating drug or drug combination response. Additionally, these models are immunodeficient and hence not amenable to test cancer immunotherapies. For more information on these topics, we refer you to Risbridger et al. (2017). Central to the coclinical trial project are faithful GEMMs to steer parallel clinical studies in patients. There are numerous advantages in using GEMMs coclinically. First, GEMMs develop tumors *de novo* in the context of the native tissue environment and intact immune system. This criterion is paramount for the study of cancer treatment, particularly when considering the role of the tumor stroma in resistance to therapy and the potential for use of immunotherapy in the treatment of PCa (Kantoff et al. 2010; Boelens et al. 2014). Second, cancer phenotypes in GEMMs arise from well-defined genetic alterations that have been known to faithfully recapitulate human cancer. For example, many GEMMs of PCa are based on loss of function of *Pten*, which is among the most frequently lost or mutated tumor suppressors in human PCa (Taylor et al. 2010; Barbieri et al. 2012; Grasso et al. 2012; Robinson et al. 2015). Moreover, the specific genetic alterations enable us to simplify and deconstruct the precise genetic elements underlying resistance or sensitivity to a given treatment modality. Third, GEMMs enable both longitudinal and molecular analyses of cancerous lesions that can lead to identification of novel biomarkers of response and facilitate formulation of new combinatorial therapeutic modalities. Fourth, GEMMs can be readily expanded to enable more subjects, drugs, and drug combinations to be tested in the coclinical setting. This has shown to be of great value, in particular to the study of genetically rare cancer subtypes of which sizable patient accrual represents a major hurdle.

Like other types of cancer, PCa is characterized by a wide range of genetic complexity, such as loss and mutation of tumor suppressor genes (e.g., *PTEN*, *NKX3.1*, *P53*, *SPOP*, *SMAD4*, *LRF*, *PLZF*, and *BRCA1/2*), amplification and mutation of oncogenes (e.g., *c-MYC*, *PI3KCA/B*, and

AR), and fusion genes (e.g., *ERG/ETV1/ETV4*). In recent years, this disheartening complexity has been successfully modeled in GEMMs (Irshad and Abate-Shen 2013), allowing for the generation of a unique genetic platform of proxy-PCa in the mouse, which has proven to be essential in elucidating the role of specific genetic perturbations in the onset and progression of human PCa. Importantly, substantial efforts have been devoted to systematically applying this resource in the preclinical/coclinical setting to test new drugs and drug combinations for targeted therapies, to dissect the mechanisms underlying *de novo* or acquired tumor resistance to SOC treatments, and to discover new treatment combinations (Carver et al. 2011; Lunardi et al. 2013; Schwartz et al. 2015; Yamamoto et al. 2015). However, challenges still remain in the development of these models. Notably, PCa is made up of complex genetic landscapes, and we currently have a limited ability to model tumor heterogeneity. Moreover, metastasis has proven to be particularly challenging to model, and the progression to metastasis from either a primary indolent or advanced-stage disease is rarely observed for the majority of the GEMMs of PCa (Ittmann et al. 2013), but we can envisage that technological developments such as CRISPR-Cas9 (clustered regularly interspaced short palindromic repeats-CRISPR-associated 9) technology (Hsu et al. 2014) would facilitate modeling PCa genetic diversity, complexity, and metastasis in GEMMs and address these challenges. Despite the current limitations, the use of GEMMs is a powerful tool for both preclinical and coclinical evaluation and fine-tuning of treatment protocols in specific genetic make-ups precisely for its ability to simplify and deconstruct the critical genetic elements dictating differential responses to treatment.

Components and Requirements for the Coclinical Trial: The Mouse Hospital and Beyond

The simultaneous execution of preclinical trials in mice and phase I/II trials in humans is incredibly resource-intensive (Fig. 2). To ensure the coclinical trial project success, a “mouse hospi-

Figure 2. Fundamental components of the coclinal effort and their relationship to infrastructure and pharmaceutical companies. To ensure that preclinical and coclinal trials are most effective, mouse trials are performed in a manner and environment similar to those for human patients. Infrastructural and operational procedures need to be in place to ensure that data are readily transferable to the clinic. Acquisition of relevant therapeutic agents for coclinal evaluation is coordinated by the mouse pharmacy, which might obtain agents directly from industry or independently coordinated synthesis of relevant agents.

tal” with a state-of-the-art infrastructure for in vivo mouse imaging and a mouse pharmacy is required for a consistent and reproducible pipeline to be maintained. This hospital-type animal facility is staffed by experts in levels of mouse care (i.e., husbandry, treatments, and surgery) with access to procedure rooms and small animal in vivo–imaging technologies (bioluminescence, magnetic resonance imaging, micro-computed tomography, micro-positron emission tomography, ultrasonogram, fluorescence-mediated tomography), as well as instruments needed for blood and urine tests. All of the above are fundamental to determining local and systemic tumor responses to treatments in GEMMs. Moreover, several other important elements are also required for integration and implementation of mouse–human coclinal trials (such as bioinformatics consortia, comparative pathology centers, and central databases). Even further, to maintain consistent workflows and generate reproducible and reliable high-quality results, a system of work needs to be established, ensuring that procedures, data collection, and informatics analyses are performed consistently throughout the mouse hospital and among mouse hospitals. This is essential to be able to

share and compare mouse data across different mouse hospitals and human clinical centers. Most important, this platform requires a sustained and coordinated effort between basic scientists and clinicians (Fig. 2).

The Coclinal Trial Efforts in PCa

Proof-of-principle examples of the efficacy of the coclinal trial platform have been recently reported in PCa (Lunardi et al. 2013). The mainstay of therapy for advanced and metastatic PCa for more than 70 years has been androgen-deprivation therapy (ADT) (Huggins and Hodges 1941). Despite the majority of patients experiencing an initial positive response to ADT, most tumors eventually acquire resistance to this treatment and, as a result, patients inevitably relapse to a lethal stage of the disease. However, there is a wide variability in the durability of ADT efficacy, with responses varying from months to years. Not much is known about the possible determinants of ADT response. Given the genetic heterogeneity of PCa, we have hypothesized that different genetic lesions driving the tumorigenic process may also affect ADT efficacy.

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Accordingly, in a coclinical trial effort, we have enrolled a cohort of 84 PCa patients and a panel of GEMMs bearing common human PCa genetics ($Pten^{pc-/-}$; $Pten^{pc-/-}p53^{pc-/-}$; $Pten^{pc-/-}Lrf^{pc-/-}$), in which we rapidly identified $Pten$ - $p53$ and $Pten$ - Lrf double-null tumors in mice as resistant to ADT. We anticipated the response of patients with PCa characterized by the same genetic lesions using these coclinical data sets. Importantly, a comparative genetic and molecular analysis of $Pten$ -null (ADT-sensitive) and $Pten$ - $p53$ and $Pten$ - Lrf double-null (ADT-resistant) mouse prostate tumors showed up-regulation of the steroid-5- α reductase type 1 gene (*Srd5a1*) and deregulation of the X-linked inhibitor of apoptosis protein (XIAP)-associated factor 1(Xaf1)/XIAP apoptotic pathway as two potential mechanisms of ADT resistance. A double combination of embelin (a natural XIAP inhibitor) and dutasteride (an SRD5A1 inhibitor currently being studied in clinical trials for ADT-resistant PCa, ClinicalTrials.gov identifier: NCT01393730, NCT00673127, NCT00553878, NCT00403000), formally proved the functional role of deregulation of both pathways in ADT resistance by rescuing ADT efficacy in vivo in both $Pten$ - $p53$ and $Pten$ - Lrf double-null mouse prostate tumors and in vitro in human PCa cell lines. This, in turn, paved the way for the clinical testing of such a combinatorial approach, and of novel predictive biomarkers and therapeutic targets for ADT resistance in men. Coclinical efforts have by now also shown significant promise toward a real-time optimization of patient stratification in the effective treatment of several other types of cancers (Chen et al. 2012, 2014; Kwong et al. 2015; Whitley et al. 2016).

TACKLING CHALLENGES AND OPPORTUNITIES FOR PRECISION MEDICINE IN PCa THROUGH COCLINICAL EFFORTS

The Clinical Challenges

The survival of mCRPC patients has been improved substantially by the use of abiraterone and enzalutamide as single-agent therapies

against the AR pathway (Scher et al. 2012; Ryan et al. 2013; Beer et al. 2014). Unfortunately, to date, no single agent has been able to cure PCa on its own, largely owing to tumor heterogeneity and the adaptive responses that support the development of de novo or acquired drug resistance. Another challenge is the relief of profound feedback inhibition of multiple oncogenic pathways triggered by anti-AR drugs that limit the success of targeted therapies (Chandarlapaty 2012). As such, it is therefore essential to be able to continue to identify additional effective strategies to combine with SOC treatment toward a curative regimen.

Therapeutic Opportunities for Personalized Therapies in mCRPC

Recent mCRPC genomic landscape studies have identified multiple molecular alterations in other actionable pathways, aside from AR, that drive mCRPC progression (Fig. 3), including PI3K/AKT/PTEN, RAF, DNA repair, WNT, cell cycle, and cellular metabolism (Taylor et al. 2010; Grasso et al. 2012; Robinson et al. 2015). To this end, the coclinical trial using GEMMs is an ideal investigational platform to rapidly test drug combinations targeting these molecular lesions in different genetic makeups to ensure rapid implementation of personalized treatment. Here, we highlight future therapeutic opportunities for mCRPC that should be further explored through using the coclinical platform, and, for information regarding AR-targeted therapies, we refer you to Centenera et al. (2017).

The Phosphoinositide-3 Kinase (PI3K)/AKT/Mechanistic Target of Rapamycin (mTOR) Pathway

The PI3K/AKT/mTOR pathway is altered in ~50% of mCRPC through either *PTEN* inactivation or/and aberrant activation in *PIK3CA/B* (Taylor et al. 2010; Grasso et al. 2012; Robinson et al. 2015). Several studies have also shown cross talk between the AR pathway and PI3K signaling, in which these pathways regulate each other by reciprocal negative feedback, such that inhibition of one activates the other,

Figure 3. Therapeutic targets in metastatic castration-resistant prostate cancer (mCRPC). AR, Androgen receptor; CDK, cyclin-dependent kinase; PARP, poly(ADP-ribose) polymerase; AMPK, AMP-activated protein kinase; FASN, fatty acid synthase.

therefore providing rationale for combination therapy in mCRPC patients (Carver et al. 2011; Schwartz et al. 2015). Phase I/II trials assessing the combination of next-generation AR therapy with a PI3K/AKT/mTOR inhibitor are currently ongoing (ClinicalTrials.gov identifier: NCT02407054 and NCT02215096).

Ras/Raf/MEK/ERK Mitogen-Activated Protein Kinase (MAPK) Pathway

The MAPK signaling pathway is often found aberrantly activated in advanced metastatic PCa (Kinkade et al. 2008; Taylor et al. 2010). However, little is known about the underlying molecular mechanisms leading to MAPK activation, because activating mutations or gene rearrangements involving the MAPK cascade are rare in human PCa. Nevertheless, aberrant KRAS/B-RAF/RAF1 fusions have been identified in 3% of mCRPC, representing a rare subset of mCRPC (Palanisamy et al. 2010; Wang et al. 2011; Beltran et al. 2013; Robinson et al. 2015). Previous studies have shown that expression of RAF fusion protein in prostate cells induces a neoplastic phenotype that is sensitive to RAF and MEK inhibitor, suggesting that tumors

from mCRPC patients harboring RAF fusion might be sensitive to pan-RAF inhibitors or MEK inhibitors (Palanisamy et al. 2010). Because the available cohort of mCRPC patients with RAS/RAF fusion is small, the coclinical trial platform with faithful GEMMs in this rare subtype of PCa will be able to better evaluate targeted combinatorial therapies. Notably, the PI3K/AKT/mTOR and ERK-MAPK signaling pathways have been shown to cooperate in PCa progression and contribute to the development of castration-resistant and metastatic PCa (Gao et al. 2006). On this basis, dual targeting of the PI3K/AKT/mTOR and ERK-MAPK signaling has been proven highly effective in suppressing mCRPC in preclinical studies (Kinkade et al. 2008; Floc'h et al. 2012; Toren et al. 2016).

DNA Repair Pathway

Another common subset of mCRPC is characterized by defects in DNA repair, accounting for 20%–30% of sporadic mCRPC, as exemplified by germline and somatic mutations in genes that encode proteins in DNA repair pathways, such as *BRCA1/2* and ataxia-telangiectasia mutated (*ATM*) (Grasso et al. 2012; Beltran et al. 2013;

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Robinson et al. 2015). *BRCA*-defective cells are unable to repair double-strand DNA breaks by homologous recombination, and thus become highly sensitive to inhibitors of the DNA repair enzyme, poly(ADP-ribose) polymerase (PARP), suggesting that treatment with a PARP inhibitor may exploit a synthetically lethal interaction. In a single-agent phase II trial, one such inhibitor, olaparib, led to a high response rate in mCRPC patients whose cancers were defective in DNA repair pathways (Mateo et al. 2015). Interestingly, not only the aberrations in DNA repair pathway, but also the overexpression of *ETS*, *PTEN* loss, or *SPOP* mutations have been associated with sensitivity to PARP inhibitor in human PCa (Brenner et al. 2011; Gonzalez-Billalabeitia et al. 2014; Boysen et al. 2015). Considering the potential use of olaparib in combinatorial therapies against various common genetic alterations in mCRPC, the coclinical platform should be used to explore responsiveness and sensitivity within distinct molecular subclasses of PCa (see the example of Gonzalez-Billalabeitia et al. 2014).

Wnt Signaling

Wnt signaling is one of the key oncogenic pathways in multiple types of cancer and is altered in 18% of mCRPC (Grasso et al. 2012; Robinson et al. 2015). Dysregulation of WNT signaling occurs through mutations of *APC* and *CTNNB1*, overexpression of Wnt ligands, such as *R-spondin 2/3*, *R-spondin* fusions, mutations of the R-spondin coreceptors *RNF43/ZNFR3*, or aberrant *SOX9* activation. Recent studies have shown that cancer cells with aberrations in *RNF43*, *ZNRF3*, or *RSPO2/3* (overall 5% of mCRPC patients) or *SOX9* overexpression, have a high response rate to the porcupine inhibitor (Liu et al. 2013; Ma et al. 2016), suggesting that a subset of mCRPC patients would benefit from Wnt-targeted therapies. In addition, R-spondins are emerging as an important cancer target (Storm et al. 2016). Therefore, targeting the R-spondin pathway in mCRPC tumors harboring Wnt pathway alterations or specifically R-spondin fusions may be worthy of further investigations in preclinical and coclinical settings.

Cell-Cycle Control and Prosenescence Therapy

The cyclin/cyclin-dependent kinase (CDK)/retinoblastoma (RB) axis is a critical modulator of cell-cycle entry and is aberrant in more than 20% of mCRPC. Cell-cycle derangement, such as through *CCND1* amplification or *CDKN2A/B* loss, results in enhanced response to CDK4/6 inhibitors in preclinical mCRPC models (Comstock et al. 2013), suggesting that the selective CDK4/6 inhibition as a potential therapeutic regime in mCRPC warrants further examination to evaluate its clinical efficacy in a coclinical setting.

By modulating the cell-cycle machinery and status, prosenescence therapy for tumor prevention and suppression can also be developed. Cellular senescence has emerged as an intrinsic tumor-suppressive mechanism, which can be enhanced by stabilizing p53, Rb, p27, and induction of *PTEN*-loss-induced senescence (PICS) (Chen et al. 2005; Takahashi et al. 2006; Alimonti et al. 2010). Enhancing the prosenescence response in PCa tumors has had a substantial therapeutic effect in preclinical studies (Alimonti et al. 2010; Lin et al. 2010). A recent study has also shown that a population of CD11b⁺Gr-1⁺ myeloid cells from the innate immune system protect a fraction of proliferating *Pten*-null mouse prostate tumors from senescence, thus sustaining tumor growth (Di Mitri et al. 2014). Therefore, it remains to be further determined in what other genetic makeups prosenescence can be engaged, and what types of targeted therapies and/or immune therapies can be used to enhance cellular senescence on the one end, and to morph it into full-blown apoptosis on the other end.

Cellular Metabolism

Cancer cells are known to undergo metabolic reprogramming to support their rapid proliferation and growth. It is now well known that metabolic reprogramming is a hallmark of cancer cells (Hanahan and Weinberg 2011). Unlike other solid tumors, the majority of PCa does not show classical aerobic glycolysis; rather, *de novo*

fatty acid and cholesterol synthesis is the hallmark of human PCa (Zadra et al. 2013). Many lipogenic enzymes are found deregulated in PCa, such as fatty acid synthase, acetyl-CoA-carboxylase, and HMG-CoA reductase (Zadra et al. 2013). Additionally, it is widely postulated that a high-fat/cholesterol Western diet can promote PCa progression, and contributes to the aggressiveness of disease (Gronberg 2003; Yang et al. 2015), in support of the notion that the metabolism of lipids, whether exogenous or endogenously produced, contributes to prostate tumorigenesis and its progression. Importantly, these lipogenic enzymes can be suppressed/inhibited by small pharmacological molecules and hence represent ideal therapeutic targets. Indeed, this metabolic dimension has been explored in preclinical studies (Kridel et al. 2004; Zadra et al. 2014) and should be now investigated coclinically.

CONCLUDING REMARKS

In conclusion, the coclinical trial approach provides multiple new opportunities and tools to accelerate and facilitate the entire process of drug testing toward accelerated drug positioning and approval for precise and personalized medicine. It allows researchers to disentangle the daunting clinical complexity of cancer. In light of the potential of immunotherapy to fulfill the promise of personalized cancer treatment, future efforts will also focus on the immune system in mice, the tumor microenvironment, and how they are impacted by the genetic makeup of the GEMMs used. Integration of preclinical with clinical data in real time, which represents the cornerstone of the coclinical trial platform, promises to quickly yield important insights into issues of resistance and sensitivity, and can provide predictive guidance for both the optimization of concurrent human clinical trials and the correct interpretation of trial results. It is important to note that a specific characteristic that enables the coclinical trial approach to be so effective is its power to identify the mechanisms of tumor resistance in humans (either de novo or acquired), a common hurdle that affects the vast majority of single-agent targeted therapies

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and quickly tests the efficacy of new drug combinations to overcome such resistance.

With the evolving landscape of cancer research and therapy for mCRPC along with our unprecedented ability to model cancer in the mouse, further bolstered by the advent of CRISPER-Cas9 technology (Hsu et al. 2014), coclinical trials will greatly help in testing novel treatments. Many oncogenic pathways involved in cell survival, in addition to the AR pathway, are actionable and can be suppressed/inhibited by current available inhibitors. Immune therapies are also coming of age. There is little doubt that, in the years to come, the coclinical trial platform and the concept of the mouse hospital will be critical in designing new personalized combinations toward the prevention and effective treatment of mCRPC.

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