

Santa Maria della Misericordia di Udine



[®] Image-guided SBRT of the prostate, 42 Gy in 7 fractions, for localized disease: dosimetric report of a phase-II study

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Introduction

We present our experience of the institutional prostate SBRT phase-II study, which has involved 40 patients, with a focus on treatment planning dosimetry and compliance with the study protocol.

Materials and Methods

Since 2012, 40 patients at low/intermediate risk for localized prostate cancer have been planned for SBRT. Fraction size is 6 Gy for 7 fractions scheduled to be delivered twice a week for a total dose of 42 Gy. For planning the GTV includes prostate with the 1/3 proximal seminal vesicles without margin; a margin of 3 mm in all directions around the GTV is used to define the PTV, as the SBRT protocol is based on 3 IGRT intra-prostatic fiducial markers with daily online checks by CBCT. Treatment is delivered with a VMAT technique, with 2 arcs using 6MV photons from a Varian 2300 iX linac. Planning was performed with Eclipse 10.0 TPS with the AAA algorithm by 3 different planners. Dose prescription is the average dose to PTV with the request V95% \geq 95%.

Results

Main dosimetric results of the first 40 clinical plans are presented in the following table together with their compliance to the protocol. The PTV is covered by the 95% isodose for all patients, and only the constraint for the PTV near-minimum dose $D_{98\%}$ is not met in 2 cases. As for the OARs, most dosimetric parameters are well within the protocol constraints, with the notable exception of the maximal doses of rectum and bladder ($D_{1\%}$, equal to 95% and 100% of the prescription dose, respectively) for which the constraints are exceeded in about 20% of cases. The mean value of the CN is 0.90 with a coefficient of variation (CV) of 5.5%. The mean value of total MU's corresponds to a delivery of 3.3 min at a doserate of 600 MU/min.



Figure 1. Example of PTV coverage on an axial (a), coronal (b) and sagittal (c) slice. The 95% isodose is shown in blue.

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	Structure		mean	CV	Constraint	deviations
	PTV	D_{median}	100.0%	0.8%		
	Vol=115.6 cc	V _{95%}	97.4%	1.2%	≥ 95%	0
	(CV=26.9%)	D _{98%}	94.7%	1.3%	≥ 93%	2
		D _{2%}	104.9%	1.4%	≤ 108%	0
	Rectum	D _{1%}	37.9 Gy	18.1%	≤ 40 Gy	9
	Vol=57.8 cc	V _{37Gy}	3.2%	47.5%	≤ 5%	3
	(CV=28.0%)	V _{32Gy}	7.7%	31.3%	≤ 10%	2
		V _{20Gy}	30.6%	11.2%	≤ 35%	0
	Bladder	D _{1%}	40.7 Gy	3.8%	≤ 42 Gy	7
	Vol=139.7 cc	V _{38Gy}	5.4%	122%	≤ 13%	3
	(CV=52.2%)	V _{33Gy}	8.2%	52.2%	≤ 30%	2
		V _{21Gy}	22.4%	40.5%	≤ 40%	0
	Rt Femoral Head	D_{mean}	6.6 Gy	28.4%	≤ 10 Gy	3
		D _{1%}	13.5 Gy	25.9%	≤ 20 Gy	0
	Lt Femoral Head	D_{mean}	6.5 Gy	30.7%	≤ 10 Gy	1
		D _{1%}	13.6 Gy	24.2%	≤ 20 Gy	0
	Penile Bulb	D _{mean}	10.2 Gy	53.3%	≤ 20 Gy	2
	CN95%)	0.90	5.5%	≥ 0.80	0
	MU		1956	14.9%		

Table. Dosimetric results and number of patients with minor deviations in the 40 patient sample.

Finally, a significant correlation between van't Riet $CN_{95\%}$ and the number of MUs was observed (r=0,61; p<0,001) as shown in Figure 3. This means that for most of the plans the degree of conformality increases ($CN_{95\%}$ going to 1) at the expense of total MU's.

MU vs CN_{95%}

The DVH constraints for OAR's have been derived from literature and local experience. Constraints for acute urinary toxicity have been recently introduced too after the paper of Cozzarini et al [1]. In addition, the plan quality have been evaluated by van't Riet [2] dose conformation Number (CN) and by the intermediate dose spillage (Spill50%), defined as the ratio of the volume of 50% isodose to the volume of the PTV.

Dosimetric analysis focused on PTV coverage and OAR's sparing based on key DVH parameters corresponding to protocol constraints.



MAT PLAN EVALUATION ractionated radiotherapy for prostate canc rostate + Seminal Vescicles = 6 Gy x 7 fx= 42 Gy Dose prescription: average dose to PTV Result Comments Objective D_{median} (%) V95% > 95% PTV V_{105%} < 5% /ol (cc)= D_{98%} > 94% D_{2%} < 108% V20 < 35% V₃₂ < 10% V37 < 5% D1% < 40Gy D_{mean} < 18Gy Overlap rectum-PTV 90% < Dmean < 95% V21 < 40% cute Urinary Toxi V₃₃ < 30% Van < 56 cc: V₃₈< 13% V43 < 5cc: _ D1% < 40 Gy At least V40 < 10%:_ Dmean < 14 Gy Dmean < 10 Gy Rt Femoral Head D_{1%} < 18 Gy D_{mean} < 10 Gy Lt Femoral Head D1% < 18 Gy Penile Bulk Dmean < 10 Gy [V_{95%} ^{PTV}(cc)]² / [V_{95%} ^{Body}(cc)· V^{PTV}(e CN95% > 0.90 V50 Body(cc) / VPTV(cc) SPILL50%

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Figure 3. Total MU's as a function of Van't Riet CN.

Conclusions

Analysis has shown a good compliance with most of the protocol constraints for PTV and OAR's, except for the maximal doses $D_{1\%}$ of rectum and bladder when these organs are significantly included in the PTV. As for the measure of plan quality, the average value of van't Riet CN is well above the 0.80 threshold, while its small CV% suggests a consistent application of the planning protocol among the different planners involved in the study.

References

V. Carillo, C. Cozzarini, T. Rancati *et al.* Relationships between bladder dosevolume/surface histograms and acute urinary toxicity after radiotherapy for prostate cancer. *Int J Radiat Oncol Biol Phys,* April 2014, Vol 111 (1):100–105.

Figure 2. a) Example of cumulative DVH for PTV and OAR's;

b) Worksheet for plan evaluation and quality control.

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