Combined Tumor Suppressor Defects Characterize Clinically Defined Aggressive Variant Prostate Cancers

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Abstract

Purpose: Morphologically heterogeneous prostate cancers that behave clinically like small-cell prostate cancers (SCPC) share their chemotherapy responsiveness. We asked whether these clinically defined, morphologically diverse, aggressive variant prostate cancer (AVPC) also share molecular features with SCPC.

Experimental Design: Fifty-nine prostate cancer samples from 40 clinical trial participants meeting AVPC criteria, and 8 patient-tumor derived xenografts (PDX) from 6 of them, were stained for markers aberrantly expressed in SCPC. DNA from 36 and 8 PDX was analyzed by Oncoscan for copy number gains (CNL) and losses (CNL). We used the AVPC PDX to expand observations and referenced publicly available datasets to arrive at a candidate molecular signature for the AVPC.

Introduction

Understanding the therapeutically relevant biologic heterogeneity of common adult solid tumors led to the successful introduction of molecularly targeted agents. This is most notable in breast cancer (ER and HER2 overexpression; refs. 1, 2), lung cancer (EGFR mutations and EML4-ALK translocations; refs. 3, 4), melanoma (BRAF mutations; ref. 5), and colon cancers (RAS mutations; ref. 6). In prostate cancer, however, the prevalent “prognostic model” based on clinical features continues to frame clinical decision making (7). This can be attributed in part to the perception that the majority of men with prostate cancer uniformly benefit from androgen ablation and, perhaps more so, to the limited access to metastatic tissue needed for clinical, pathologic, and molecular correlations. However, the diverse responses to recently approved therapies with distinct mechanisms of action (e.g., second-generation androgen biosynthesis inhibitors) has exposed the biologic heterogeneity of prostate cancer and illustrates the urgent need to apply our understanding of prostate carcinogenesis to clinical decision making.

To overcome the challenge of tissue acquisition in metastatic prostate cancer and gain insight into the various clinical phenotypes observed, we initiated a comprehensive approach for tissue collection, clinical annotation, and development of patient–derived xenografts (PDX). Here we describe the molecular characterization of a subset of prostate cancer with aggressive clinical behavior. Virulent prostate cancers with atypical (variant) clinical features have long been associated with small-cell prostate carcinoma (SCPC) morphology (8). More recently, we extended these observations in a prospective study (NCT00514540) that showed that aggressive prostate cancers of various morphologies sharing clinical features with the SCPCs (termed anaplastic prostate cancers, AVPCs) also displayed a high responsiveness to chemotherapy (9). The shared clinical and chemotherapeutic response profiles led us to hypothesize that these morphologically heterogeneous cancers share an underlying biology. In parallel, we had observed that

Results: Irrespective of morphology, Ki67 and Tp53 stained ≥10% cells in 80% and 41% of samples, respectively. RB1 stained <10% cells in 61% of samples and AR in 36%. MYC (surrogate for 8q) CNL and RB1 CNL showed in 54% of 44 samples each and PTEN CNL in 48%. All but 1 of 8 PDX bore Tp53 missense mutations. RB1 CNL was the strongest discriminator between unselected castration-resistant prostate cancer (CRPC) and the AVPC. Combined alterations in RB1, Tp53, and/or PTEN were more frequent in the AVPC than in unselected CRPC and in The Cancer Genome Atlas samples.

Conclusions: Clinically defined AVPC share molecular features with SCPC and are characterized by combined alterations in RB1, Tp53, and/or PTEN. Clin Cancer Res; 1–11. ©2015 AACR.
Translational Relevance
Prostate cancer is a heterogeneous disease yet available therapies continue to be applied homogeneously. Biomarkers that identify the clinically meaningful, therapeutically relevant prostate cancer subsets are needed to allow for optimal therapy development. Here we show that the aggressive variant prostate cancer (AVPC), a clinically defined subset of the disease that shares virulent and atypical clinical features and chemotherapy sensitivity with the small-cell prostate carcinomas (SCPC), are morphologically heterogeneous but can be identified by common molecular features (also shared with SCPC) and should be considered as a distinct clinical–biologic subset of prostate cancer with therapeutic implications. A candidate molecular signature consisting of joined alterations in RB1, Tp53, and/or PTEN is proposed which may account for the shared clinical features, resistance to AR inhibition and chemotherapy sensitivity and should serve as the foundation for a biologically defined therapeutically relevant classification of prostate cancer.

PDXs generated from cancers of men meeting AVPC criteria were frequently AR-negative and bore alterations in tumor suppressors, as well as increases in the expression of mitotic genes and proneural transcription factors (in line with others’ observations; refs. 10–13). Here we combined our findings in PDX models with the clinically annotated NCT00514540 tumor tissues to more completely characterize AVPC and to identify markers of their underlying biology that can be deployed clinically to predict and prognosticate.

Materials and Methods
Patient tumor samples
We collected all available specimens stored in the MD Anderson Department of Pathology from patients registered (n = 121) to the clinical trial, a phase II study of caboplatin plus docetaxel in patients with anaplastic prostate carcinoma (NCT00514540). To be eligible, men with metastatic castration–resistant prostate cancer (mCRPC) had to display ≥1 of the “anaplastic” or AVPC eligibility criteria (Supplementary Table S1; ref. 9). Institutional review board (IRB)-approved written consent was obtained from participants before enrollment for the collection, banking, and use of tumor samples in research. IRB approval was obtained separately for the analyses performed in this study and are available on the UT MDACC Research.

A total of 114 patients were eligible for the NCT00514540 clinical trial. One patient was ineligible for treatment on trial due to thrombocytopenia but otherwise met AVPC criteria thus making 115 patients evaluable for the current study. We identified 70 surgically excised and biopsy specimens from 47 of the 115 evaluable patients. Of these, 59 (from 40 patients) had sufficient tumor to attempt a full immunohistochemical (IHC) panel (Table 1).

PDXs
We also used 8 PDX lines derived from 6 of the NCT00514540 trial participants (Supplementary Tables S2 and S3). MDA prostate cancer (PCa) 155-2, 144-4, and 144-13 have been reported previously (10). The latter two were derived from the same donor tumor. MDA PCa 177-0 and MDA PCa 189-1 were derived from the prostate of patient #83 – 0.1 months before and 11.9 months after chemotherapy, respectively. Two (MDA PCa 163-A and MDA PCa 177-0) were obtained before chemotherapy and MDA PCa 163-A was obtained while patient had noncastrate levels of testosterone. The rest were obtained >5 months after the start of chemotherapy (timeline shown in Fig. 4). The donor tumors for MDA PCa 177-0, 189-1, and 166-1 were not available for study.

DNA extraction-Oncoscan
Tumor areas within samples were marked by the pathologist on a hematoxylin–eosin (H&E)-stained slide to guide the acquisition of a 0.6 mm core biopsy from the FFPE block. After xylene extraction and ethanol wash, DNA was isolated using the Recover All Total Nucleic Acid Isolation kit for FFPE (Life Technologies, catalog-AM1975). The average DNA recovered was 36.6 ng/mL (range, 2.6–203.3 ng/mL). All samples (de-identified) were placed on sealed 96-well plates provided by Affymetrix (as per Affymetrix Services Guide for Oncoscan FFPE Assay Kit) and shipped on dry ice.

IHC, quantitative real-time reverse transcriptase-PCR and Western blot analyses are described in Supplementary Material. Supplementary Tables S4 and S5 list the antibodies and primer sequences used.

Statistical analyses
Supervised hierarchical clustering of gene expression profiles. Gene expression of 6 PDX profiles using the U133A2.0 Plus array (Affymetrix, Inc.) was previously described. (11) Here, genes within categories of interest (luminal-epithelial-androgen, neural development, and cell-cycle-proliferation; Supplementary Fig. S1) were selected from RMA-processed and log2-transformed gene expression data to cluster samples with use of Euclidean distance and complete linkage.

Analyses of patient characteristics and IHC studies. Patient characteristics were tabulated by donor status. Fisher exact tests and Wilcoxon rank sum tests were used to test differences between covariates. Uncertain values of serum markers were replaced by their upper limits (for example, <0.1 was replaced by 0.1). The standardized ChrA was redefined as (ChrA-36.4)/36.4 or (ChrA-s-225)/225. Relationships among the markers in donor patients were evaluated using Pearson correlations. A corrgram was applied for exploratory visual display (14).

The effect of the markers on variance was investigated using Principal component analysis (PCA) and component scores were grouped as ≥0 vs. <0. The association between component score groups and SCPC, overall survival (OS), or progression-free survival (PFS) were investigated using Fisher test or log-rank tests, respectively. Cox proportional hazards regression models were fit to assess the association between AVPC markers and OS or PFS (15). Predictive factors were transformed as appropriate based on martingale residual plots (16). Statistical analyses were performed with SAS 9.3 (17) and S-Plus (18) software.

Bioinformatics analyses
Reports and detailed descriptions of the bioinformatics analyses performed in this study are available on the UT MDACC
Department of Bioinformatics and Computational Biology
website http://bioinformatics.mdanderson.org/main/Publications_Supplements_Data. Data for 44 (36 prostate cancer patient samples; 8 PDX samples) plus 2 Affymetrix control samples were submitted producing 92 CEL files (two arrays per sample: AT and GC channels) and have been deposited in the Gene Expression Omnibus (GEO) under accession number GSE73902. Raw data were processed in Affymetrix OncoScan Console to yield log R ratios (LRR) and B allele frequencies (BAF) for each single nucleotide polymorphism (SNP) in each sample by using the TuScan algorithm. Computations were performed relative to the normal reference provided by Affymetrix. LRR and BAF plots were generated along each chromosome for some sample with use of ASCAT 2.1 (allele-specific copy number analysis of tumors, version 2.1). Two quality control (QC) metrics provided by the software, the MAPD (median of the absolute values of all pairwise differences) and the ndSNP/QC (SNP QC of Normal Diploid Markers) were within bounds in 23 (52.3%) and 18 (40.9%) of the 44 samples, respectively. Given that these QC markers were developed for normal diploid genomes and that heterogeneouscopy number changes, we used LRR values relative to (a) absolute cutoffs (marker based changes, we used LRR values relative to (a) absolute cutoffs and (b) the median segmentation of the corresponding chromosome arm (p or q). For regions with multiple segments, the extreme value away from zero was used to make the copy number call. To determine how gene-specific events related to regional/chromosomal integrity, we plotted the LRR values for each gene of interest versus the LRR values for the corresponding chromosomal arm. To study the correlation between copy number (median of segmentation) and labeling indices, we performed both Pearson and Spearman analyses. The Broad Institute GISTIC 2.0 tool was used for clustering of the median LRR values for each chromosome arm. We used both unsupervised (PCA) and supervised display methods to determine the degree to which the AVPC samples stood out from the unselected TCGA prostate cancer population. These analyses used level 3 copy number estimates from Affymetrix SNP6 arrays (n = 404) and somatic mutation calls for Tp53 retrieved from cBioPortal (n = 253 of the 404; refs. 19, 20). To define our final classification rule, we used logistic regression with 10 potential exploratory covariates and used top-forward regression coupled with cross-validation to arrive at a solution with high sensitivity. Copy number alterations and Tp53 mutations within the data reported by Robin-son and colleagues (21) were queried from cBio.

Results
Interactions between tumor suppressors, AR-signaling, neurodevelopmental and mitotic immunohistochemistry, and serum markers in the AVPC
In our previously reported PDX gene expression profile (11), we found that SCPCs segregated from typical CRPC adenocarcinomas based on RB loss and expression of luminal epithelial markers, genes implicated in mitosis and pronuclear transcription factors (Supplementary Fig. S1). We also found (as others) frequent Tp53 alterations in the SCPCs (10, 13). To examine whether the clinically defined AVPC shared molecular features with SCPC, we used IHC analysis to query the expression of RB1, Tp53, AR, NKX3.1, ASC1, BRN2, AURKA, UBE2C, and Ki67 in the 59 tumor samples and 8 PDX lines obtained from 42 participants in clinical trial NCT00514540 for men with mCRPC meeting AVPC criteria (Supplementary Table S1; ref. 9). Of the 59 samples (Table 1), 5 from 5 patients (early) were obtained >13 months prior to registration (median 24.1; range -131.6 to -14.0); 38 from 31 patients (baseline) between -13 and +1 months from registration (median -0.18; range -12.18 to +1); and 16 from 13 patients (postchemotherapy) >1 month after registration and chemotherapy (median 7.43; range 4.04 to 28.46). Supplementary Tables S6 and S7 summarize the patient donor characteristics and serum marker levels at registration. Supplementary Figure S2 shows their morphologic spectrum. The 31 patient donors of the "Baseline samples" were similar to "nondonor" participants of the NCI00514540 trial except for having worse Eastern Cooperative Oncology Group (ECOG) Performance Status (PS) scores (P < 0.001) and higher lactate dehydrogenase (P < 0.001) levels. Supplementary Tables S2 and S3 summarize the clinicopathologic features of the donor patients and tumors for the PDX, their growth rates and their PSA production.

Shown in Fig. 1A are the percent of cells with positive stains for each marker in the "Baseline samples." Where repeated measurements were available for a single patient, only the measurement taken closest to the registration date was considered (n = 31). Of these 80.6% (25/31) displayed Ki67 labeling indices ≥10%, as did 71.0% (22/31) for nuclear (N) AURKA (AURKA-N). 51.6% (16/31) for cytoplasmic (-C) AURKA (AURKA-C), 35.5% (11/31) for both AURKA-N and AURKA-C, 44.8% (13/29, 2 uninterpretable) for UBE2C-N, 48.3% (14/29) for UBE2C-C, 37.9% (11/29) for both UBE2C-N and UBE2C-C, and 32.2% (10/31) for Tp53. In 62.1% (18/29, 2 uninterpretable) and 41.9% (13/31) RB1 and AR stained <10% nuclei, respectively. Results for the remaining samples are in Supplementary Fig. S3.

RB1-negative samples appeared more likely to be AR-negative, to be Tp53-positive, and to have higher Ki67 and UBE2C staining. Indeed, Pearson correlations of the 10 tissue markers of the "Baseline samples" and the levels of 6 serum markers at registration (PSA, PAP, CEA, chromogranin A, calcitonin, and somatostatin; Fig. 1B; Table 2) showed strong positive correlations between AR and NKX3.1, RB1, AURKA-N, and serum PSA. Strong positive correlations were also observed between NKX3.1 and RB1; Tp53 and UBE2C-N and Ki67; UBE2C-N and Ki67 and serum somatostatin; UBE2C-C and Ki67; and PSA and calcitonin. A strong negative correlation was observed between NKX3.1 and Ki67.

To determine which marker(s) best explained the variance in the data and examine their relationship with outcome, PCA was applied to the 16 serum and IHC tumor marker set of the "Baseline" samples from 21 patients (10/31 were excluded due to missing values). The first principal component had high positive correlations with PSA, AR-N, NKX3.1-N, RB1, and AURKA-N, and negative correlations with Tp53, UBE2C-N, Ki67, and somatostatin (Supplementary Fig. S4A). A strong association between small cell morphology (n = 6 samples) and first component score ≥0 versus <0 was observed (P = 0.01), but clustering of the samples was not (Supplementary Fig. S4B). Neither overall survival (OS) nor progression-free survival (PFS) were
Table 1.

| Patient number | Sample Name | Early Samples (n=5) | Late Samples (n=22) | SCPC (n=18) | IHC & Serum Pearson (n=31) | IHC & Serum PCA (n=21) | CNA Cluster (n=29) | Sample names indicate patient or PDX line number and the procedure and site of origin. Samples are ordered by patient. Early samples were obtained more than 13 months prior to registration to NCT00514540. Baseline values for the median of the chromosome arm 10q, suggesting 21 samples showed CNL for PTEN and copy number for each marker. Using the relative cut-off values decreased to 2 (1 loss and 1 gain) and for MYCN amplification (12). To expand on the IHC findings and further investigate the potential heterogeneity within the AVPCs, we submitted DNA from 44 tumor samples (36 patient samples and 8 PDXs) for Oncoscan analysis. Assuming 60% tumor purity and using absolute cut-off values, a total of 197,971 copy number alteration (CNA) calls (58,845 losses and 139,126 gains) were made across all samples: 22,279 of 23,355 genes (99.7%) displayed ≥1 CNA in ≥1 sample. However, tumor samples varied widely with respect to CNA: between 29 and 17,249 (median 3,650.5) CNA calls made per sample (Supplementary Fig. S5, full list at http://bioinformatics.mdanderson.org/main/Publications_Supplements_Data). The gene showing the highest CNA rate across the AVPC tumors was ADCY8 (8q24.22), amplified in 27 of 44 samples (61.4%). MYC (8q24.21) was amplified in 24 (54.5%). The gene that showed the highest rate of copy number loss (CNL) was RB1 (13q14.2), lost in 24 samples (54.5%), followed by PTEN (10q23.31), lost in 21 (47.7%). Also using absolute cut-off values, copy number gains (CNG) were called for AURKA (20q13.31) in 11 samples (25.0%), MYCN (2p24.3), UBE2C (2q11.12) and AR (Xq12) in 9 (20.4%) each, and ASC1L1 (12q23.32) in 6 (13.6%). CNG for NXX3-1 (8p21.2) were called in 14 samples (31.8%) and for TP53 (17p13.1) in 11 (25.0%). Seven samples (15.9%) from 5 patients displayed concurrent AURKA and MYCN CNG, 4 of which (from 2 patients) showed SCPC morphology. Because a CNA that involves a specific gene segment (as opposed to a region containing multiple genes) reinforces the functional significance of the gene we used cut-off values relative to the corresponding chromosome arm to ask whether the CNA involving each of the genes of interest was a "segmental" versus a "regional" alteration. As shown in Fig. 2A, 19/21 samples showed CNI for PTEN despite copy number neutral values for the median of the chromosome arm 10q, suggesting a gene-specific alteration. This was also true for RB1 in 11 of the 24 samples with CNI and for MYCN in 4 of 9 samples with CNG. However, for TP53 and AURKA the number of CNA calls using the relative cut-off values decreased to 2 (1 loss and 1 gain for both) and for MYC to 0. We examined the correlation between the IHC labeling indices and copy number for each marker. RB1 CNA showed a positive correlation with its labeling index (Fig. 2B) but a number of copy number neutral tumors showed ≤10% RB1-positive cells, indicating CNI is not the sole mechanism of RB silencing in these significantly associated with the dichotomized first component scores within this sample set (data not shown). These data suggest that, irrespective of morphology, AVPCs express high levels of mitotic markers, are frequently RB1 negative and that RB1 loss within them is often associated with mutant TP53, loss of AR, and high Ki67. The observed heterogeneity may be due to limitations in the specificity of the clinical criteria used for patient selection, to the shortcomings of IHC analysis of a single marker to evaluate a full pathway, and/or to actual biologic heterogeneity within AVPC.

Copy number alterations in AVPCs

Absence of a full-length RB1 protein in SCPC is commonly a consequence of genomic deletions or microdeletions in RB1 (11, 13). The mechanisms of TP53 alteration are more diverse and include gene mutations or deletions (13). Also, AURKA amplification has been reported in 40% of SCPC, with concurrent MYCN amplification (12). To expand on the IHC findings and further investigate the potential heterogeneity within the AVPCs, we submitted DNA from 44 tumor samples (36 patient samples and 8 PDXs) for Oncoscan analysis. Assuming 60% tumor purity and using absolute cut-off values, a total of 197,971 copy number alteration (CNA) calls (58,845 losses and 139,126 gains) were made across all samples: 22,279 of 23,355 genes (99.7%) displayed ≥1 CNA in ≥1 sample. However, tumor samples varied widely with respect to CNA: between 29 and 17,249 (median 3,650.5) CNA calls made per sample (Supplementary Fig. S5, full list at http://bioinformatics.mdanderson.org/main/Publications_Supplements_Data).

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tumors. For the remaining markers, there was no correlation between CNA and their labeling indices (data not shown).

These data indicate that the most common CNA across the AVPC samples are 8q amplification and RB1 and PTEN CNL (which occurred irrespective of morphology, i.e., in both small-cell and non–small cell histologies alike), and also that CNA are insufficient to detect full pathway alterations in these tumor samples (12, 22).

Pathway analysis in PDX
To gain insight into the significance and the functional consequences of the alterations observed by IHC and copy number analysis, we examined associated and downstream molecular events in the PDX models.

Loss of RB1 RNA and protein expression was a frequent finding, consistent with the CNL observations (Fig. 3). Two of the 3 RB-positive PDX showed high levels of phospho-RB by Western blot analysis, an inactive RB state (23). We interrogated levels of representative genes from published RB-loss signatures (24), by qRT-PCR (Supplementary Fig. S6A). In all but one (MDA PCa 163A) of the RB-negative and hyperphosphorylated RB PDX, this partial RB-loss signature was expressed at intermediate to high levels relative to the RB-positive PDX MDA PCa 189-1.

We found Tp53 mutations in 7 of the 8 PDX lines, 6 of them within the DNA-binding domain (Fig. 3). In MDA PCa 163-A, a truncating mutation (R65/C3) was associated with loss of expression, exemplifying the fact that absent Tp53 staining does not signify an intact gene and suggesting that Tp53 alterations are more common in the AVPC samples than was evident by IHC analysis. At least two of the Tp53 mutants found (R273C and Y163N) have been associated with gain-of-function activity (25, 26). Whether distinct Tp53 mutants are linked to different clinical phenotypes remains to be determined.

**Figure 1.**
IHC analysis of NCT00514540 unique “Baseline” samples. A, labeling indices (% of cells staining positive for the marker listed on the y-axis, Li) for RB1, Tp53, AR, NKX3-1, AURKA, UBE2C, and Ki67 in unique patient and PDX baseline samples (only one sample per patient is shown of the set obtained between −13 and +1 months from registration). Note, 177-0-XENO is included in the graph (n = 32) but not in the analyses described in the text because the donor-patient tumor was not available for analysis. The BRN2 stains did not work and are not reported. Black bars depict nuclear staining. For AR, AURKA, and UBE2C cytoplasmic stains are shown in light gray. Each individual sample is listed on the x-axis. Numbers indicate accession number on the protocol or PDX line. Letters indicate site of origin. Blue arrowheads indicate SCPC morphology. B, Corrgram of selected tissue and serum markers among the 31 baseline samples’ donors. Red represents positive values and blue represents negative values. The intensity of the color is proportional to the magnitude of the correlation. Asterisk (*) indicates the IHC did not work. Abbreviations: RB, retinoblastoma 1; p53, tumor suppressor Tp53; NKX3-1, NK3 homeobox 1; AURKA, aurora kinase A; UBE2C, ubiquitin conjugating enzyme 2C; Ki67, antigen identified by monoclonal antibody Ki-67; RXPX, radical prostatectomy; BMBX, bone marrow biopsy; TURP, transurethral resection of prostate; CRAN, craniotomy; RCBX, rectal biopsy; XENO, xenograft; PVBX, pelvic mass biopsy; LNBX, lymph node biopsy; BOBX, bone biopsy; LIBX, liver biopsy; PAP, prostatic acid phosphatase; PSA, prostate-specific antigen; -N, nuclear staining; -C, cytoplasmic staining; StdCHRA, standardized chromogranin A; CEA, carcinoembryonic antigen.

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Table 2. Correlations between selected measurements of tissue and serum markers among "baseline samples" (n = 31)

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NOTE: Strong correlations (≥0.5) are highlighted in black. Abbreviations are described in the Fig. 1B legend.
promiscuity that is re-functionalized and account for the increased migratory and invasive properties of CRPC. Their expression patterns also suggest that, as in neural development, different mental pathways are coopted in the latter stages of tumor evolution. They also suggest that, as in neural development, different neurodevelopmental pathways are coopted in the latter stages of tumor evolution.

Integration of IHC, CNA, and PDX data and refinement of marker selection with reference to publicly available datasets

With the overall goal of arriving at a candidate molecular signature to distinguish the AVPC and guide therapy allocation in men with CRPC, we combined the IHC, CNA, and PDX data and used The Cancer Genome Atlas (TCGA) and a recently published unselected CRPC dataset (21) to refine our marker selection.

We examined the rate of CNA in the genes of the markers initially queried by IHC (RBI, TP53, AR, NKX3-1, ASCL1, AURKA, and UBE2C) in the 404 samples from TCGA analyzed by Affymetrix SNP 6.0 Array and the 150 unselected CRPC

As in the patient samples, AR expression was heterogeneous in the PDX and its presence tightly associated with canonical downstream signaling markers (such as PSA) and markers of luminal epithelial differentiation (such as HOXB13 and the pioneer factor GATA2; Supplementary Fig. S6B). NKX3-1 and FOXA1 were expressed in all AR-positive tumors but also in some of the AR-negative tumors. These data suggest that, when AR is expressed it remains active and within the context of a prostate luminal epithelial program. AR splice variants were also expressed in the AR-positive tumors but at much lower levels than the full-length AR.

An inverse correlation between the expression of AR and that of the proneural transcription factors (TF) tested (ASCL1, BRN2, NEUROD1, INSM1, MYCN) was observed (Fig. 3 and Supplementary Fig. S6C). However, there was no single proneural TF that showed increased expression across all models. Even within PDX lines derived from the same tumor, MDA PCa 144-13 and MDA PCa 144-4 (10), the expression of proneural TF was heterogeneous. REST was absent in the PDX lines with small cell carcinoma morphology, which had the highest levels of proneural TF. However, in MDA PCa 177-0, BRN2 and MYCN were expressed despite the presence of REST (Supplementary Fig. S6C). These findings are consistent with the hypothesis that neural developmental pathways are coopted in the latter stages of tumor evolution. They also suggest that, as in neural development, different combinations of proneural TF result in distinct corresponding functions and account for the increased migratory and invasive promiscuity that is reflected in the unconstrained metastatic patterns of AVPC.

Levels of AURKA mRNA were concordant with those of UBE2C while levels of AURKB mRNA for the most part paralleled those of PLK1 and were higher in the AR-negative PDX (Supplementary Fig. S6D). Protein levels of all three mitotic kinases and UBE2C were lowest in MDA PCa 205-6 and MDA PCa 189-2. MDA PCa 205-6 was derived from sample 118-RXPX (Supplementary Fig. S3), which was donated by the patient with the longest survival (Supplementary Table S2) of the PDX donors. It was also the only PDX in which a TP53 mutation was not found. MDA PCa 189-1 was derived, after chemotherapy, from the same patient as MDA PCa 177-0. Both PDX retained the Y163N TP53 mutation but while the prechemotherapy MDA PCa 177-0 PDX was RB-negative/AR-negative/proneural TF-positive high, the postchemotherapy MDA PCa 177-0 PDX was RB-negative/AR-negative/proneural TF-positive high, the postchemotherapy MDA PCa 189-1 was RB-positive/AR-positive/proneural TF-negative and expressed lower levels of TP53, suggesting either clonal selection or tumor plasticity as a result of treatment. Protein and mRNA levels of the mitotic kinases and UBE2C were not correlated, corroborating the importance of regulatory posttranslational events for these markers.

Figure 2.

Copy number alterations at genes of interest and correlation with IHC results. A, segmental (values corresponding only to the gene of interest) versus regional (median values for the corresponding chromosome arm) copy number calls for PTEN, RBI, MYCN, AURKA, Tp53, and MYC. “n” in parenthesis following the gene name indicates the number of gene-specific probes present on the Oncoscan chip. The blue and red dashed lines indicate the absolute cutoff values. The solid blue line represents the corresponding chromosome arm copy number. Red dots indicate samples with SCPC morphology. B, correlation between labeling indices (y-axis) and copy number (x-axis) for RBI.

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Copy number alterations at genes of interest and correlation with IHC results. A, segmental (values corresponding only to the gene of interest) versus regional (median values for the corresponding chromosome arm) copy number calls for PTEN, RBI, MYCN, AURKA, Tp53, and MYC. “n” in parenthesis following the gene name indicates the number of gene-specific probes present on the Oncoscan chip. The blue and red dashed lines indicate the absolute cutoff values. The solid blue line represents the corresponding chromosome arm copy number. Red dots indicate samples with SCPC morphology. B, correlation between labeling indices (y-axis) and copy number (x-axis) for RBI.
samples recently reported by Robinson and colleagues (21). We also queried PTEN and MYC (the latter taken as a surrogate for 8q amplification), given their frequent alteration in the AVPC samples, as well as MYCN, given its reported frequent amplification in SCPC (12), and Repressor element (RE)-1 silencing transcription factor (REST), given its role as a master transcriptional regulator of neural development and its reported down-regulation in SCPC (ref. 22; Fig. 4A). An unsupervised PCA (Supplementary Fig. S7) showed that of the 10-CNA-marker(s) (RB1, PTEN, Tp53, MYC, MYCN, NKX3-1, ASCL1, REST, AURKA, and UBE2C) MYC and NKX3-1 best explained the variance between the three sample sets (Supplementary Table S8). As our primary objective is to identify the AVPC within an unselected CRPC population, we applied a supervised linear discriminant analysis to examine the segregating ability of the 10-CNA marker (Fig. 4B). In this analysis, 38 (25.3%) of the 150 unselected CRPC samples segregated with the AVPC samples (an expected proportion based on our prior experience) and RB1 CNL carried the largest weight in this distinction (Supplementary Table S9).

Hence alterations in RB1 (both CNL and/or negative staining by IHC), Tp53 (CNL, positive staining by IHC and/or detectable mutation), PTEN CNL, and MYC CNG appeared to be the most common molecular alterations of the AVPC, with MYC CNG seemingly more frequent in the unselected CRPC than in the AVPC. These are prevalent alterations in CRPC (21, 27), and have been described at various rates in primary prostate tumors (27), but genetically engineered mouse models (GEMM) have shown that each of these alterations individually is insufficient to result in lethal prostate cancer (28–32). In contrast, the concurrent loss of RB1 and Tp53 resulted in highly metastatic neoplasms of various morphologies that are resistant to androgen depletion from early stages of development both in Tag, TRAMP, and PB-Cre4 (PB-Cre4, p53loxP/loxP, RbloxP/loxP) driven models (33, 34) and haploinsufficiency of the Pten gene increased the rates of tumor progression in the TRAMP mouse model (35).

We thus asked how often the alterations in RB1, Tp53, and PTEN occurred together in TCGA, the AVPC and the unselected CRPC samples (21) using the available information. Here we included Tp53 mutations identified by sequencing in the unselected CRPC and TCGA samples (70 (46.7%) of 150 and 24 (9.5%) of 253, respectively (Supplementary Table S10). In the AVPC samples, Tp53 sequencing was only available for the PDX and high confidence calls using the Oncoscan data could only be made for 3 of the 29 samples (therefore, here the rate of Tp53 mutations in the AVPC is underestimated). Alterations in 2 of the 3 markers were found in 25 (9.9%) of the 253 TCGA samples for which Tp53 mutation status was available, in 39 (26.0%) of the 150 unselected CRPC and in 14 (48.3%) of the 29 AVPC samples.

Taken together, our observations in patient tumors and PDX models suggest that, irrespective of morphology, clinically defined AVPC are characterized by joint alterations in two or more of the following: RB1, Tp53, and PTEN.
The findings we report link the prospectively clinically defined AVPC and their corresponding PDX to a molecular marker profile not predicted by morphology and suggest that the aberrancy of multiple tumor suppressors (RB1, Tp53, and/or PTEN) is a dominant “driver” leading to the development of this phenotype. Whether different combinations of the alterations are linked to distinct clinical behaviors within the AVPC remains to be determined. Adding significance to the findings is the similarity of the molecular alterations to those observed in SCPC. Furthermore, these alterations have been linked to increased DNA damage response signaling (36) which, in a parallel study, we have identified in the AVPC (26), and may represent an actionable therapy target for this entity. The results from our dataset and those of others (12, 13, 22), will lead to the identification of predictive markers that enable the development and use of effective therapy combinations and sequences in the biologically distinct subsets of men with metastatic CRPC.

It has been proposed that AURKA amplification and overexpression are implicated in the pathogenesis of SCPC (12). We observed a high frequency of increased nuclear and/or cytoplasmic AURKA expression as well as 20q amplification in 7 (24.1%) of 29 unique AVPC samples irrespective of morphology. Intriguingly, in the AVPC, AURKA nuclear expression correlated with AR expression. Indeed, the interactome of AURKA is large (37) and includes cell cycle and transcriptional regulators such as AR (38), Tp53 (39), MYC (40), and BRCA1 (41). In turn, Tp53 mutants increase AURKA expression and cooperate with AURKA in the induction of genomic instability (ref. 42; also a likely feature of the AVPC suggested by the high rate of CNA observed) and in the regulation of DNA damage repair (43). Although these data are congruent with a role for AURKA in the pathogenesis of the AVPC, two concerns remain. First, amplification of AURKA is most often regional and thus accompanied by the enhanced expression of multiple additional genes, making the attribution of outcomes to AURKA overexpression alone difficult (37). Second, there is a lack of a standard IHC methodology and of a consensus as to what constitutes abnormal AURKA expression (37, 44). Thus, additional investigation is needed to determine the significance of AURKA expression in the AVPC and its value as a marker of this phenotype.

The loss of AR in a large proportion of the AVPC is consistent with the resistance to AR signaling inhibition. Of interest is...
the finding that AR expression is inversely correlated with that of neurodevelopmental pathway markers. If confirmed, this observation would be in contrast to the frequent coexpression of AR with markers of neuroendocrine differentiation such as chromogranin A (43). The presence of neuroendocrine differentiation is often considered a precursor or even synonymous with SCPC. However, neuroendocrine marker expression in prostate tumors is associated with a wide range of clinical behaviors (46) and is not predictive of chemotherapy responsiveness (47). A master regulator of the neural-progenitor phenotype or a consistent signature that reflects it remains to be fully characterized. Loss of REST has been proposed (22), although in our PDX models, the expression of proneural TF was not consistently linked to loss of REST mRNA. Experiments are ongoing to determine whether these sequences correspond to the known dominant-negative REST splice variant (48). Additional studies in larger numbers of patients will be needed to determine whether adoption of a neural-progenitor program is but one path of progression to the clinical phenotype. Ongoing studies will determine the prevalence of APVC, prospectively evaluate the utility of the proposed marker signature, characterize the potential heterogeneity within them, and test specific therapies tailored to their unique biology.

The data we report demonstrate that some castration-resistant prostate adenocarcinomas share molecular as well as clinical features with SCPC. The consistent marker profile of the morphologically heterogeneous cancers in this dataset comprises combined alterations in RB1, Tp53, and/or PTEN. This marker profile can account for the observed cell-cycle alterations, AR signaling loss, and neural progenitor program adoption, which in turn can explain the clinical phenotype. Ongoing studies will determine the presence of APVC, prospectively evaluate the utility of the proposed marker signature, characterize the potential heterogeneity within them, and test specific therapies tailored to their unique biology.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Authors’ Contributions

Conception and design: A.M. Aparicio, S.N. Maity, C.J. Logothetis
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): A.M. Aparicio, E.L.N. Tapia, G. Wu, P. Troncoso, P. Corn, S.N. Maity
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): A.M. Aparicio, L. Shen, H.-C. Chen, J. Zhang, X. Wang, P. Troncoso, P. Corn, B. Broom, K. Baggerly, S.N. Maity
Writing, review, and/or revision of the manuscript: A.M. Aparicio, L. Shen, E.L.N. Tapia, H.-C. Chen, J. Zhang, P. Troncoso, P. Corn, T.C. Thompson, B. Broom, S.N. Maity, C. Logothetis
Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): J.-F. Lu, C. Logothetis, A.M. Aparicio
Study supervision: A.M. Aparicio, S.N. Maity, C.J. Logothetis

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References

## Combined Tumor Suppressor Defects Characterize Clinically Defined Aggressive Variant Prostate Cancers

Ana M. Aparicio, Li Shen, Elsa Li Ning Tapia, et al.

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