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Stereotactic Ablative Radiation Therapy for Oligometastatic Prostate Cancer Delays Time to Next Systemic Treatment --Manuscript Draft--

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Abstract:	<p>Purpose: Local ablative treatment to oligometastatic patients can result in long-term disease-free survival in some cancer patients. The importance of this treatment paradigm in prostate cancer is a rapidly evolving field. Herein we report on the safety and preliminary clinical outcomes of a modern cohort of oligometastatic prostate cancer (OPC) patients treated with consolidative stereotactic ablative radiation (SABR). Methods: Records of men with OPC who underwent consolidative SABR at our institution were reviewed. SABR was delivered in 1-5 fractions of 5-18 Gray. Kaplan-Meier estimates of local progression-free survival (LPFS), biochemical progression-free survival (bPFS; PSA nadir+2), distant progression free survival (DPFS) and time-to-next intervention (TTNI) were calculated.</p> <p>Results: In total, 66 OPC patients were identified with consolidative SABR delivered to 134 metastases: 89 bone, 40 nodal, and 5 viscera. The majority of men (49/66) had hormone sensitive prostate cancer (HSPC). Crude Grade 1 and 2 acute toxicities were 36% and 11% respectively with no \geq Grade 3 toxicity. At 1-year LPFS was 92% and bPFS and DPFS were 69%. Of the 18 men with HSPC who had hormone therapy deferred, 11 (56%) remain disease free following SABR (1-year ADT-FS was 78%). In 17 castration resistant men, 11 had $> 50\%$ PSA declines with 1-year TTNI of 30%.</p> <p>Conclusions: Consolidative SABR in OPC is feasible and well tolerated. The heterogeneity and small size of our series limits extrapolation of clinically meaningful outcomes following consolidative SABR in OPC, but our preliminary data suggests this approach warrants continued prospective study.</p>	
Response to Reviewers:	<p>Response to reviewer's comments:</p> <p>Reviewer #1: A retrospective analysis of 66 patients with oligometastatic prostate cancer treated with consolidative SBRT. Importantly, and unique to this study, 53% of patients actually had 2 or more lesions. The majority of the lesions were osseous, as expected, and the majority of men (74%) had hormone sensitive disease. There were no grade 3 or higher toxicities of any kind, and average time to failure of any kind was 35 weeks. Of the 18 men with hormone sensitive disease who sought to avoid ADT, 61% were able to remain free of ADT at 1 year. Among patients with mCRPC, the majority (11/17) had $>50\%$ decline in PSA. This is important as the ability to delay chemotherapy in men with CRPC is an important goal.</p> <p>Overall, this paper warrants publication in the World Journal of Urology. It is well written and the data are presented clearly. The manuscript does not overstate its findings and is framed appropriately. A major strength is that the majority of patients had 2 or more lesions. The clear description of outcomes in patients with mCRPC vs. hormone-sensitive disease is timely as well.</p> <p>Strengths: The manuscript is well written and timely. The authors present the data in a clear, concise, and straightforward manner.</p> <p>Weaknesses: Limited by small numbers and single institution nature. These limitations are acknowledged by the authors.</p> <p>REPLY - We appreciate this reviewer's comments and time dedicated to assisting in the publication of this paper.</p> <p>Reviewer #2: This is a retrospective study evaluating the tolerability and efficacy of a focal SBRT approach for oligo-metastatic patients with prostate cancer. The article is well written and clearly understandable.</p> <p>The message is of interest in the literature, in particular because it takes into account patients with more than 2 metastatic sites, but also because it includes some notions still confidential about the place of loco-regional treatment of patients presenting oligo-metastatic castration resistant disease.</p> <p>MINOR COMMENTS:</p> <p>- In the material and methods section, please define how the next intervention was decided?</p> <p>For example, what type of criteria were retained to introduce ADT? Same question in the castration resistant population?</p>	

For example , in the STOMP trial : The indication to start ADT was symptomatic progression, progression to more than three metastases, or local progression of baseline-detected metastases.

REPLY - We appreciate the suggestion to clarify intervention indications. With the STOMP trial, the protocol for intervention indication and modality was clearly delineated from the start of the trial. However, with the retrospective nature of our study, the absolute indications for intervention timing and modality were subject to patient and provider variability within the scope of best-practices. Fortunately, the majority of patients were treated by a limited number of Hopkins medical oncologists will similar practice patterns. So broadly, the indication for next intervention was determined by disease objective evidence of disease progression by PSA and/or radiographic testing or symptomatic progression. We have placed this description in the Methods section.

- As proposed in the STOMP trial, authors should performed an analysis taking into account doubling time (< 3 or 6 months versus >3 or 6 months)

REPLY - We have performed the analysis as suggested and we were able to retrieve PSADT using the MSKCC on-line calculator for 33 of our patients (50%). However, we did not see any differences between rapid or slow PSADT for bPFS, LPFS or DMPS. This is likely too few events and is underpowered so we did not include this analysis in the paper.

- In the castration resistant arm, is there any treatment concomitantly given with SABR? Or patients were free of new agents?

REPLY - The patients of the castration resistant arm of our study received additional treatments as appropriate for their clinical disposition and disease burden. As the focus of the study centered on clinical outcomes of OPC patients receiving SABR, patient treatments did not explicitly influence inclusion in our retrospective review.

Perhaps it could be relevant to cite the brilliant paper from Beauval et al in this population to highlight the rationale and limitations of this approach (Loco-regional treatment for castration-resistant prostate cancer: Is there any rationale? A critical review from the AFU-GETUG. Beauval JB et al. Crit Rev Oncol Hematol. 2018 Feb;122:144-149).

REPLY - We appreciate the suggestion and have cited the paper from Beauval et al.

- A short paragraph of imaging limitations, especially concerning negative predictive value of classical imaging modalities, rending a focal approach discutable may be added in the discussion section. PSMA must be cited.

REPLY - Thank you for pointing this deficiency out. We have added text to address this issue in the Discussion.

[Click here to view linked References](#)

Stereotactic Ablative Radiation Therapy for Oligometastatic Prostate Cancer Delays Time to Next Systemic Treatment

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Abstract

Purpose: Local ablative treatment to oligometastatic patients can result in long-term disease-free survival in some cancer patients. The importance of this treatment paradigm in prostate cancer is a rapidly evolving field. Herein we report on the safety and preliminary clinical outcomes of a modern cohort of oligometastatic prostate cancer (OPC) patients treated with consolidative stereotactic ablative radiation (SABR).

Methods: Records of men with OPC who underwent consolidative SABR at our institution were reviewed. SABR was delivered in 1-5 fractions of 5-18 Gray. Kaplan-Meier estimates of local progression-free survival (LPFS), biochemical progression-free survival (bPFS; PSA nadir+2), distant progression free survival (DPFS) and time-to-next intervention (TTNI) were calculated.

Results: In total, 66 OPC patients were identified with consolidative SABR delivered to 134 metastases: 89 bone, 40 nodal, and 5 viscera. The majority of men (49/66) had hormone sensitive prostate cancer (HSPC). Crude Grade 1 and 2 acute toxicities were 36% and 11% respectively with no \geq Grade 3 toxicity. At 1-year LPFS was 92% and bPFS and DPFS were 69%. Of the 18 men with HSPC who had hormone therapy deferred, 11 (56%) remain disease free following SABR (1-year ADT-FS was 78%). In 17 castration resistant men, 11 had $> 50\%$ PSA declines with 1-year TTNI of 30%.

Conclusions: Consolidative SABR in OPC is feasible and well tolerated. The heterogeneity and small size of our series limits extrapolation of clinically meaningful outcomes following consolidative SABR in OPC, but our preliminary data suggests this approach warrants continued prospective study.

Key words: Oligometastatic prostate cancer, stereotactic ablative radiation therapy, androgen deprivation therapy

Introduction

Originally proposed by Hellman and Weichselbaum [1], the oligometastatic or low volume metastatic state is currently defined as having fewer than 3 to 5 metastatic lesions [2]. As the collective understanding of this state evolves, biological parameters may someday inform or supplant this descriptive clinical definition [3-5].

Improvements in imaging and metastasis-directed therapies (MDT) have placed new emphasis on the oligometastatic state [6]. Local ablative treatment for limited metastatic disease has been correlated retrospectively with improvements in long term disease-free survival in lung, breast, colorectal, and sarcoma patients [6] and mounting prospective evidence suggests improvements in progression-free survival for oligometastatic lung cancers and hormone sensitive oligorecurrent prostate cancer [7-10].

Traditionally, metastatic prostate cancer has been, regardless of tumor burden, treated systemically with androgen deprivation therapy (ADT) and/or chemotherapy [11]. The importance of consolidating all macroscopic tumor deposits in men with oligometastatic prostate cancer is an area of active investigation [9,12,13], but the observation that men with lower numbers of metastases have better outcomes [14-16] supports a fundamental difference between oligometastatic prostate cancer (OPC) and higher volume disease. By this logic, recent studies have explored MDT of OPC by ablative and surgical treatments [17-20], as well as stereotactic ablative radiation (SABR) [21], also known as stereotactic body radiation therapy (SBRT) [22]. This highly targeted, high-dose approach provides millimeter-scale precision while minimizing collateral tissue damage and has shown efficacy in the management of OPC [17-19,21,23,24].

Studies to date have been conducted predominately in patients with a single metastasis [18,19,17,23,24,9]. Herein we review our early institutional outcomes with SABR for OPC in 66 consecutively treated patients with the majority having ≥ 2 metastases. We demonstrate that SABR is well tolerated and can significantly delay time to subsequent intervention.

Methods

Criteria:

With approval from of our institutional review board, we reviewed our retrospectively collected Johns Hopkins prostate cancer (PCa) radiation oncology database for consecutively SABR treated patients from 02/07/2011 to 1/15/2017. Inclusion criteria were: biopsy-proven PCa, diagnostic imaging consistent with ≤ 5 metastases, and follow-up with post-SABR prostate specific antigen (PSA) testing. Three patients with ≤ 5 initially observed metastases who were found during treatment planning to have 6 metastases were included. Of the 84 total men during this period with OPC treated with SABR we only examined the first 66 men to allow for a minimum of 4.5-months of follow-up.

SABR technique:

During CT-simulation patients were immobilized using an Alpha Cradle (Smithers Medical Products, Inc, North Canton, Ohio) or equivalent device. Magnetic resonance imaging (MRI), bone scan, NaF or DCFPyL positron emission tomography [25]/CT images were used to delineate the gross tumor volume (GTV) and when possible fused with the planning CT. As per our institutional policy, if patients had ≥ 3 -mm breathing motion on 4-dimensional CT, active breathing control (ABC) was used for motion management. If breathing was < 3 mm, patients were treated free breathing (FB) with an internal target volume based on the 0% and 60% phases of the breathing cycle. GTV was defined as the sum of the abnormalities noted on PET, MRI T1 post-gadolinium sequences, bone scan and/or CT. In general, the clinical tumor volume (CTV) was equal to the GTV. Most commonly, the planning target volumes (PTV) were the GTV with the addition of 3-5 mm used as the volume to which the prescription was assigned [17]. If men were treated with CyberKnife, then image-guidance was per that device. During non-CyberKnife SABR delivery, a cone beam CT scan was co-registered (spine) with the FB or ABC simulation scan. To verify tumor positioning before SABR, patients were shifted to align with the GTV for each beam based on fluoroscopy, cone beam CT, or kV images.

Follow-up:

Patients were scheduled for serial follow-up exams every 3-6 months with history and physical, PSA, testosterone and imaging every 6-12 months unless a shorter interval was clinically indicated. Repeat imaging with NaF or DCFPyL PET/CT, MRI, or bone scans was conducted at 3-6 month intervals after SABR. The majority of patients were treated by a limited number of Johns Hopkins medical oncologists with similar practice patterns. So broadly, the indication for next intervention was determined by disease objective evidence of disease progression by PSA and/or radiographic testing or symptomatic progression.

Statistical analysis:

Clinical endpoints reported include biochemical progression-free survival (bPFS), local progression-free survival (LPFS), distant progression-free survival (DPFS), overall survival (OS) [25], ADT-free survival (ADT-FS), and time-to-next intervention (TTNI); all were calculated from the date of SABR. Biochemical failure was defined as: (1) PSA rise ≥ 2 above nadir following SABR or (2) an increase of PSA following SABR with failure to decline initially. LPFS was defined as the lack of tumor progression within the treated planning tumor volume (PTV) margin on follow-up imaging. DPFS was defined as the lack of new metastases on follow-up imaging. For both local and distant progression-free survival, clinically significant events (e.g. bone fractures) in a previously stable lesion were considered as clinical progression. TTNI was defined as the interval to next local or systemic intervention due to progression. The Kaplan-Meier method was used to estimate the rates of bPFS, LPFS, DPFS, OS, ADT-FS and TTNI. Analysis of LPFS was performed with each treated lesion considered individually ($n = 134$), while all other endpoints were analyzed with each patient considered individually.

Results

Patient and disease characteristics:

Sixty-six men with 134 metastatic PCa lesions were identified and analyzed. Prior treatment of metastatic tumors (including radiation and surgery) did not exclude patients from this review. Patients were generally asymptomatic or had minimal pain associated with their metastatic burden. Thirty-eight percent of men presented with synchronous or de novo oligometastases while 62% developed metachronous oligometastases between the conclusion of definitive treatment for their prostate cancer and consultation for SABR treatment. Prior chemotherapy was allowed with 55% ($n = 36$) of patients receiving chemotherapy prior to or concurrently with treatment. Forty-nine (or 74%) of men had hormone sensitive prostate cancer (HSPC) at the time of SABR and 17 (or 26%) had castration resistant prostate cancer (CRPC). Initial Gleason scores (GS) were as follows: GS 6 ($n = 7$), GS 7 ($n = 16$), GS 8 ($n = 18$), GS 9 ($n = 22$), and GS 10 ($n = 3$). Additional patient and disease characteristics are shown in **Table 1**.

The median and mean PSA reported prior to SABR were 1.2 ng/mL and 9.0 ng/mL, respectively. The majority of patients, 68% ($n = 45$), underwent SABR with a PSA ≤ 5 . Fifty-three percent of men ($n = 35$) had ≥ 2 metastases. Consolidative SABR was delivered to a total of 134 metastases, with 89 osseous (66%), 40 nodal (30%) and 5 visceral (4%) lesions. The mean GTV was 9.1 cm³ (range 0.1-152.3 cm³). Additional patient characteristics at time of SABR are shown in **Table 2**. SABR was delivered to all lesions with a total dose in the range of 15 Gy to 55 Gy. The majority of lesions (72%) were treated with total doses > 15 Gy with varied fractionation; while 28% of lesions were treated with 15 Gy as a single fraction (**Supplemental Table 1**).

Clinical outcomes:

Median and mean follow-up were 61 and 66 weeks, respectively. Acute toxicities were noted in 46% ($n = 30$) of patients; whereas late toxicities were noted in 27% ($n = 18$) of patients with the most common toxicities being fatigue and hot flashes (**Supplemental Table 2**). In total, grade 1 toxicities were noted in 24 patients and grade 2 toxicities were noted in 7. There were no grade 3 or higher toxicities observed.

The average time to any failure was 35 weeks. Overall, LPFS at 1-year was 92% (**Figure 1A**) with LPFS of 100% for SABR biologically equivalent doses (BED with $\alpha/\beta = 10$) of > 50 . The bPFS and DPFS for the entire

cohort at 1-year were each 69% (**Figure 1B-C**). For all types of failure, men with oligometastatic HSPC had better outcomes than those with CRPC (**Figure 1A-C**). Twenty-two men had a new intervention secondary to progression of their disease post-SABR, with median TTNI not reached overall with our current follow-up of 15-months (**Figure 2A**). Of the 18 men with HSPC who were not treated with adjuvant hormone therapy, 11 (61%) remain free of disease following SABR corresponding to a 1-year ADT-FS of 78% (**Figure 2B**) and a median ADT-FS that was not reached. Of the 17 men with oligometastatic CRPC treated with SABR, 11 had > 50% PSA declines with a 1-year TTNI of 38% (**Figure 2A**) and a median TTNI of 45 weeks. This was congruent with a 1-year bPFS and DPFS of 43% and 33%, respectively (**Figure 1B-C**) in men with CRPC. During the study period, two patients died: one of complications related to their metastatic CRPC and the other from unconfirmed causes not likely related to prostate cancer.

Discussion

This study demonstrates consolidative SABR in men with OPC is well tolerated and appears to have a clinical signal adding to a growing literature of evidence validating its use. We report 1-year LPFS of 92% and bPFS and DPFS of 69% and in those with HSPC not treated with hormone therapy, 61% remain free of disease following SABR. Furthermore, treatment was very well tolerated with 24 patients experiencing grade 1 toxicity, 7 with grade 2 toxicities and no grade 3 or higher toxicities observed.

Perhaps the most notable contributor to the subjective wellbeing of a man with metastatic HSPC is the delaying or deferring of ADT, the side effects of which can significantly hamper a patient's lifestyle and systemic health [11]. With ADT, in addition to decreases in subjective feelings of wellness associated with androgen loss, there has been reported increased risk of osteoporosis, cardiovascular-related mortality and diabetes [26]. Consistent with prior retrospective reports [12,27] and the recently reported randomized phase II STOMP trial [9], our study also suggests that SABR can substantially delay initiation of ADT in men with oligometastatic HSPC for up to a year. Additionally, unlike ADT, SABR is associated with mild side effects without an associated decrement in quality of life measured in both acute and long term follow up [9].

STOMP represents the first published prospective randomized trial of MDT in OPC and validated the efficacy of SABR for MDT by demonstrating a superior ADT-FS (21 months vs 13 months) compared to observation [9,13]. Justification for its advent was supported by an accumulation of retrospective and observational studies reporting durable LPFS, DPFS, and ADT-FS with the use of MDT [17-19,23,24]. The results of our institutional experience are concordant with these reports and in addition add novel information to the already established body of literature. Specifically, the majority of men in this report (53%) had ≥ 2 metastases and thus our study examines a population of OPC with a heavier disease burden than has been explored in the current literature [9,17,19,28]. Rates of two year distant and progression free survival reported in the literature following SABR range from 30-40% and in a subset of those with oligometastatic progression retreated with SABR, the proportion of individuals remaining progression free is as high as 50% at last follow up [28]. Herein we report a one-year DPFS rate of 69%, which compares favorably to the aforementioned studies and therefore, may justify the more aggressive use of MDT in men with several oligometastatic foci.

Thus far the outcomes of interest and treatment intent of SABR in OPC have, for the most part, been aimed at delaying the start of palliative ADT. With growing support for its efficacy in this realm, the aims and goals MDT may shift from a palliative nature to a definitive one by offering these men a second (and in the context of salvage radiation a third) chance for cure. In alignment with the Hellman-Weichselbaum spectrum theory of metastasis, the validation and subsequent adoption of more sensitive imaging modalities such as NaF, Choline C-11, and DCFPyL PET/CT scan can allow for earlier detection of low volume metastatic disease that subsequently that can be treated with local therapy, possibly with definitive intent. Justification for such a paradigm is also supported by retrospective evidence that MDT improves prostate cancer cause specific survival in a propensity matched cohort [29]. Therefore, treatment intensification aimed at simultaneously eradicating other sites of microscopic metastatic disease through the use of concurrent ADT, which is retrospectively associated with improved DPFS when combined with SABR [17], non-castrating antiandrogens such as enzalutamide, chemotherapies such as docetaxel and radiopharmaceuticals may be justified.

In addition to preventing the site-specific complications associated with local progression of metastases, SABR is a complimentary local treatment to maximize durability of systemic approaches. SABR has already been shown to improve disease control in lung cancer patients treated with systemic oncogene-directed therapies through ablation of resistant clones [7,30,31]. The ability to maintain men with CRPC on 2nd-line hormonal agents that are oligoprogressing and delay chemotherapy adds significantly to the clinical options for these patients. Our data showing that SABR can delay distant progression and time to initiation of the next line of therapy (median 42-45 weeks) compares very favorably to any of the approved systemic agents in the CRPC space [32,35]. Finally, the highly conformal nature of SABR limits radiation-associated immunosuppression [33] and may be well suited for concurrent use with immune therapies.

Our study has all the intrinsic limitations of a small retrospective series, heterogeneity, and inclusion of shorter follow-up timeframes in our series limits definitive conclusions following consolidative SABR in OPC. In addition, a compelling weakness is that attempts to define the oligometastatic state by radiographic enumeration of lesions will always be limited by the sensitivity and specificity of the imaging modality and other patient-disease factors that are not well understood. As molecular imaging modalities such as prostate specific membrane antigen (PSMA)-targeted imaging mature, the value of MDT in the oligometastatic state may increase further as patients

previously defined as oligometastatic based on conventional imaging may have additional detectable tumor deposits and require reclassification as polymetastatic [36]. Regardless, our preliminary data suggests SABR in OPC men warrants continued prospective study. In addition to the recent reporting of the Belgian STOMP trial, more definitive conclusions await the completion and reporting of other prospective randomized studies such as our Baltimore ORIOLE trial (NCT02680587).

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Author contributions

Moyer: Data collection or management, Data analysis, Manuscript writing/editing

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Deek: Manuscript wiring/editing

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Disclosure of Potential Conflicts of Interest

Conflict of Interest: The authors declare that they have no conflict of interest

Research involving Human Participants and/or Animals

For this type of study formal consent is not required

Informed Consent

For this type of study formal consent is not required

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Tables

Table 1 Patient and disease characteristics at diagnosis (n=66)

Characteristic	Value
Age at diagnosis (y), median (range)	65 (47-84)
Gleason score, n (%)	
6	7 (10.6)
7	16 (24.2)
8	18 (27.3)
9	22 (33.3)
10	3 (4.5)
PSA	
Initial value (ng/mL), median (range)	8.7 (1.9-2155)
PSA value (ng/mL), n (%)	
0-5	9 (13.6)
>5-10	24 (36.4)
>10-20	9 (13.6)
>20	14 (21.2)
Unknown	10 (15.2)
pT stage, n (%)	
pT1	0 (0.0)
pT2	15 (22.7)
pT3	31 (47.0)
pT4	0 (0.0)
Unknown	3 (4.5)
cT stage, n (%)	
cT1	5 (7.6)
cT2	6 (9.1)
cT3	5 (7.6)
cT4	1 (1.5)
N stage, n (%)	
N0	50 (75.8)
Synchronous N1	16 (24.2)
M stage, n (%)	
M0, Metachronous	41 (62.1)
Synchronous M1	25 (37.9)
Primary treatment, n (%)	
RP	48 (72.7)
RT	17 (25.8)
Post-RP XRT	30 (45.5)
RT and salvage RP	0 (0.0)
Chemotherapy, n (%)	
Prior or concurrent	36 (54.5)
Adjuvant androgen-deprivation therapy, n (%)	
Neoadjuvant	53 (80.3)
Concurrent	53 (80.3)
Adjuvant	48 (72.7)
Sensitivity to hormone therapy, n (%)	
HSPC	49 (74.2)
CRPC	17 (25.8)

Abbreviations: PSA = prostate-specific antigen; RP = radical prostatectomy; RT = radiation therapy; XRT = external beam radiation therapy; HSPC = hormone sensitive prostate cancer; CRPC = castration resistant prostate cancer

Table 2 Patient and disease characteristics at time of stereotactic ablative radiotherapy

Characteristic	Patient (n=66)	Lesion (n=134)
Pre-SABR PSA (ng/mL), median (range) and mean	1.2 (0.01-95.8) 9.0	
PSA (ng/mL), n (%)		
0-5	45 (68.2)	
>5-10	5 (7.6)	
>10-20	7 (10.6)	
>20	9 (13.6)	
Total no. of metastases, no. of patients (%)		
1	31 (47.0)	
2	19 (28.8)	
3	7 (10.6)	
4	4 (6.1)	
5	2 (3.0)	
6	3 (4.5)	
Regions treated, n (%)		
Bone	41 (62.1)	89 (66.4)
Node	24 (36.4)	40 (29.9)
Other	3 (4.5)	5 (3.7)
Lesion gross tumor volume, cm ³ (n=134)		
Mean (Range)		9.09 (0.1-152.3)
Median		2.45

Abbreviations: SABR = stereotactic ablative radiotherapy; PSA = prostate specific antigen

Figure Legend

Figure 1. The effect of stereotactic ablative radiotherapy on clinical progression in men with oligometastatic prostate cancer. (A) Local progression-free survival of the entire cohort (blue), HSPC [34] and CRPC (green). (B) Biochemical progression-free survival of the entire cohort (blue), HSPC [34] and CRPC (green). (C) Distant progression-free survival of the entire cohort (blue), HSPC [34] and CRPC (green).

Figure 2. The effect of stereotactic ablative radiotherapy on delaying subsequent therapy in men with oligometastatic prostate cancer. (A) Time to next therapy of the entire cohort (blue), HSPC [34] and CRPC (green). (B) Time to ADT initiation in men with oligometastatic HSPC.

Figure 1

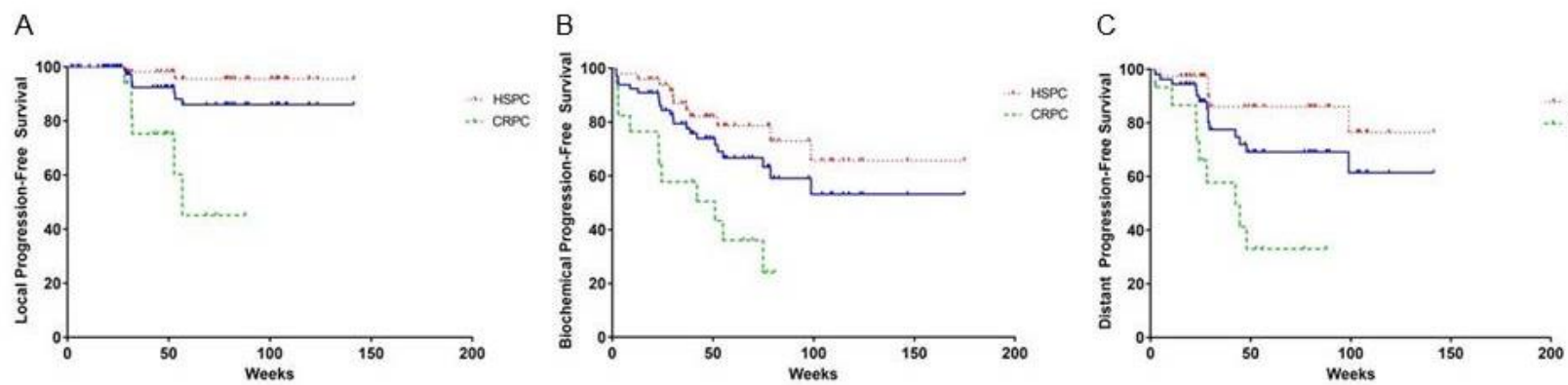
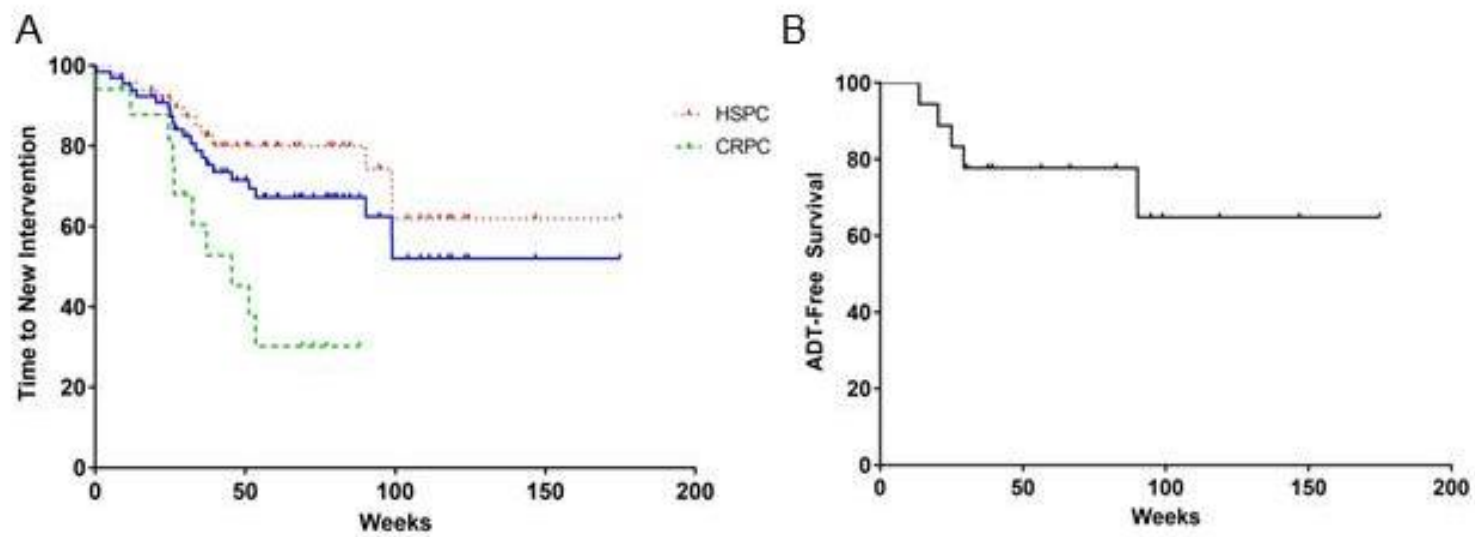


Figure 2



Supplemental Materials

Supplemental Table 1 Specific stereotactic body radiation therapy fractionation patterns by lesion

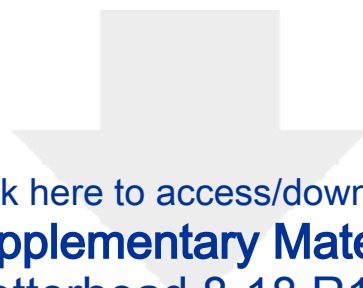
Specific PTV dose (Gy)/no. of fractions*	Lesions treated, n (%) (n=134)	GTV (cm ³), mean (range)
<8/5	22 (16.4)	30.4 (0.5-152.3)
8/5	2 (1.5)	8.8 (3.8-13.9)
9/3	18 (13.4)	6.4 (0.2-19.6)
10/3	18 (13.4)	5.6 (0.1-43.2)
10/5	9 (6.7)	6.4 (0.3-29.8)
11/5	1 (0.7)	Unavailable
12/2	3 (2.2)	27.2 (0.8-53.5)
12/3	6 (4.5)	2.3 (0.3-10.0)
15/1	38 (28.4)	6.3 (0.2-25.3)
16/1	2 (1.5)	1.1 (0.4-1.8)
18/1	15 (11.2)	1.6 (0.1-5.3)

Abbreviations: PTV = planning tumor volume; GTV = gross tumor volume

Supplemental Table 2 Toxicity

Characteristic	Value	
Toxicity presentation, n (%)		
Grade 1	24 (36.3)	
Grade 2	7 (10.6)	
Grade 3	0 (0.0)	
Acute toxicity, n (%)	30 (45.5)	
Late toxicity, n (%)	18 (27.3)	
Toxicity regions, n (%); mean grade (range) and median		
GI	1 (1.5)	2 (2-2), 2
GU	6 (9.1)	1.2 (1-2), 1
MSK	2 (3.0)	1 (1-1), 1
Other	20 (30.3)	1.3 (1-2), 1

Abbreviations: GI = gastrointestinal; GU = genitourinary; MSK = musculoskeletal



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Supplementary Material

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