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BOLOGNA, 27-29 OTTOBRE 2023

PALAZZO DEI CONGRESSI

Radioterapia Oncologica: l'evoluzione al servizio dei pazienti





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STEREOTACTIC BODY RADIATION THERAPY (SBRT) WITH MRI-DEFINED FOCAL SIMULTANEOUS INTEGRATED BOOST FOR PATIENTS WITH LOCALIZED PROSTATE CANCER: ACUTE TOXICITY AND DOSIMETRY RESULTS FROM A PROSPECTIVE OBSERVATIONAL STUDY

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CONFLICT OF INTEREST

Nothing to declare

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PROSTATE SBRT

✓ Stereotactic body radiotherapy (SBRT) represents an effective curative option for localized prostate cancer

Van As, ASTRO, 2023

✓ After prostate SBRT, local recurrence typically occurs in the pretreatment multiparametric magnetic resonance imaging (mpMRI) region of a PI-RADS 4 or 5 dominant intra-prostatic lesion (DIL)

Gorovets, EurUrol Oncol. 2022

✓ The addition of a focal boost to the DIL is an emerging strategy to potentially improve tumor control in patients with organ-confined prostate cancer

Kerkmeijer, J Clin Oncol. 2021 Draulans, Radiother Oncol. 2020

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STUDY DESIGN & AIM

- ✓ IRB-approved prospective observational study of 5-fraction prostate SBRT with a SIB to the DIL, starting in January 2022
- ✓ Subset data cut off December 2022, statistical analisys October 2023



- \checkmark Assess the early toxicity and dosimetry results of prostate SBRT-SIB_{DIL}
- Results, in terms of toxicity and dosimetry, were compared with a group of patients treated with prostate SBRT without boost in the same period



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PATIENT WORKFLOW



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TREATMENT PLANNING



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- ✓ DILs were defined using mpMRI T2-weighted, diffusion and perfusion images
- ✓ The prescribed dose was 36.25 Gy to the prostate with an isotoxic SIB_{DIL} of 40 Gy (dose range 100%-120%)



AIRO2023 RESULTS (1)

- ✓ 18 patients treated with prostate SBRT from January 2022 to December 2022 with mpMRI-defined DIL received SBRT-SIB_{DIL}
- ✓ 83% had cT2 disease
- ✓ Median iPSA 7 ng/ml (range 3.6-19.2)
- ✓ Baseline IPSS score mild/moderate in 61%/39%
- ✓ Most patients had either 3+4 (39%) or 4+3 (33%) Gleason

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The majority of DILs were in the peripheral zone (89%)



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RESULTS (2)

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- ✓ At a median follow up of 16.4 months (range 9.8-20.1), no G≥3 toxicity was observed
- ✓ Cumulative acute G1-2 GU and GI toxicity rates of 44.4% and 16.7%, respectively
- ✓ All patients experienced biochemical response

Table 1. SBRT-SIB treatment plan analysis for major organs at risk (OARs)			
	Dosimetric parameters	Objectives	Results (mean \pm SD)
	V40Gy	<2cc	$0.64 \text{ cc} \pm 0.97 \text{ cc}$
Bladder	V37Gy	<10cc (optimal 5 cc)	$6.15 \text{ cc} \pm 3.82 \text{ cc}$
	V18.1Gy	<40%	$17.94\% \pm 8.49\%$
	V40Gy	<1cc	$0.03 \text{ cc} \pm 0.10 \text{ cc}$
Rectum	V36Gy	<2cc (optimal 1 cc)	$0.35 \text{ cc} \pm 0.68 \text{ cc}$
	V24Gy	<50%	$11.09\% \pm 10.17\%$





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RESULTS (3)



- ✓ In comparison with the non-boost SBRT group (25 patients) treated in the same period, rectal V40, 36 and 24Gy, and bladder V40, 37 and V18.1Gy did not significantly differ (all p > 0.05)
- ✓ No differences were found between groups in terms of acute G2 GU (16.7% boost vs 20.8% no-boost; p = 0.74) and GI toxicity (5.6% boost vs 4.2% no-boost; p =0.83).





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CONCLUSIONS

- ✓ Our results demonstrate that treating PCa with five-fractions mpMRI-defined SBRT-SIB_{DIL} is safe and effective, with excellent adherence to the planning protocol
- ✓ Longer follow-up is needed to evaluate tumor control and late toxicity. In this respect, further enrollment of patients is ongoing





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THANK YOU FOR THE ATTENTION

QUESTIONS?



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