NF-κB2/p52:c-Myc:hnRNPA1 Pathway Regulates Expression of Androgen Receptor Splice Variants and Enzalutamide Sensitivity in Prostate Cancer

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Abstract

Castration-resistant prostate cancer (CRPC) remains dependent on androgen receptor (AR) signaling. Alternative splicing of the AR to generate constitutively active, ligand-independent variants is one of the principal mechanisms that promote the development of resistance to next-generation antiandrogens such as enzalutamide. Here, we demonstrate that the splicing factor heterogeneous nuclear RNA-binding protein A1 (hnRNPA1) plays a pivotal role in the generation of AR splice variants such as AR-V7. hnRNPA1 is overexpressed in prostate tumors compared with benign prostates, and its expression is regulated by NF-κB2/p52 and c-Myc. CRPC cells resistant to enzalutamide exhibit higher levels of NF-κB2/p52, c-Myc, hnRNPA1, and AR-V7. Levels of hnRNPA1 and AR-V7 are positively correlated with each other in prostate cancer. The regulatory circuit involving NF-κB2/p52, c-Myc, and hnRNPA1 plays a central role in the generation of AR splice variants. Downregulation of hnRNPA1 and consequently of AR-V7 resensitizes enzalutamide-resistant cells to enzalutamide, indicating that enhanced expression of hnRNPA1 may confer resistance to AR-targeted therapies by promoting the generation of splice variants. These findings may provide a rationale for cotargeting these pathways to achieve better efficacy through AR blockade.

Introduction

Prostate cancer remains the second most lethal disease for males in western countries. The development of abiraterone and enzalutamide marked the continuing success of androgen deprivation therapy (ADT) practiced for more than 70 years, reinforcing the concept that androgen receptor (AR) is the key factor for metastatic castration-resistant prostate cancer (CRPC) progression and lethality. However, like earlier ADT, these new therapies have a short efficacy due to primary or acquired resistance. A major form of ADT-resistance in prostate cancer is the generation of AR splicing variants that lack the ligand-binding domain, thus evading binding of antiandrogens such as bicalutamide and enzalutamide. Several reports have documented the expression of alternatively spliced AR-Vs lacking the C-terminal ligand binding domain in prostate cancer cells, which are constitutively nuclear and active even in the absence of androgens, thus indicating their potential role in the acquisition of the CRPC phenotype. Expression of these variants arises from the inclusion of cryptic exons located in intron 2 and 3 of the AR gene, which inserts premature stop codons and termination sites, yielding shorter AR proteins of 75 to 80 kDa lacking the androgen-binding domain (1, 2). Truncated AR-Vs such as AR-V7 (AR3) and ARV6/7 can function independently of full-length AR and their selective knockdown can suppress androgen-independent growth of CRPC cells. Alternatively, AR-Vs may play important roles in activating the full length AR in a ligand-independent manner (3). AR-Vs confer resistance to not only AR-targeted therapies (4, 5) but to conventional chemotherapy such as taxanes used as first line therapies against CRPC (6). These splice variants are rapidly induced after androgen deprivation and are suppressed after restoration of androgen supply (7). The mechanisms mediating increased expression of aberrant AR-Vs in prostate cancer are still largely unknown. One possible cause of defective splicing is the genomic rearrangement and/or intragenic deletions of the AR locus in CRPC (8). Alternatively, aberrant expression of specific splicing factors in prostate cancer cells may also contribute to unbalanced splicing and aberrant recognition of cryptic exons in the AR gene. Understanding the molecular mechanism of AR-Vs production will facilitate the design of mechanism-based inhibitors, extending the efficacy of current ADT, and possibly treating progression of CRPC and prolonging patient survival. The importance of alternative messenger RNA splicing in regulatory circuits is underscored by the fact that >90% of human genes encode transcripts that undergo at least one alternative splicing event with a frequency higher than 10% (9, 10). Alternative splicing plays important roles in development, physiology, and disease and is often disturbed in inflammatory disorders and cancers (11, 12). Alternative splicing modulates the generation of protein isoforms with distinct structural and functional properties.

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or affects mRNA stability, by the insertion of premature stop codons, and translatability, by altering microRNA target sites (13). Two nuclear RNA-binding protein families, heterogeneous nuclear ribonucleoproteins (hnRNP) and serine/arginine-rich proteins (SR), play pivotal roles in regulation of alternative splicing. The hnRNP family consists of ~20 members which bind to splicing silencers located in exons or introns to promote exon exclusion and act as splicing repressors (13). The best characterized proteins of this group are hnRNP A1 and hnRNP A2, which share a high degree of sequence and functional homology (14). HnRNP A1 and hnRNP A2 are overexpressed in various kinds of tumors and serve as early tumor biomarkers (15–17). The SR family includes >20 members, which bind to splicing enhancers and predominantly function to counterbalance the activity of hnRNP proteins (18). Splicing factor 2/alternative splicing factor (SF2/ASF), the best-characterized member of the SR family, is upregulated in multiple human cancers, including lung and cervical cancers, and plays important roles in the establishment and maintenance of cellular transformation (19). During tumor progression, stimuli from the tumor microenvironment may affect the expression and/or activity of splicing regulatory factors, thus perturbing the physiological splicing program of genes involved in cellular processes. An increasing body of evidence indicates that splicing variants of many cancer-related genes can directly contribute to the oncogenic phenotype and to the acquisition of resistance to therapeutic treatments (11, 12, 20). Hence, understanding the functional role(s) of cancer-associated alternative splicing variants and the mechanisms underlying their production offers the potential to develop novel diagnostic, prognostic, and more specific anticancer therapies.

In this study, we investigated the mechanisms involved in aberrant splicing of AR transcripts in a constitutively occurring setting as well as in response to chronic treatment with enzalutamide. Our results show that the splicing factor hnRNP A1 plays a major role in generation of AR-Vs. We also demonstrate that enhanced expression of hnRNP A1 may be mediated by c-Myc and NF-kB2/p52, thus paving the way for increase in transcript numbers of constitutively active splice variants and contributing to CRPC therapy resistance.

**Materials and Methods**

**Cell lines and reagents**

LNCaP, CWR22Rv1, and VCaP cells were obtained in 2001 from the American Type Culture Collection (ATCC) and were cultured in RPMI containing 10% complete FBS and penicillin/streptomycin. ATCC uses short tandem repeat (STR) profiling for testing and authentication of cell lines. C4-2B cells were kindly provided and authenticated by Dr. Leland Chung, Cedars-Sinai Medical Center, Los Angeles, CA, USA, in 2006. All experiments with these cell lines were performed within 6 months of resuscitation after cryopreservation. LNCaP cells stably expressing NF-kB2/p52 were generated by stable transfection of LNCaP cells with plasmids expressing NF-kB2/p52 as described previously (21) and were not authenticated further. 22Rv1 and C4-2B cells resistant to enzalutamide (22Rv1-Enza-R and C4-2B-Enza-R) were generated by chronic culture of 22Rv1 and C4-2B cells in enzalutamide as described previously (22, 23) and were not authenticated further. Antibodies against NF-kB2/p52 (K-27), AR (441; mouse monoclonal), HA, tubulin, U2AF65, and ASF/SF2 were from Santa Cruz Biotechnologies. Antibodies against splicing factors hnRNP A1 (9H10) and hnRNP A2B1 (DP3B3) were from Sigma-Aldrich and AbCam, respectively. Sso Fast Eva Green qPCR Supermix was from Bio-Rad. All other reagents were of analytical grade and obtained from local suppliers.

**Cell growth assays**

Plasmid transfections were performed using Attractene transfection reagent (Qiagen). Oligonucleotide siRNA transfections were performed using Lipofectamine 2000 transfection reagent (Invitrogen). Viable cell numbers were determined using a Coulter cell counter (Beckman Coulter).

**Western blot analysis**

Cells were lysed in high salt buffer containing 50 mmol/L Hepes pH 7.9, 250 mmol/L NaCl, 1 mmol/L EDTA, 1% NP-40, 1 mmol/L PMSF, 1 mmol/L Na Vanadate, 1 mmol/L NaF, and protease inhibitor cocktail (Roche). Total protein was estimated using the Coomassie Protein Assay Reagent (Pierce). Equal amounts of protein were loaded on 10% SDS-PAGE and transferred to nitrocellulose membranes. The membranes were blocked with 5% nonfat milk in PBST (1 × PBS + 0.1% Tween-20) and probed with indicated primary antibodies in 1% BSA. The signal was detected by ECL (Millipore) after incubation with the appropriate horseradish peroxidase-conjugated secondary antibodies.

**Real-time quantitative RT-PCR**

Total RNAs were extracted using TRIzol reagent (Invitrogen). One microgram of total RNAs were subjected to digestion with RNase-free RQ1 DNase (Promega) and cDNAs were prepared using ImPromIII Reverse Transcriptase (Promega) according to manufacturer’s instructions. The cDNAs were analyzed by real-time reverse transcription-PCR (RT-PCR) using Sso Fast Eva Green Supermix (Bio-Rad) as described previously (24). Each reaction was normalized by coamplification of actin. Triplicates of samples were run on a Bio-Rad CFX-96 real-time cycler.

**RNA immunoprecipitation assay (RIP)**

RIP assays were performed as described earlier (25). RNA-protein complexes were cross-linked using 1% formaldehyde. Nuclear extracts were immunoprecipitated with antibodies against indicated RNA-binding splicing factors. Isotype-matched IgG was used as control. Bound RNAs were purified, reverse transcribed, and the levels of indicated transcripts were analyzed by qPCR. Splice sites in the full length AR pre-mRNA were detected using ESRsearch and ESEFinder programs (Supplementary Fig. S1).

**Human clinical specimens**

Paired benign and tumor prostate tissue extracts were described previously (26). Total RNAs from human clinical specimens used for measurement of splicing factor transcript levels were described previously (27).

**Gene expression omnibus analysis**

Two separate data sets from NCBI GEO were screened independently for expression levels of hnRNP A1, hnRNP A2, U2AF65, and SF2/ASF. GDS1439 (28) compared specimens of benign prostatic hyperplasia with clinically localized primary prostate cancer and metastatic prostate cancer. GDS2545 (29) compared normal prostate specimens without any pathology, normal prostate adjacent to tumor, primary prostate tumor, and metastatic...
prostate cancer. Significant differences between groups were determined using Microsoft Excel Tools.

**Oncomine analysis**
Data generated from four comparisons of normal prostate tissue with prostate carcinoma: Lapointe prostate (30), Wallace prostate (31), Singh prostate (32), and Yu prostate (33) were analyzed using the differential expression function of Oncomine.

**Statistical analyses**
Data are shown as means ± SD. Multiple group comparison was performed by one-way ANOVA followed by the Scheffe procedure for comparison of means. $P \leq 0.05$ was considered significant.

**Results**

HnRNPA1 regulates the expression of AR variants
To test whether generation of AR variants by alternative splicing is dependent upon expression of hnRNPs, we analyzed expression levels of full-length AR and of variants such as AR-V7, AR-V1, AR-V5, AR-1/2/2b, and AR-1/2/3/2b using qRT-PCR in 22Rv1 and VCaP prostate cancer cells transfected with siRNAs against hnRNPA1 and hnRNPA2. Downregulation of hnRNPA1 and hnRNPA2 decreased the expression levels of AR variants in 22Rv1 and VCaP cells (Fig. 1A and B; Supplementary Fig. S2A and S2B). insets in Fig. 1A and B confirm the downregulation of hnRNPA1 and hnRNPA2 by specific siRNAs. The downregulation of hnRNPA1 and the resultant suppression of AR-V7 protein levels were confirmed by Western blot analysis (Fig. 1C). The protein levels of AR-V7 variant were decreased in both 22Rv1 and VCaP cells transfected with hnRNPA1 siRNA. These results indicate that hnRNPA1 may regulate the generation of AR-Vs in prostate cancer cells.

Next, we tested whether overexpression of hnRNPA1 affects the expression levels of AR-Vs. LNCaP cells were transfected with full-length hnRNPA1 cDNA and levels of AR-Vs were analyzed by Western blotting and qRT-PCR. Overexpression of hnRNPA1 enhanced AR-V7 protein levels in LNCaP cells, which possess undetectable endogenous levels of AR-V7 protein (Fig. 1D). qRT-PCR confirmed that overexpression of hnRNPA1 significantly enhanced the mRNA levels of AR-V7, AR-V5, AR-1/2/2b, and AR-1/2/3/2b variants in LNCaP cells (Fig. 1E; Supplementary Fig. S2C). Inset in Fig. 1E confirms the overexpression of hnRNPA1 after transfection in LNCaP cells. These results using downregulation as well as overexpression of hnRNPA1 suggest that hnRNPA1 plays an important role in the generation of AR splice variants.

Recruitment of hnRNPA1 to splice sites in AR pre-mRNA is increased in enzalutamide-resistant cells
We found hnRNP binding sites (UAGGGA) in the full-length AR mRNA using sequence analysis and ESRSearch program (Supplementary Fig. S1). To determine whether hnRNPA1 is recruited to splice sites in the AR pre-mRNA, we performed RNA immunoprecipitation (RIP) assays using specific antibodies against hnRNPA1 and hnRNPA2 in 22Rv1 versus 22Rv1-Enza-R and C4-2B versus C4-2B-Enza-R cell lines. The 22Rv1-Enza-R and C4-2B-Enza-R cell lines were generated by chronic exposure to enzalutamide and display resistance to enzalutamide (22, 23). The degree of recruitment of hnRNPA1 to AR-V7 splice sites was significantly higher in 22Rv1-Enza-R cells compared with parental 22Rv1 cells (Fig. 2A), indicating that hnRNPA1 may promote
generation of AR-V7 in prostate cancer cells resistant to enzalutamide. Even though the recruitment of hnRNPA2 to AR-V7 splice sites was also enhanced in 22Rv1-Enza-R cells compared with 22Rv1 cells, the recruitment of hnRNPA1 was several fold higher than that of hnRNPA2 (Fig. 2A). No significant differences were observed in the recruitment of either hnRNPA1 or hnRNPA2 to FL AR splice sites between 22Rv1 parental and 22Rv1-Enza-R cells (Fig. 2B). We also analyzed recruitment of hnRNPA1 and hnRNPA2 to splice sites for other AR-Vs such as AR-V1, AR-V5, AR-1/2/2b, and AR-1/2/3/2b (Supplementary Fig. S3A–S3D). Recruitment of hnRNPA1 to AR-1/2/2b splice sites was significantly higher in 22Rv1-Enza-R cells (Supplementary Fig. S3C), indicating that hnRNPA1 may play a selective role in generation of AR splice variants. In addition, recruitment of hnRNPA2 to AR-1/2/3/2b splice sites was significantly higher in 22Rv1-Enza-R cells (Supplementary Fig. S3C), indicating that hnRNPA1 may play a selective role in generation of AR splice variants leading to enzalutamide resistance.

To confirm the above results in another enzalutamide-resistant cell line, we analyzed the C4-2B versus C4-2B-Enza-R cell line pair. Our results showed that recruitment of hnRNPA1 and hnRNPA2 to AR-V7 splice sites was significantly enhanced in C4-2B-Enza-R cells compared with parental C4-2B cells (Fig. 2C), whereas, similar to 22Rv1-Enza-R cells, recruitment of either hnRNPA1 or hnRNPA2 to FL AR splice sites was not altered significantly in C4-2B-Enza-R cells compared with parental C4-2B cells (Fig. 2D). In all cases, the fold enrichment of hnRNPA1 at splice sites on AR pre-mRNA was much higher compared with hnRNPA2, indicating that hnRNPA1 may play a more central role in promoting the expression of AR-Vs (Supplementary Fig. S3E–S3H). These results collectively demonstrate that different splicing factors may play context- and cell type-dependent roles in prostate cancer cells in alternative splicing of the AR.

Expression levels of hnRNPA1 are elevated in prostate cancer tissues

To determine whether increased expression of splicing factors and AR-Vs is associated with prostate cancer, we examined the expression levels of hnRNPA1 and hnRNPA2 by immunoblotting in lysates from 27 archived paired benign and tumor prostate clinical samples. Levels of hnRNPA1 and hnRNPA2 were elevated in ~44% of tumor tissues compared with matched benign tissues (Fig. 3A; Supplementary Fig. S4A). These results were correlated positively with the protein expression levels of AR-V7, which were enhanced in ~48% of tumor tissues compared with their benign counterparts (Fig. 3A and Supplementary Fig. S4A). In addition, expression levels of hnRNPA1 and hnRNPA2 were low or undetectable in 9 of 12 and 6 of 12 donor prostates, respectively. These observations
were also correlated with expression levels of AR-V7, which were low or undetectable in 8 of 12 donor tissues (Table 1).

We also analyzed the mRNA levels of hnRNPA1, hnRNPA2, and AR-V7 in archival total RNAs extracted from 10 pairs of matched benign and tumor clinical prostate specimens (27, 34). Transcript levels of hnRNPA1 were elevated in 5 of 10 tumor tissues compared with matched benign tissues with no appreciable differences between tumor and benign being observed in the other 5 of 10.
samples (Fig. 3B). Transcript levels of AR-V7 were elevated in 6 of 10 tumor tissues compared with their matched benign counterparts (Fig. 3C), demonstrating that expression of hnRNPA1 and AR-V7 may be positively correlated with each other in human prostate cancer. No significant differences were observed in mRNA levels of hnRNPA2 between matched tumor and benign prostate tissues (Supplementary Fig. S4B).

To further confirm our findings, we analyzed expression levels of hnRNPA1 and hnRNPA2 in clinical prostate cancer tissues using publicly available data sets from Gene Expression Omnibus (GEO) and Oncomine. Results from an analysis of Oncomine data sets revealed that expression levels of hnRNPA1 and hnRNPA2 are significantly elevated in prostate tumor tissues compared with benign prostate tissues in 17 of 21 and 15 of 21 data sets, respectively (Supplementary Fig. S4C). Results from a representative data set, Singh_prostate (n = 102) from Oncomine are shown (Fig. 3D and E). An analysis of GEO revealed that expression levels of hnRNPA1 and hnRNPA2 were elevated in primary as well as metastatic prostate cancer compared with benign prostate tissues (Fig. 3F and G and Supplementary Fig. S4D). Data regarding expression levels of AR-Vs were not available in these data sets, but nonetheless, these results indicate that elevated levels of hnRNPA1 may contribute to prostate cancer development and progression. Our findings correlate well with studies showing that elevation of AR-V7 is one of the transcription factors which regulate transcription via alternative splicing. Previous studies also showed that c-Myc expression transcriptionally, whereas hnRNPA1 regulates c-Myc expression levels of splicing factors (2, 35), indicating that elevated expression of hnRNPA1 in prostate tumors may contribute to generation of higher levels of AR-Vs.

Expression of hnRNPA1 is regulated by c-Myc

Previous studies indicated that hnRNPA1 and c-Myc exhibit positive reciprocal regulation (36). C-Myc enhances hnRNPA1 expression transcriptionally, whereas hnRNPA1 regulates c-Myc via alternative splicing. Previous studies also showed that c-Myc is one of the transcription factors which regulate transcription of the AR (37). Hence, we analyzed the status of c-Myc or hnRNPA1 when the expression of either was downregulated in prostate cancer cells. Lysates from 22Rv1 and VCaP cells transfected with shRNA against c-Myc were analyzed using specific antibodies against hnRNPA1. Downregulation of c-Myc reduced protein levels of hnRNPA1 significantly (Fig. 4A). Similarly, lysates from LNCaP, 22Rv1, and VCaP cells transfected with siRNA against hnRNPA1 were examined by immunoblotting using specific antibodies against c-Myc. Downregulation of hnRNPA1 reduced protein levels of c-Myc (Fig. 4B). These results confirm that hnRNPA1 and c-Myc exhibit reciprocal regulation in prostate cancer cells. We also analyzed whether reduction in hnRNPA1 levels by c-Myc shRNA affects expression of AR-Vs in 22Rv1 and VCaP cells. Western blotting and qRT-PCR analyses showed that levels of AR-Vs, including that of AR-V7, were abrogated due to depletion of hnRNPA1 caused by downregulation of c-Myc (Fig. 4A–D; Supplementary Fig. S5). These findings support an important role for c-Myc in the generation of AR splice variants.

To analyze whether overexpression of hnRNPA1 can overcome the effects of downregulation of c-Myc, we suppressed expression of c-Myc using shRNA in 22Rv1 cells followed by overexpression of hnRNPA1. The results showed that even though suppression of c-Myc reduced mRNA as well as protein levels of both FLAR and AR-V7, subsequent overexpression of hnRNPA1 restored the expression of AR-V7 fully while having minimal effect on FLAR (Fig. 4E and F). These results indicate that hnRNPA1 primarily regulates generation of alternative splice variants of the AR and not the generation of the FLAR.

Table 1. Summary of the Western blot analysis of levels of hnRNPA1, hnRNPA2, and AR-V7 in 27 paired benign and tumor prostate clinical samples

<table>
<thead>
<tr>
<th>Gene</th>
<th>T &gt; B</th>
<th>T &lt; B</th>
<th>T = B</th>
<th>High in donor</th>
<th>Low in donor</th>
</tr>
</thead>
<tbody>
<tr>
<td>hnRNPA1</td>
<td>12 (44%)</td>
<td>6 (29%)</td>
<td>7 (26%)</td>
<td>3/12</td>
<td>9/12</td>
</tr>
<tr>
<td>AR-V7</td>
<td>13 (48%)</td>
<td>7 (25%)</td>
<td>7 (26%)</td>
<td>4/12</td>
<td>8/12</td>
</tr>
<tr>
<td>hnRNPA2</td>
<td>12 (44%)</td>
<td>8 (29%)</td>
<td>7 (26%)</td>
<td>6/12</td>
<td>6/12</td>
</tr>
</tbody>
</table>

NF-κB2/p52 regulates AR-V7 expression via hnRNPA1 and c-Myc

We reported earlier that activation of NF-κB2/p52 promotes progression to CRPC and enzalutamide resistance via the generation of AR variants, specifically AR-V7 (21, 23, 24). Our previous findings also indicated that NF-κB2/p52 may regulate c-Myc expression. Hence, we examined whether NF-κB2/p52 plays a role in the elevated expression of hnRNPA1 and c-Myc in prostate cancer using lysates from LNCaP cells stably expressing p52 (LN-p52). Protein levels of AR-V7, hnRNPA1, and c-Myc were elevated in LN-p52 cells, whereas no appreciable differences were found in the expression of hnRNPA2 (Fig. 5A, left). These results were confirmed using LNCaP cells expressing p52 under the control of a Tet-inducible promoter (LN/TR/p52). Induction of p52 expression led to increases in expression levels of AR-V7, hnRNPA1, and c-Myc (Fig. 5A, right), indicating that upregulation of AR-V7 by p52 may be mediated by hnRNPA1 and c-Myc. To examine the relationship between hnRNPA1, c-Myc and AR-V7 in LN-p52 cells, we analyzed levels of AR-V7 by Western blotting in LN-p52 cells transfected with hnRNPA1 or hnRNPA2 siRNAs. Downregulation of hnRNPA1 abrogated the expression of AR-V7 in LN-p52 cells, whereas downregulation of hnRNPA2 did not have an appreciable effect on AR-V7 protein levels (Fig. 5B, left). Protein levels of c-Myc were also downregulated, keeping in line with earlier findings that hnRNPA1 and c-Myc regulate each other (36). In addition, expression of hnRNPA1 and AR-V7 were abolished as a result of downregulation of c-Myc expression in LN-p52 cells transfected with c-Myc shRNA (Fig. 5B, middle).

To confirm these findings in a cell line with constitutive expression of both p52 and AR-V7, we analyzed levels of AR-V7, hnRNPA1, and c-Myc by immunoblotting in 22Rv1 cells transfected with p52 shRNA. Downregulation of p52 in 22Rv1 cells abrogated expression of AR-V7, hnRNPA1, and c-Myc (Fig. 5B, right). These results demonstrate that NF-κB2/p52 may modulate generation of AR-Vs by regulation of hnRNPA1 and c-Myc.

Prostate cancer cells resistant to enzalutamide exhibit higher levels of splicing factors

As our results demonstrate that expression of hnRNPA1 and AR variants may be positively correlated with each other in prostate cancer cells, and AR-V7 expression has been shown to be involved in the acquisition of resistance to enzalutamide (3, 23), we tested the correlation between levels of AR variants and hnRNPA1 in
prostate cancer cells that have acquired resistance to enzalutamide. 22Rv1-Enza-R and C4-2B-Enza-R cell lines exhibited higher levels of AR-V7 and hnRNPA1 (Fig. 5C), indicating that expression of hnRNPA1 may be positively correlated with expression of AR-Vs. Furthermore, expression levels of c-Myc and NF-kB2/p52 were also elevated in enzalutamide-resistant cells, confirming the importance of the NF-kB2/p52:c-Myc:hnRNPA1:AR-V7 axis in enzalutamide resistance. No significant differences were observed in the expression of hnRNPA2 in 22Rv1-Enza-R and C4-2B-Enza-R cells compared with their parental cells (Fig. 5C, right and middle). To confirm these results in vivo, we analyzed extracts from xenograft tumors derived from C4-2B and C4-2B-Enza-R cells using antibodies against AR-V7 and hnRNPA1. Higher levels of AR-V7 were observed in xenografts derived from C4-2B-Enza-R cells, which was correlated well with higher levels of hnRNPA1 and c-Myc (Fig. 5C, right), confirming our observations that expression of AR-Vs is positively correlated with expression of hnRNPA1 in prostate cancer cell lines resistant to enzalutamide.

Next, we tested whether downregulation of hnRNPA1 affects endogenous levels of AR-Vs in 22Rv1-Enza-R cells. Transfection of hnRNPA1 siRNA abrogated expression levels of AR-V7 in 22Rv1-Enza-R cells (Fig. 5D, left). Similarly, downregulation of c-Myc by specific shRNA reduced expression levels of AR-V7 and hnRNPA1 in 22Rv1 and 22Rv1-Enza-R cells (Fig. 5D, right), confirming the c-Myc:hnRNPA1:AR-V7 axis in prostate cancer cells.

To confirm the importance of the link between NF-kB2/p52, c-Myc, hnRNPA1, and AR-V7 in prostate cancer, we analyzed the correlation between their expression levels at mRNA and protein levels in paired benign and tumor prostate clinical samples from Fig. 3. Transcript (left) and protein (right) levels of NF-kB2/p52, c-Myc, hnRNPA1, and AR-V7 were positively correlated with each other (Fig. 5E), demonstrating that the NF-kB2/p52:c-
Myc:hnRNPA1:AR-V7 axis plays a vital role in prostate cancer and in the development of castration and therapy resistance.

Suppression of hnRNPA1 resensitizes enzalutamide-resistant prostate cancer cells to enzalutamide

To examine the functional relevance of regulation of AR alternative splicing by hnRNPA1, we tested whether down-regulation of hnRNPA1 resulting in decreased levels of AR-V7 resensitizes enzalutamide-resistant cells to enzalutamide. We examined cell survival in 22Rv1 and 22Rv1-Enza-R cells transfected with hnRNPA1 or hnRNPA2 siRNAs and treated with 0 and 20 μmol/L enzalutamide for 48 hours. Reduced expression of hnRNPA1 enhanced the sensitivity of enzalutamide-resistant 22Rv1-Enza-R cells to enzalutamide (Fig. 6A), indicating that upregulation of AR-V7 expression by hnRNPA1 may be required to sustain the acquired resistance of 22Rv1-Enza-R cells to enzalutamide. Downregulation of hnRNPA1 and hnRNPA2 also enhanced the sensitivity of LN-neo and LN-p52 cells transfected with hnRNPA1 or hnRNPA2 siRNAs and treated with 20 μmol/L enzalutamide to enzalutamide (Fig. 6B). Suppression of hnRNPA1 expression reduced cell survival by ~40% to 50% when enzalutamide-resistant LN-p52 cells were treated with enzalutamide. Suppression of hnRNPA1 expression also reduced survival of VCaP cells when treated with enzalutamide (Fig. 6C), confirming the essential nature of AR variants in these cells.
Figure 6.
Suppression of hnRNPA1 restores enzalutamide sensitivity of enzalutamide-resistant prostate cells. A, left, cell survival in 22Rv1-Enza-R cells transfected with hnRNPA1 or hnRNPA2 siRNAs and treated with vehicle or 20 μmol/L enzalutamide. Cell numbers were counted after 48 hours; right, immunoblots confirm the downregulation of hnRNPA1 or hnRNPA2 and of AR-V7. B, left, cell survival in LN-p52 cells transfected with hnRNPA1 or hnRNPA2 siRNAs and treated with vehicle or 20 μmol/L enzalutamide. (Continued on the following page.)
Our findings collectively demonstrate that hnRNPA1 plays a major role in the generation of AR splice variants. Expression of hnRNPA1 is modulated by NF-κB2/p52 via c-Myc (Fig. 6D). Our results point to the enhanced expression of hnRNPA1 in prostate tumors being instrumental in inducing alternative splicing of the precursor AR mRNA.

Discussion

A large number of previous studies have shown that C-terminally truncated AR-Vs are expressed in prostate cancer cells and even in normal prostate epithelial cells (35, 38) and promote CRPC progression under androgen deprivation (39, 40). In addition, enhanced expression of AR-V7 confers resistance to next-generation therapeutics such as enzalutamide and abiraterone (4, 5). These studies attest to the importance of AR-Vs in CRPC and cotargeting the mechanisms which contribute to their generation may increase the efficacy of currently used AR-targeted therapies and prolong time to development of resistance. Our findings in this study demonstrate that the splicing factor hnRNPA1 plays a major role in the alternative splicing of AR mRNA.

HnRNPA1 is a multifunctional RNA-binding protein involved in the regulation of RNA biogenesis. HnRNPA1 is under the transcriptional control of the c-Myc proto-oncoprotein and modulates the splicing of PKM2, activating the metabolic switch to aerobic glycolysis that is a hallmark of cancer cells (11, 36). HnRNPA1 also regulates alternative splicing of genes involved in invasion and metastasis such as Rac1 and Ror (11). Increased expression of hnRNPA1 has been documented in proliferating and transformed cells (20) and in lung, breast, colon, renal cell carcinomas, and gliomas (41–46). HnRNPs cooperate with other splicing factors to promote pro-oncogenic and proinflammatory molecules in cancers (41, 47). Silencing of hnRNPA1 and A2 promotes apoptosis in human and mouse cancer cell lines, while having no effect on normal epithelial and fibroblastic cell lines (48).

In this study, hnRNPA1 binding sites (UAAGGA) were identified in AR mRNA using sequence analysis and ESRSearch program. Downregulation of hnRNPA1 significantly reduced the expression of AR-Vs such as AR-V7, while not affecting full-length AR. HnRNPA1 binding sites were not detected at full length AR splice sites in the AR pre-mRNA. Moreover, the slight decrease seen in either full length AR mRNA or protein levels in VCaP or 22Rv1 cells respectively (Fig. 1) was not observed consistently in all experiments. In consideration of these observations, we concluded that hnRNPA1 does not play a significant role in splicing of full length AR from AR pre-mRNA. RNA-binding assays revealed that hnRNPA1 is recruited to splice sites for AR splice variants. Enhanced expression of hnRNPA1 was observed in prostate cancer tissues compared with their benign counterparts, which was correlated positively with expression of AR-V7. Increased expression of hnRNPA1 was also correlated with higher levels of AR-V7 in prostate cancer cells with acquired resistance to enzalutamide. Exploration of the mechanisms revealed that c-Myc and NF-κB2/p52 contribute to the development of therapy resistance in prostate cancer cells by inducing hnRNPA1 expression and thereby ligand-independent AR-Vs. Downregulation of hnRNPA1 sensitized enzalutamide-resistant cells to enzalutamide, indicating that suppression of hnRNPA1 resulting in suppression of AR-Vs reversed the acquired resistance to enzalutamide. These data led us to conclude that hnRNPA1 is the central player in a splicing regulatory circuit involving c-Myc, NF-κB2/p52, and AR.

Our study demonstrates that elevated levels of splicing factors such as hnRNPA1 promote expression of alternative splice forms of AR. Relative amounts of splicing factors have been proposed to determine alternative splicing (45). A recent study by Liu and colleagues (49) showed that recruitment, and not expression, of splicing factors SF2/ASF and U2AF65 determines the generation of AR splice variants in enzalutamide resistance. We found higher levels of hnRNPA1 in prostate cancer clinical samples compared with SF2/ASF (data not shown), indicating that higher expression of a splicing factor, and not simply its recruitment to splice sites under certain conditions, may determine the levels of alternative splice forms. These results are supported by previous studies which showed that the relative levels of hnRNPA1 expression increased to a greater extent than those of SF2/ASF in lung tumors (44). Of note, analysis of expression levels of U2AF65 and SF2/ASF in prostate cancer tissues using GEO and Oncomine data sets revealed that expression levels of both splicing factors are elevated in prostate cancer tissues compared with benign counterparts (Supplementary Fig. S6), lending credence to our observations that enhanced expression of splicing factors may be one of the principal mechanisms driving generation of AR-Vs. Elevated levels of hnRNPA1 may conceivably change the splicing milieu of a broad spectrum of proteins in addition to that of the AR, with splicing of CD44 being an example. But splice variants of AR have been demonstrated to play major roles in resistance to enzalutamide, underlining the importance of our results in splicing regulation of the AR. Furthermore, earlier studies indicated that the splice variant AR-1/2/3/2b is also generated by intragenic genomic rearrangement of the AR gene due to duplication of exon 3 (1). Our studies consistently found transcripts corresponding to this splice variant in LNCaP, C4-2B, and VCaP cells, albeit at extremely low levels. These observations warrant further exploration of the mechanisms involved in generation of this splice variant and further validation.

In summary, we demonstrated that hnRNPA1, in concert with NF-κB2/p52 and c-Myc, regulates the generation of AR-Vs in prostate cancer cells and that the NF-κB2/p52:c-Myc:hnRNPA1:AR-V7 axis (Fig. 6D) plays a pivotal role in the development and maintenance of resistance to androgen blockade. These findings may have important implications in targeting AR-Vs and the splicing factors responsible to overcome acquired enzalutamide resistance in prostate cancer.

(Continued.) Cell numbers were counted after 48 hours; right, immunoblots confirm the downregulation of hnRNPA1 or hnRNPA2 and AR-V7. C, left, cell survival in VCaP cells transfected with hnRNPA1 or hnRNPA2 siRNAs and treated with vehicle or 20 μmol/L enzalutamide. Cell numbers were counted after 48 hours; right, immunoblots confirm the downregulation of hnRNPA1 or hnRNPA2 and of AR-V7. Results are presented as means ± SD of three experiments performed in triplicate. *P < 0.05. D, schematic representation of the alternative splicing of AR mRNA regulated by the NF-κB2:c-Myc:hnRNPA1 axis.
Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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