Androgen Withdrawal in Patients Reduces Prostate Cancer Hypoxia: Implications for Disease Progression and Radiation Response

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Abstract

Hypoxia is a feature of many human malignancies, and leads to aggressive clinical behavior and recurrence after treatment. Here, we show for the first time that androgen withdrawal reduces prostate cancer hypoxia in patients. Oxygen measurements were done in 248 patients with clinically localized prostate cancer prior to radiotherapy, and showed hypoxia of potential biological and clinical significance. In 22 of these patients, prostate oxygen levels were measured both before and after 30 to 145 days of the androgen antagonist bicalutamide. There was a significant reduction in tumor hypoxia with androgen withdrawal (P = 0.005). The median pO2 increased from 6.4 to 15 mm Hg, and the hypoxic proportion decreased from 40% to 31%. However, the response was heterogeneous, with improvement in 12 patients, stable oxygen readings in 9 patients and worsening hypoxia in 1 patient. Among the responding patients, the median pO2 increased from 4.9 to 33 mm Hg, and the hypoxic proportion decreased from 51% to 23%. There was no apparent relationship between the change in oxygenation and baseline prostate volume, T category, Gleason score, prostate-specific antigen levels, the duration of treatment with bicalutamide, or the change in prostate-specific antigen levels with bicalutamide. These results might, in part, explain the improved patient outcome that has been observed in clinical trials of radiotherapy and hormones, and suggest a role for novel therapeutic agents that block the molecular response to hypoxia in prostate cancer either alone or in combination with other established treatments. [Cancer Res 2007;67(13):6022–5]

Introduction

Prostate cancer is the most common malignancy among North American men. At diagnosis, most patients have tumors that are clinically confined to the prostate gland. Depending on comorbidities and individual preferences, they are candidates for potentially curative treatment with either radical prostatectomy or radiotherapy. Despite technical advances in both of these treatments, ~25% of radically treated patients will develop progressive disease either locally in the pelvis or at remote sites, most commonly in bone (1). This underscores the importance of better understanding the biological factors that are responsible for malignant progression, the development of metastases, and the failure of currently available treatments.

Androgens play an important role in the growth and progression of prostate cancer. Androgen withdrawal through surgical or medical castration causes tumor regression in most patients. Several phase III studies have shown improved tumor control and patient survival when radiotherapy is combined with androgen withdrawal to treat prostate cancer (2), and combined treatment is widely used in clinical practice. Despite this, the biological mechanisms responsible for the favorable interaction between radiotherapy and androgen deprivation remain ill-defined.

Hypoxia is a feature of many human malignancies. In general, patients with hypoxic primary tumors at diagnosis are at greater risk of developing progressive disease and dying regardless of whether initial treatment is with surgery or radiotherapy (3). These clinical observations are consistent with hypoxia-mediated changes in DNA repair, genomic instability, and abnormal expression of genes that promote both malignant progression and metastasis formation (4). Faulty DNA repair is an important determinant of genetic instability and contributes to chromosomal rearrangement, oncogene activation, and tumor suppressor gene inactivation (5). Repair-deficient cells are more likely to acquire a mutator phenotype characterized by greater biological and clinical aggressiveness. Indeed, preclinical data have shown that hypoxia could lead to the selection of aggressive cancer cells with decreased sensitivity to apoptotic and DNA repair pathways and increased genetic instability, angiogenesis, proliferation, and metastatic capability (4, 5).

We and others have previously reported that many prostate cancers are hypoxic (6–8). Here, we show for the first time that androgen withdrawal reduces prostate cancer hypoxia in patients, and discuss the implications of this for prostate cancer development and progression, metastasis formation, and response to radiotherapy.

Materials and Methods

A total of 248 patients with \( cT_{1c}-T_{2a}, N_0, M_0 \) (UICC-TNM Classification, 6th Edition, 2002) prostate cancer participated in this prospective study of tumor hypoxia prior to treatment with escalated-dose conformal or intensity-modulated radiotherapy. The characteristics of these patients are summarized in Table 1. The study was approved by the University Health Network Research Ethics Board, and informed consent was obtained in all cases.

Prostate cancer hypoxia was measured prior to any treatment using an ultrasound-guided transrectal needle-electrode technique, as previously described (7). Between 40 and 80 individual oxygen readings were obtained along two to four linear measurement tracks 1.5 to 2 cm in length through regions of the prostate gland likely to contain tumor based on information from previous diagnostic biopsies, digital rectal examination, and real-time
Table 1. Characteristics of patients with prostate cancer oxygen measurement

<table>
<thead>
<tr>
<th></th>
<th>Patients with oxygen measurements prior to treatment only</th>
<th>Patients with oxygen measurements both pre- and post–androgen withdrawal</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>226</td>
<td>22</td>
</tr>
<tr>
<td>Age*</td>
<td>71 (55–82)</td>
<td>71 (55–79)</td>
</tr>
<tr>
<td>T category</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T1</td>
<td>87 (39%)</td>
<td>12 (54%)</td>
</tr>
<tr>
<td>T2</td>
<td>135 (60%)</td>
<td>10 (46%)</td>
</tr>
<tr>
<td>T3</td>
<td>4 (2%)</td>
<td></td>
</tr>
<tr>
<td>Gleason score</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>67 (30%)</td>
<td>3 (14%)</td>
</tr>
<tr>
<td>7</td>
<td>146 (65%)</td>
<td>18 (82%)</td>
</tr>
<tr>
<td>8, 9</td>
<td>13 (6%)</td>
<td>1 (4%)</td>
</tr>
<tr>
<td>PSA* (ng/mL)</td>
<td>8.0 (0.9–33)</td>
<td>7.8 (3.3–30)</td>
</tr>
<tr>
<td>Prostatic volume* (cm³)</td>
<td>40 (13–179)</td>
<td>37 (23–171)</td>
</tr>
<tr>
<td>Grand median pO2* (mm Hg)</td>
<td>6.8 (0–75)</td>
<td>5.5 (1.8–61)</td>
</tr>
<tr>
<td>Median HP5* (%)</td>
<td>33 (0–100)</td>
<td>46 (0–93)</td>
</tr>
</tbody>
</table>

*Median and range.
†Clinical T category. All patients were clinically N0, M0 (UICC-TNM Classification, 6th Edition, 2002).
‡Values prior to androgen withdrawal.

Results and Discussion

This is the largest reported study of patients with prostate cancer who have undergone assessment of tumor hypoxia. A total of 14,038 individual oxygen measurements were done in 226 patients prior to treatment, and includes the cohort of 55 patients that we previously described (7). In addition, an additional 3,388 measurements were done in 22 patients who were evaluated both before and after androgen withdrawal. In the larger group, the pretreatment grand median pO2 was 6.8 mm Hg (range, 0–75 mm Hg) and the median HP5 was 33% (range, 0–100%), as indicated in Table 1. These are similar to previously reported results for prostate cancer (8) and other human tumors (3), and indicate hypoxia of potential clinical and biological significance. The pretreatment grand median pO2 and median HP5 values in the androgen withdrawal group were in the same range at 5.5 mm Hg (range, 1.8–61 mm Hg) and 46% (0–93%), respectively.

There were statistically significant (P < 0.0001) differences in pretreatment hypoxia among the four measurement tracks and as a function of depth in the gland. In the 226 patients who only had pretreatment assessment of hypoxia, the median pO2 values for the four tracks varied from 6.2 to 9.8 mm Hg. The individual oxygen readings were designated as arising from either the superficial (subcapsular) or deep prostate gland by dividing the measurements along each linear track into two equal groups. The deeper regions of the gland were more hypoxic than the superficial regions (median pO2 of 7 versus 9 mm Hg, respectively). Similar results were obtained when the analysis was repeated using the pretreatment oxygen values for the 22 patients in whom measurements were done both before and after androgen withdrawal. Therefore, measurement track number and depth were included as covariates in the analysis of androgen withdrawal to minimize the potential for bias in the results.

Androgen withdrawal with bicalutamide produced a significant (P = 0.005) reduction in prostate hypoxia among the 22 patients who had both pretreatment and posttreatment measurements. The median pO2 increased from 6.4 to 15 mm Hg and the HP5...
Androgens seem to directly stimulate angiogenesis in hormone-sensitive prostate cancer through activation of growth factor receptors and the phosphatidylinositol-3-kinase/AKT signaling pathway, leading to hypoxia-independent up-regulation of HIF-1α and VEGF expression (17–19). This probably occurs early in prostate cancer development (18), and triggers a cascade of increasing angiogenesis and hypoxia. Hypoxia in turn may sensitize tumor cells to very low levels of androgens through a receptor-mediated mechanism (20). The variable oxygen response to bicalutamide that we observed in this study may in part be due to differences in the relative importance of androgenic and hypoxic stimulation of angiogenesis in individual hormone-sensitive tumors. Our future research will focus on better understanding this heterogeneity among patients.

The reduction in prostate cancer hypoxia with androgen withdrawal shown in this study might in part explain the improved patient outcome that has been observed in many clinical trials of combined treatment with radiotherapy and hormones (2). However, there may be more far-reaching implications. We hypothesize that androgen-induced angiogenesis and hypoxia in nonmalignant regions of the prostate gland may be important in prostate carcinogenesis. Our group has observed aberrant DNA repair in hypoxic prostate cancer cells, which supports hypoxia as a mediator of genetic instability in this disease (5). Therefore, novel therapeutic strategies, which inhibit HIF-1α or other downstream molecular targets, might not only reduce morbidity and mortality in men with diagnosed prostate cancer but might also play an important role in prostate cancer prevention.

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