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Single and Dual Source Strength Focal Boost Planning in Low Dose Rate Prostate Brachytherapy: Feasibility Study

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Corresponding Author:	Anna Kristina Clark, MSc Leeds Teaching Hospitals NHS Trust Leeds, West Yorkshire UNITED KINGDOM
Corresponding Author Secondary Information:	
Corresponding Author's Institution:	Leeds Teaching Hospitals NHS Trust
Corresponding Author's Secondary Institution:	
First Author:	Anna Kristina Clark, MSc
First Author Secondary Information:	
Order of Authors:	Anna Kristina Clark, MSc
	Gavin Wright, MSc
	Josh Mason, PhD
	Oliver Hulson, MBChB
	Ann Henry, MD
	Sree Rodda, MD
	Peter Bownes, MSc
Order of Authors Secondary Information:	
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	seed displacement was similar to the clinical plans. Conclusions Treatment planning using Variseed to produce SSS and DSS focal boost plans is feasible for LDR prostate brachytherapy to achieve a median escalation in sector D90 of 27% without exceeding local urethral and rectal constraints. SSS and DSS plan dosimetric robustness was similar to clinical plan dosimetric robustness.

Single and Dual Source Strength Focal Boost Planning in Low Dose Rate Prostate Brachytherapy: Feasibility Study

A. K. Clark[a*], MSc, G. Wright[a], MSc, J. Mason[b], PhD, O. Hulson[c], MBChB, A. Henry[d], MD, S. Rodda[d], MD, P. Bownes[a], MSc.

[a]Medical Physics, Leeds Cancer Centre, Leeds Teaching Hospitals NHS Trust
[b]Radiation Physics & Radiobiology, Imperial College Healthcare NHS Trust
[c]Radiology, Leeds Cancer Centre, Leeds Teaching Hospitals NHS Trust
[d]Clinical Oncology, Leeds Cancer Centre, Leeds Teaching Hospitals NHS Trust

*Corresponding author. Department of Medical Physics, St James's University Hospital, Leeds LS9 7TF, UK. Tel.:44-113-20-67537.
Name: Anna K Clark
Email: anna.clark11@nhs.net

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Abstract

Introduction: This study investigates the dose escalation to dominant intra-prostatic lesions (DILs) that is achievable using single-source-strength (SSS) and dual-source-strength (DSS) low-dose-rate (LDR) prostate brachytherapy and a sector-based plan approach.

Methods: 20 patients were retrospectively analysed. Image registration and planning were undertaken using Variseed v9.0. SSS and DSS boost plans were produced and compared to clinical plans. Dosimetric robustness to seed displacement for SSS and DSS plans was compared to clinical plans using Monte Carlo simulations.

Results: 14/20 patients had DIL identifiable on MRI. Median increase in sector D90 of 27% ($p<0.0001$) and sector V150 of 31% ($p<0.0001$) was achieved with SSS planning without exceeding local rectum and urethra dose constraints. DSS plans achieved dose distributions not statistically significantly different from the SSS plans with a median of 8 fewer seeds and 2 fewer needles. SSS and DSS plan sensitivity to random seed displacement was similar to the clinical plans.

Conclusions: Treatment planning using Variseed to produce SSS and DSS focal boost plans is feasible for LDR prostate brachytherapy to achieve a median escalation in sector D90 of 27%

without exceeding local urethral and rectal constraints. SSS and DSS plan dosimetric robustness was similar to clinical plan dosimetric robustness.

Introduction

Prostate cancer is the most common cancer in UK men with most presenting with localised disease. Low-dose-rate (LDR) brachytherapy is a treatment option as monotherapy or combined with external beam radiotherapy (EBRT) as a boost treatment in patients with higher risk localised disease [1]. LDR brachytherapy involves permanent implantation of radioactive seeds into the prostate, most commonly using iodine-125 (I-125).

Prostate cancer can be a heterogeneous disease and there is evidence that clinically significant disease spreads from a dominant intra-prostatic lesion (DIL) [1]. Local recurrence can occur after radiation; usually at the same site as the DIL [2,3].

Prostate cancer displays a dose response to radiation, hence escalating the dose to the DIL is expected to improve local control [4]. A randomised trial using EBRT alone to escalate dose to the DIL demonstrated improved 5-year biochemical control with no increase in normal tissue toxicity [5]. In focal boost treatments the therapeutic aim is to deliver the prescription dose to the whole prostate gland and escalate dose to the DIL to improve the tumour control probability (TCP) whilst maintaining organ at risk (OAR) constraints [1,6].

The literature supports escalation of dose using different treatment and DIL localisation techniques. Gaudet et al. [7] treated 120 patients with LDR brachytherapy focal boost with DILs identified by sextant biopsies and increased the mean coverage of the DIL by 150% of the prescription by 9% in comparison to 70 standard plans with no difference in acute and late toxicities at follow-up. Mason et al. [6] compared focal boost optimisation methods for High-dose-rate (HDR) prostate brachytherapy boosting a focal planning target volume (F-

PTV) or sector to 150% of the prescription and maintaining coverage of the whole prostate. Both optimisation methods were achievable without compromising OAR tolerances.

Conventionally, LDR plans use seeds of a single-source-strength (SSS). When escalating dose to focal volumes, increased seed density leads to an increase in number of needles and subsequent prostate trauma [8]. Seed density could be reduced by utilising a mixture of standard source strength and higher source strength (HSS) seeds. Mahdavi et. al. [8] investigated the use of dual-source-strength (DSS) planning for treating focal-only targets to the prescription dose and sparing the rest of the prostate gland and achieved acceptable coverage with approximately half the number of needles and sources compared to SSS plans.

Positional errors in seed placement and the migration of seeds post-implant reduces prostate coverage and increases OAR doses on average [9]. Kaplan et al. found an average radial migration of stranded seeds of 3.7 mm from intended positions [10]. SSS focal boost planning improves plan robustness as a greater number of seeds are used; however, there are necessarily fewer seeds on the opposite side of the prostate to the involved sectors. Random shifts in those individual seeds could cause a significant loss of coverage of the prostate**.** DSS plans are likely to have reduced numbers of seeds with respect to SSS plans, therefore it must be established if the robustness of these plans is reduced and the technique infeasible.

The main outcome of this study was to evaluate the potential for dose escalation to the DIL using LDR prostate brachytherapy with SSS and DSS treatment planning prior to clinical

implementation. Dose-volume histogram (DVH) parameters for targets and OAR were compared between clinical and sector boost plans produced using a SSS and DSS approach to determine the achievable dose escalation to involved sectors without compromising OAR toxicity. Robustness of SSS, DSS, and clinical plans to seed displacement was assessed using an in-house Monte Carlo (MC) simulation.

Methods

Data Preparation

Patients previously treated with 145Gy LDR brachytherapy as a monotherapy and those treated with 110Gy LDR brachytherapy followed by a course of EBRT of 46 Gy in 23 fractions (combined therapy) were included in this retrospective study. These treatment groups were chosen to ensure a range of disease stages. Patients were selected chronologically back in time until 10 monotherapy patients and 10 combined therapy patients were identified. All 110Gy patients and 2 of the 145Gy patients received hormone therapy prior to brachytherapy. Clinical stage ranged from T1c to T3a N0M0, Gleason score ranged from 6 to 9. Presenting PSA had a median of 8.6 $ngml^{-1}$ (0.5 - 48.5 $ngml^{-1}$).

Multiparametric magnetic resonance imaging (mp-MRI) series, trans-rectal ultrasound (TRUS) imaging with prostate capsule, planning target volume (PTV), urethra and rectal wall contours, and the original clinical plan were retrieved. PTV was a 3 mm expansion of the prostate capsule clipped posteriorly at the prostate-rectal interface. All patients were

imaged at Leeds Teaching Hospitals with the same imaging protocols. Patient cases with identifiable DILs progressed to the planning stage.

Mp-MRI series included a T2-weighted fast spin echo (T2W-FSE) scan, diffusion-weighted-imaging (DWI) scan, and dynamic-contrast-enhanced (DCE) scan with a gadolinium-based contrast agent. Prostate capsule and visible DILs were contoured on the T2W-FSE by an experienced consultant radiologist and informed by the DWI and DCE scans. A rigid registration between mp-MRI prostate and TRUS prostate was performed in Variseed, which had no deformable registration solution.

Planning

All plans were produced using the Variseed v9.0 treatment planning system (TPS) and the AAPM TG-43U1S2 calculation algorithm [11]. AgX100 TheraStrand (Theragenics, Georgia, USA) stranded I-125 seeds were used. Clinically delivered plans for which a source strength other than the standard 0.453U seed strength had been used were re-planned with 0.453U seeds to reduce confounding. The aim was to produce plans boosting involved sectors. All plans were reviewed to be clinically acceptable by experienced planners.

For sector planning, prostate volumes were split into 3 sections of equal length - base, mid-gland, and apex - each with 4 transverse sectors for a total of 12 sectors centred on the urethra in one of two orientations demonstrated in figure 1. DIL volume locations after rigid registration informed the selection of sectors to boost; the orientation of the transverse sections was chosen to minimise the number of involved sectors.

SSS plans were produced with 0.453U seeds aiming to maximise the dose escalation to the boost volume (BV) whilst remaining within the local rectum, urethra and target constraints specified in table 1. DSS plans were produced to meet the same aims using 0.453U and 0.682U seeds. A strength of 0.682U was selected for the HSS seeds as this was as close to 150% of the standard source strength as could be ordered. Mahdavi et al. [8] used source strengths of 0.4 U and 0.8-0.9 U for focal DSS plans where only a focal region identified by mp-MRI was treated, however it was decided a lower source strength would be more appropriate for focal boost treatments where the whole prostate was still to receive the minimum peripheral dose in addition to the DIL dose escalation.

Planning techniques followed local planning protocols using stranded sources with one seed strength per needle and avoiding single seeds except at the apex. Sources were nominally constrained to template positions; however small shifts off-template locations were allowed for one or two needles in each plan to optimise positioning. Sources were not placed in directly adjacent template positions except within the BV. HSS needles were manually constrained to pass through the BV. HSS seeds were allowed superiorly and inferiorly to the BV to ensure coverage with only one seed strength per needle.

Local planning aims for standard seed planning are detailed in table 1. For boost plans the prostate D90, V150, and V200 statistics were expected to exceed these limits due to the escalation of dose to the DIL, which is contained within the prostate volume. This was deemed acceptable because of the expected clinical benefit from DIL dose escalation.

As data from monotherapy and combined therapy patients was used, the distribution of individual DVH parameters was not normally distributed; therefore, statistical significance was assessed using the two-tailed Wilcoxon signed-ranks test with a significance level of 5% to compare the distribution of DVH parameters from the SSS and DSS boost cases to the clinical plans and to each other.

Robustness

In-house code previously described by Al-Qaisieh et al. [12] was adapted for this project. Structure sets and planned source positions were exported from Variseed. Random positional shifts were applied to the individual seed coordinates. Dose distribution and DVH parameters were calculated in Matlab by superimposing MC dose distribution data for an AgX100 seed [13]. Resulting dose to the target and OAR volumes was quantified to evaluate the robustness of SSS, DSS, and clinically delivered plans against post-implant seed migration.

Random shifts applied to seed coordinates were based on a Gaussian distribution with a mean of zero and a standard deviation increasing from 2 mm to 5 mm in 1 mm increments as in work by Al-Qaisieh et al. [12]. For each increment 50 random shifts were applied and mean DVH parameters calculated. The MC code was previously validated against TG43 source data as described by Mason et al. [13].

Results

14 out of 20 cases had identifiable lesions on mp-MRI; 7 from each treatment group. A single lesion was identified in 11 cases and 2 lesions identified in 3 cases. 15 lesions were in the mid-gland, 1 in the base, and 1 in the apex.

These 14 patients were rigidly registered. The average mean-distance-to-agreement between the TRUS prostate contour and the registered mp-MRI prostate contour was 5.3 mm (3.87 mm - 7.95 mm). Contoured DIL were used to identify sectors to be included in the BV; the use of sectors mitigates uncertainties in delineation and registration and allows the movement of dose within the boost volume to better spare OAR [6].

Planning

Median 2.5 sectors were involved per patient case with a median BV of 6.22 cm^3 (range: 3.09 cm^3 - 13.71 cm^3) (table 2). This corresponded to a median BV of 25% (range: 8% - 40%) of the total TRUS prostate volume.

The median percentage change in key parameters from the clinical plan for SSS and DSS is detailed in table 3. Figure 2 compares isodoses from a single slice of a single case with the clinically delivered, SSS, and DSS plans.

Statistically significant increases in sector D90, V150, and V200 were obtained with both boost planning methods without compromising prostate and PTV coverage. There was a

statistically significant increase in median rectum D2 cm^3 and urethra D10 from both the SSS and DSS planning methods, but for all cases the local dose constraints were met for rectum and urethra. No statistically significant differences were found when comparing DVH parameters for the sectors, prostate, PTV, and OAR between SSS and DSS plans. The SSS plan total reference air kerma rate (TRAK) increased by median 8% from clinically delivered plans, which was statistically significant. The DSS plan TRAK was not statistically significantly different from the SSS plans.

SSS plans for monotherapy (145 Gy) had a median rectum D2cc increase of 13% whereas for combined therapy (110 Gy) the median increase was 15%. For DSS plans the rectum D2cc mean increase compared to clinical plans for monotherapy was 11% and for combined therapy was 10%. This result was not statistically significant due to the small sample size of each type of therapy.

Robustness

The MC code was validated by assessing the mean percentage change in DVH parameters from those reported by Variseed for the unchanged SSS and DSS boost and clinical plans for each patient case (table 4).

As the standard deviation of random shifts in all directions was increased overall prostate coverage was lost and OAR doses increased (figure 3) for the clinical plans, and SSS and DSS boost plans when compared to the original unchanged plan. After a random shift with a

213 standard deviation of 5 mm prostate D90 was decreased by a mean of 24% for clinical plans

214 and 20% for SSS and DSS boost plans.

Discussion

Focal boost techniques could improve outcomes for patients with localised disease and DIL, however there are no recommendations on the level of dose escalation required or the most appropriate technique [1,14]. The key dosimetric focus of this work was to ascertain the dose escalation achievable using SSS and DSS techniques while not exceeding current clinically implemented OAR constraints.

This sector boost approach proved feasible for a statistically significant escalation in sector V150 of 31% using SSS and 32% using DSS planning. When boosting mp-MRI identified DILs using HDR brachytherapy with an inverse planning optimiser Mason et al. [15] achieved an increase in DIL D90 of 16% and in DIL V150 of 48.6% for DILs with a median volume of 1.9 cm^3. Tissaverasinghe et al. [16] achieved a DIL D90 of 151% of the prescription dose for LDR monotherapy patients where the average BV was 1.9 cm^3. We boosted a larger volume of the prostate (median: 6.22 cm^3) than these cases, and therefore would not be able to achieve as high a boost without exceeding urethra and rectum constraints and risking increased toxicity.

A key clinical impact is that the escalation achieved is comparable to HDR and LDR techniques presented in the literature when the size of the boost volume is considered and was achieved using the current clinical system without significant changes to planning techniques. Consequently, implementation of this technique would not require significant additional training burden.

HDR boost treatments produce fewer severe urethral toxicities than LDR boost treatments and are indicated for more advanced disease [1]. However, there is no current evidence-based recommendation for HDR as monotherapy. LDR monotherapy treatments have the advantage of a single treatment visit and may have less impact on long-term sexual function [1]. This work demonstrates dose escalation feasibility for monotherapy treatments prescribed to 145 Gy and combined therapy treatments prescribed to 110 Gy whilst maintaining dose constraints to rectum and urethra.

The study is limited by having a single consultant radiologist for contouring and a single treatment planner, and consequently does not account for inter-operator variability. However, all plans were validated by two members of physics staff with combined planning experience of 35 years, and a selection of plans were reviewed by a consultant oncologist.

The study has demonstrated that the SSS and DSS plans were similar to clinical plans in dosimetric robustness to random seed migration (figure 3). Consequently, the post-implant dosimetry of the SSS and DSS plans would be expected to be not significantly different to that of the standard clinical plans at our centre. These results support the feasibility of both techniques. Boost plans were less robust than the clinical plans in absolute dose to OARs, however this is due to these plans starting with a higher urethra D10 and rectum D2cc.

A weakness of this dosimetric robustness assessment is that it did not account for the stranded nature of the seeds within each needle, which suggests seed motions would be likely to be systematic within each strand. This could be addressed in a future study by modifying the existing model.

Mahdavi et al. [17] investigated plan robustness to source displacement for DSS plans for focal-only LDR prostate brachytherapy treating a hemi-gland target volume. Random seed displacement was modelled for 50 simulated cases. They found DSS plans were superior in robustness of target volume coverage to the SSS planning technique used clinically at their centre. Our work adds to these findings by applying a similar robustness assessment for focal boost plans based on clinical patients.

There were statistically significantly more needles and seeds used in SSS plans than DSS plans (median[range]: 2[0 – 10] needles, 8[1 – 16] seeds) however in practice on average this might not translate into a practical time saving for every patient. This is clinically significant as achieving the same dosimetric result with a reduction in seeds and needles can result in reduced trauma to the patient and reduced time in theatre, which means the patient can be under general anaesthetic for a shorter period. Additionally reducing the number of seeds and needles can reduce the overall cost of the procedure, which is a compelling advantage considering the current economic climate in healthcare [18]. SSS boost planning would have fewer risks in implementation due to the practical aspects of handling multiple source strength seeds in one treatment.

It could have been expected that the DSS plans would be less robust than the SSS due to the reduced number of seeds and needles used, however the DSS plans were not statistically significantly different to the clinically delivered plans in numbers of seeds and needles, and this combined with the higher strength of the boost seeds resulted in similar robustness.

SSS plans for monotherapy (145 Gy) had a median rectum D2cc increase of 13% whereas for combined therapy (110 Gy) the median increase was 15%. For DSS plans the rectum D2cc median increase compared to clinical plans for monotherapy was 11% and for combined therapy was 10%. This result was not statistically significant due to the small sample size of each type of therapy; however, this observation suggests further work investigating the dosimetry of boost plans for different plan prescriptions could demonstrate the efficacy of one plan type over another with respect to rectum sparing, and lead to improved personalisation of patient treatment.

LDR focal boost techniques are feasible and produce escalations comparable to HDR techniques. Next steps are the clinical implementation of the technique and audit of long-term patient outcomes.

Conclusions

DILs were identifiable in mp-MRI in 70% of cases and informed the involvement of sectors for sector-based planning. A statistically significant median escalation in sector D90 of 27% was achieved using SSS and DSS boost planning methods. Using the DSS planning method this was achieved with a median of 8 fewer seeds and 2 fewer needles. This dose escalation was achieved without exceeding local OAR constraints or loss of prostate coverage. The robustness of SSS and DSS plans were not significantly different to clinical plans.

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Captions to Illustrations

Figure 1: A single TRUS slice from Variseed from the same case demonstrating the two orientations of transverse sectors centred on the urethra to produce 12 sectors in total with subfigures demonstrating the classification of sectors throughout the prostate in each method. The orientation of the transverse sections was chosen to minimise the number of involved sectors.

Figure 2: A single TRUS slice from a 110 Gy patient: (a) transverse sectors at midgland, (b) clinical plan, (c) SSS plan, and (d) DSS plan. Structures are: red = TRUS prostate capsule; light blue = PTV; green = urethra; dark blue = rectal wall; pink = rigidly registered DIL. Isodoses are: yellow = 100%; red = 150%; burgundy = 200%. Needle paths are shown in yellow with 0.453 U seeds filled in green and 0.682 U seeds in light pink.

Figure 3: Mean percentage change in DVH stats for SSS**,** DSS, and clinically delivered plans calculated using the MC code for increasing standard deviations of random shifts (σ_{shift}). Each case was recalculated 50 times. Error bars represent 95% confidence intervals.

Figure 1

Click here to access/download;Figure;Figure_1.tif

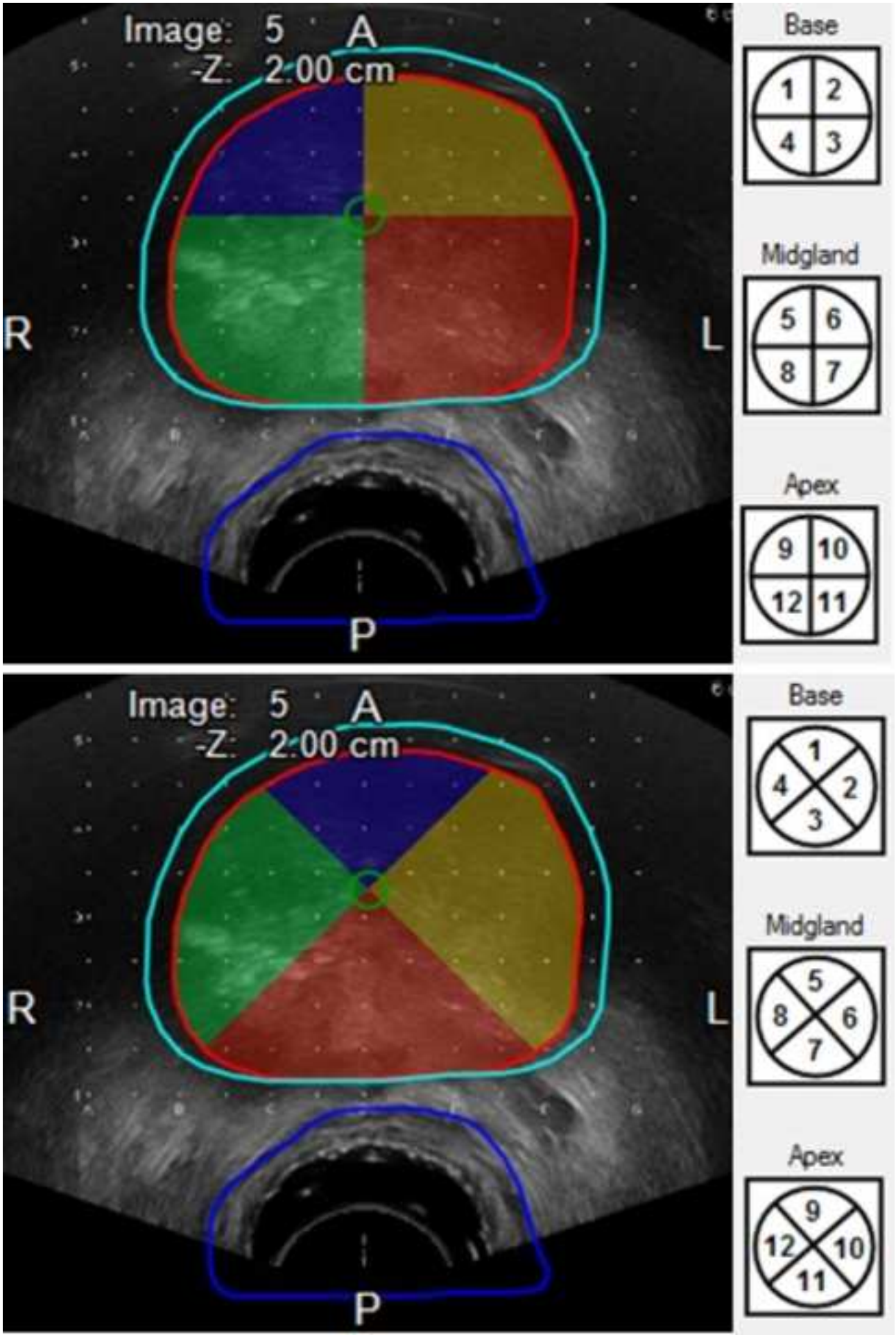

Figure 2

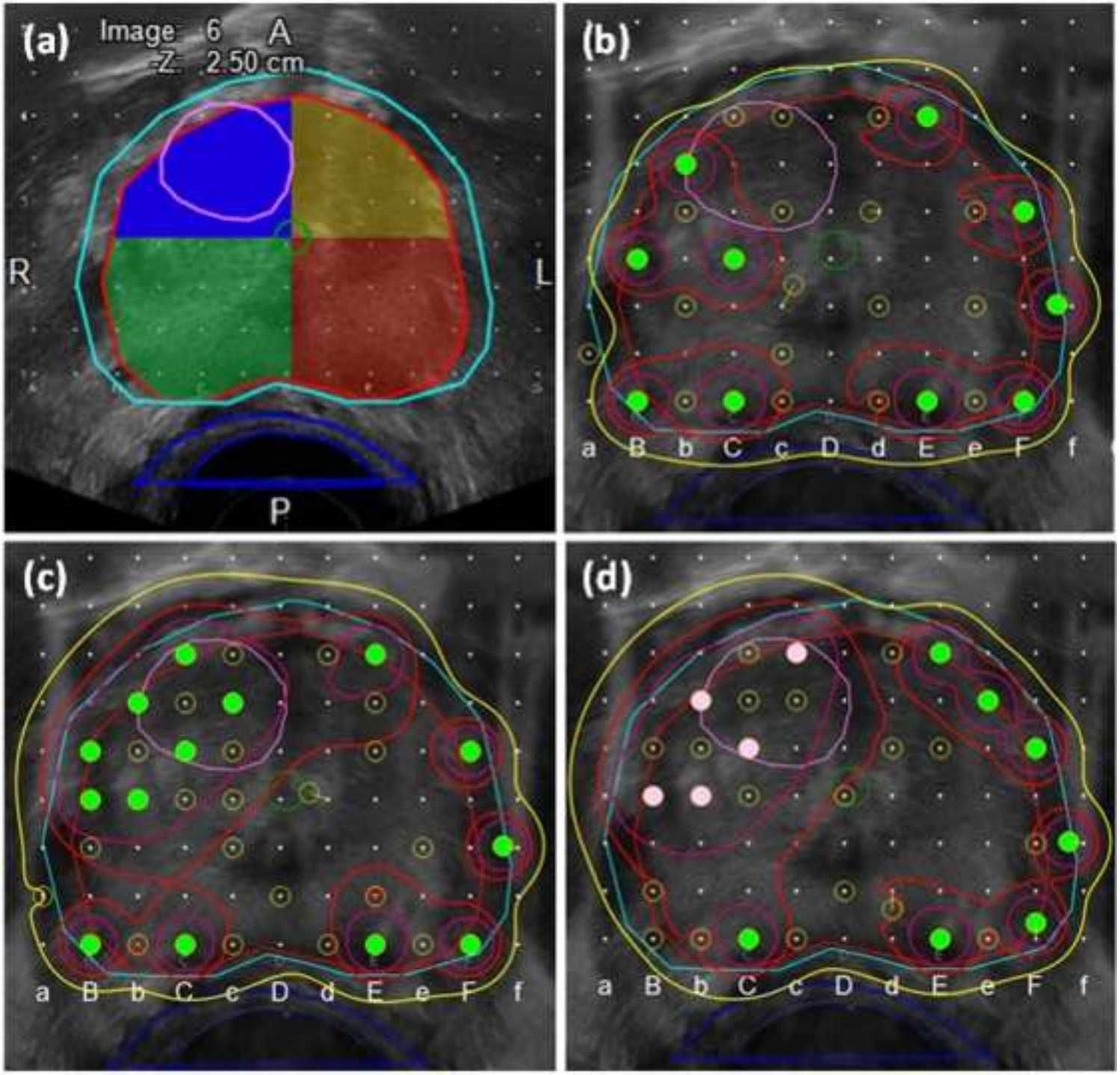

Figure 3

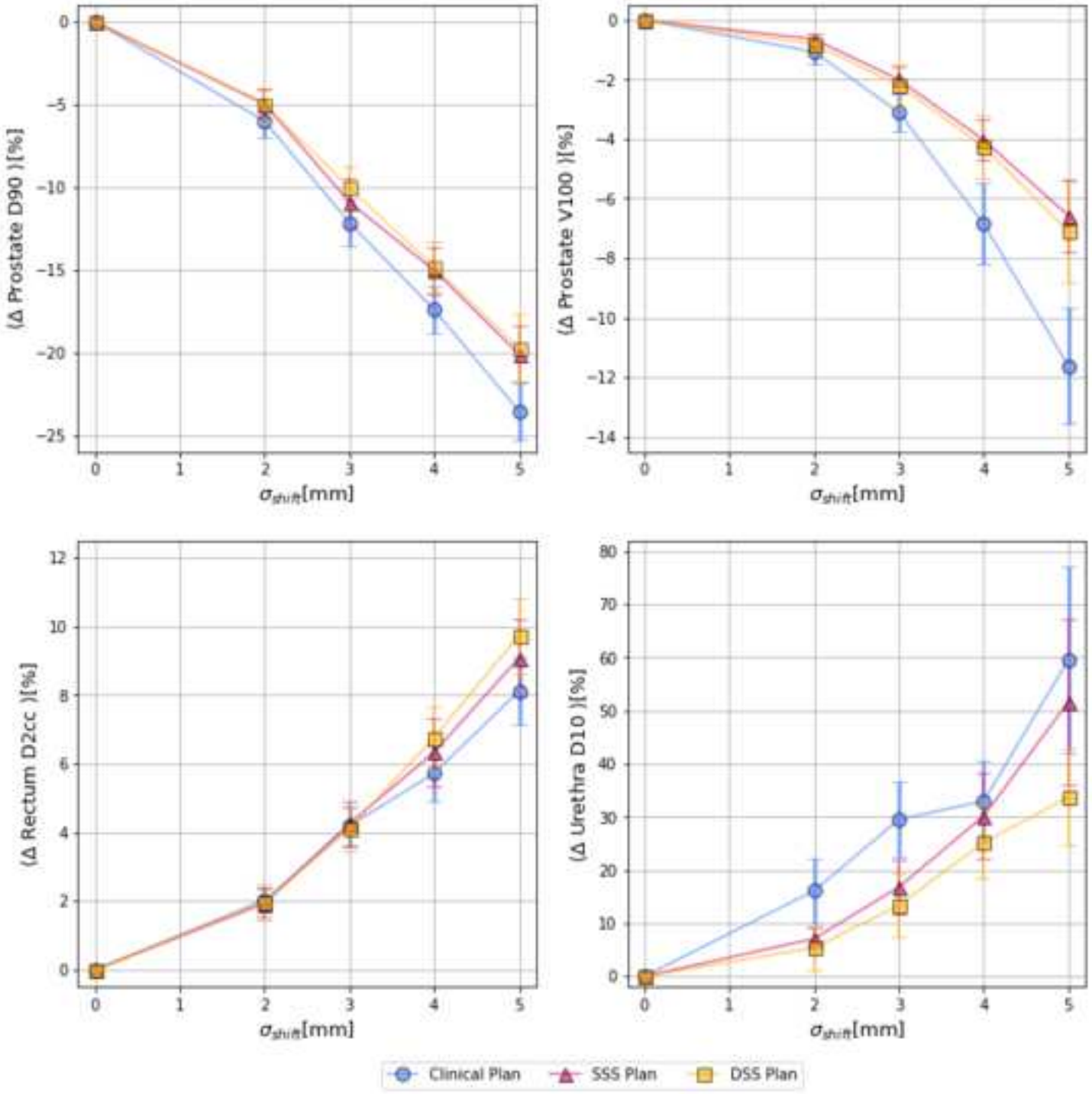

Table 1: Local planning aims for non-focal boost 145 Gy and 110 Gy LDR prostate brachytherapy.

Volume	Parameter	Monotherapy Aim 145 Gy to 100%	Combined Therapy Aim 110 Gy to 100%
Prostate	V100%	> 99.8%	> 99.8%
	V150%	55% ≤ V150 ≤ 60%	55% ≤ V150 ≤ 60%
	V200%	≤ 22%	≤ 22%
	D90 (Gy)	185Gy ≤ D90 ≤ 195Gy	140Gy ≤ D90 ≤ 148Gy
PTV	V100%	> 95%	> 95%
Rectum	D2.0cm^3 (Gy)	≤ 145Gy	≤ 110Gy
	D0.1cm^3 (Gy)	≤ 200Gy	≤ 150Gy
Urethra	D10%	≤ 165%	≤ 165%
	D30%	≤ 150%	≤ 150%

Abbreviations: Vn% = percentage of the target receiving n% of the prescription dose; Dn% = minimum dose received by n% of the target; Dncm^3 = minimum dose received by n cm^3 of the target.

Table 2: Treatment prescriptions and volumes for patients included in the planning study.

Patient ID	Prescription (Gy)	Prostate Volume (TRUS) (cm^3)	No. Involved Sectors	Total Sector Boost Volume (cm^3)
1	**145**	36.95	1	3.09
2	**145**	25.85	3	7.00
3	**145**	40.26	2	9.23
5	**145**	31.62	2	6.43
6	**145**	42.59	4	13.71
7	**145**	22.71	2	6.00
9	**145**	34.05	1	4.66
11	**110**	33.78	3	9.84
12	**110**	19.66	3	5.75
13	**110**	37.19	2	8.35
15	**110**	11.62	3	3.77
16	**110**	41.50	1	4.47
17	**110**	24.33	3	3.95
19	**110**	29.91	4	11.95

Table 3

Table 3: Median and range for DVH values and deliverability parameters for 145 Gy and 110 Gy clinical plans and the median change from clinical plans for the SSS boost, DSS boost, and OAR sparing boost plans. P-values obtained using the two-tailed Wilcoxon Signed-Rank Test. TRAK = total reference air kerma rate.

Parameter	Clinical Plan Median [range] (n=14)		SSS Median Change from Clinical [range] (n=14)	DSS Median Change from Clinical [range] (n=14)
	145 Gy (n=7)	110 Gy (n=7)		
Sector D90(Gy)	192.79 [183.47 - 204.39]	144.38 [128.82-156.46]	27% [12%-49%] p=0.0001	27% [4%-52%] p=0.0001
Sector V150(%)	72.08 [39.25-79.70]	63.85 [31.98-77.85]	31% [18%-58%] p=0.0001	32% [10%-56%] p=0.0001
Sector V200(%)	16.20 [10.54-29.52]	25.66 [12.54-32.77]	46% [26%-74%] p=0.0001	44% [24%-75%] p=0.0001
Prostate D90(Gy)	186.32 [185.15-188.72]	140.52 [137.85-143.98]	6% [-1%-12%] p=0.0004	5% [1%-14%] p=0.0001
PTV V100(%)	98.05 [95.97-98.67]	98.13 [95.78-98.81]	1% [0%-3%] p=0.002	1% [-1%-3%] p=0.003
Rectum $D2cm^3$ (Gy)	101.24 [90.37-107.09]	71.00 [67.02-90.09]	14% [-6%-29%] p=0.002	10% [-6%-34%] p=0.003
Urethra D10(%)	149.61 [141.82-157.48]	144.33 [131.44-161.97]	12% [-4%-19%] p=0.0004	12% [-5%-24%] p=0.0006
TRAK (μGym^2h^{-1})	36.693 [27.633-42.582]	27.180 [17.214-34.428]	8% [1%-16%] p=0.0001	11% [1%-19%] p=0.0001
Seeds	81 [61-94]	60 [38-76]	5[1-13] p=0.0001	-3[-9-4] p=0.01
Needles	26 [21-33]	23 [18-29]	5[0-13] p=0.0001	2[-3-7] p=0.002

Seed Density (cm^{-3})	2.38 [2.21-2.69]	2.01 [1.83-3.27]	8%[1%-16%] p=0.2	-5%[-10%-6%] p=0.02

Table 4

Table 4: Mean percentage change from TPS for selected DVH statistics when recalculated using the MC method for the SSS boost, DSS boost, and clinical plans for all patients. SD = standard deviation.

Parameter	**Prostate D90(Gy)**	**Prostate V100(%)**	**Urethra D10(%)**	**Rectum D2cm^3 (Gy)**
Mean Percentage Change (%) [SD]	-1.13 [2.32]	-0.09 [0.33]	2.73 [2.70]	1.40 [2.50]

Response to Reviewers - JRP-D-23-00002
Single and Dual Source Strength Focal Boost Planning in Low Dose Rate Prostate Brachytherapy: Feasibility Study

The following contains a summary of the revisions made in response to reviewer comments, followed by a list of all changes made to the manuscript.

Reviewer 1 comments:

"There needs to be more information to familiarise the readers with the idea of using LDR-brachy for prostate cancer, a line or two would suffice."

1. *The introduction from lines 27-43 has been edited to provide a more robust introduction to LDR-brachy for prostate cancer.*

"Both HDR and LDR have advantages over the other and I think it would be better if you highlight this and the impact it could have for other departments utilising LDR."

2. *A full comparison of HDR and LDR techniques is beyond the scope of this paper. However, I see the advantages of including this. A short comparison is now included in the discussion. See lines 238-244.*

"Was the main drive of your feasibility study to focus on improving treatments for multi-focal disease?"

3. *The main aim of the study is now clearly stated in lines 72-74. An additional comment on the aim of the work is now presented in the discussion lines 217-221.*

"Lines 76-78 - clumsy statement, try re-phrasing."

4. *This statement has been rephrased for clarity. See lines 84-86.*

"I think you should have a clear objective at the end of the introduction, you state what was compared (more like a retrospective study) but try to make it an objective (which is a core concept of feasibility studies)."

5. *See response 3.*

"Lines 114-115 - within your methodology be specific about what OARs you are referring to."

6. *OARs now specifically stated. See line 123.*

"The discussion is well considered from a technical point of view - I think it would be strengthened by adding some discussion on the overall clinical impact of dose escalation - e.g. could LDR play a bigger role for some patients who would traditionally be put forward for HDR? HDR generally have fewer acute toxicities than LDR but LDR has the advantage of a single implant - some consideration of these points would be good."

7. *More clinically focused discussion points have been added; this includes clearer statements on why the work is clinically relevant, and some comparison of HDR and LDR techniques. See lines 217-221, 233-234, 238-242.*

"Also, what is the next steps/actions as a result of this feasibility study?"

8. *A statement of the next steps, which are clinical implementation and audit of patient outcomes, has been added to the end of the discussion. See lines 295-297.*

"All your tables have Table 1 as the caption?"

9. *Table captions error corrected.*

Reviewer 2 comments:

"This is an interesting manuscript that should be of value and interest to JRP readers. My recommendation here would be a minor revision as I feel that the work would really benefit from some more specific details about the clinical implications of this work and how this can be taken forward."

10. *See responses 7 and 8.*

"The authors should also check the table captions."

11. *See response 9.*

Full List of Changes

Lines 27-42: Introduction edited to address comments requesting a more robust introduction to LDR-brachy for prostate cancer.

Line 51: Shortened statement for coherence.

Line 56: New acronym added.

Lines 72-74: This line added to address reviewer 1 request for explanation of the main drive of the study and to give a clear objective at the end of the introduction.

Lines 84-86: Rephrased per reviewer 1 to be a clearer statement.

Lines 93-94: Changed to define acronyms lost in earlier changes to introduction.

Line 123: Clarified OARs in methodology per reviewer 1.

Line 180: Edited wording to improve clarity.

Lines 217-221: Edited and expanded to improve demonstration of clinical benefit per both reviewers.

Lines 223-234: Added to improve demonstration of clinical benefit per both reviewers.

Lines 238-242: Added to address reviewer 1 request for comparison of advantages and disadvantages of LDR and HDR prostate brachytherapy.

Line 264: Edited wording to improve clarity.

Line 269: Removed statement on TRAK to focus on other important outcomes. This result is still presented in table 3.

Lines 295-297: Added to address comments from both reviewers about clinical impact and next steps.

Lines 302-303: Reworded for clarity.

Line 309: References have been added and removed where required as a result of the revisions.