



XXXII CONGRESSO NAZIONALE AIRO
XXXIII CONGRESSO NAZIONALE AIRB
XII CONGRESSO NAZIONALE AIRO GIOVANI

AIRO2022

Radioterapia di precisione per un'oncologia innovativa e sostenibile

BOLOGNA, 25-27 NOVEMBRE
PALAZZO DEI CONGRESSI



Associazione Italiana
Radioterapia e Oncologia clinica



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AIRO



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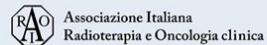
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Post-Prostatectomy Ablative Radiation Therapy (POPART): a Multicentric Prospective Italian Trial

ClinicalTrials.gov Identifier NCT04831970

Dr.ssa Raffaella Lucchini





DICHIARAZIONE

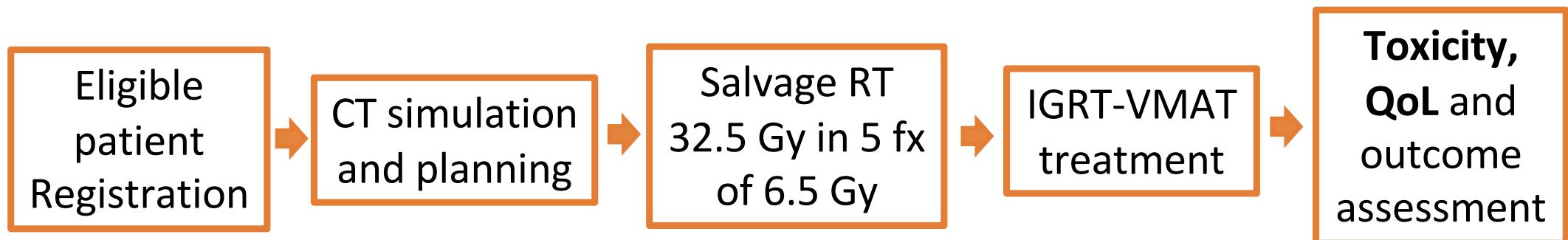
Relatore: Raffaella Lucchini

Come da nuova regolamentazione della Commissione Nazionale per la Formazione Continua del Ministero della Salute, è richiesta la trasparenza delle fonti di finanziamento e dei rapporti con soggetti portatori di interessi commerciali in campo sanitario.

- Posizione di dipendente in aziende con interessi commerciali in campo sanitario (**Niente da dichiarare**)
- Consulenza ad aziende con interessi commerciali in campo sanitario (**Niente da dichiarare**)
- Fondi per la ricerca da aziende con interessi commerciali in campo sanitario (**Niente da dichiarare**)
- Partecipazione ad Advisory Board (**Niente da dichiarare**)
- Titolarità di brevetti in compartecipazione ad aziende con interessi commerciali in campo sanitario (**Niente da dichiarare**)
- Partecipazioni azionarie in aziende con interessi commerciali in campo sanitario (**Niente da dichiarare**)



Multicentric Trial



Sistema Socio Sanitario



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Inclusion Criteria

- ❖ Prostate adenocarcinoma treated with radical prostatectomy
- ❖ Post–prostatectomy PSA of ≥ 0.1 - < 2 ng/ml and/or local relapse
- ❖ No distant metastases at **PSMA PET CT** within 60 days prior to registration
- ❖ Androgen deprivation therapy allowed



Patients and Disease Characteristics

Age

Median 72 [range 55-82]

PSA pre-prostatectomy (ng/ml)

Median 6,04 [range 3,30 -17,25]

Gleason score

6 (3+3)	4 (13%)
7 (3+4)	12 (40%)
7 (4+3)	11 (37%)
8 (4+4)	2 (6,5%)
9 (4+5)	1 (3,5%)

ISUP Grade Group

1	4 (13%)
2	12 (40%)
3	11 (37%)
4	2 (6.5%)
5	1 (3.5%)

Pathological T stage

pT2	19 (64%)
pT3a	8 (26%)
pT3b	3 (10%)

Positive Margins

R1	11 (37%)
R1 and positive apex	8 (26%)

Time from prostatectomy (months)

Median 54,5 [range 7-155]

Clinical relapse

Yes	8 (26%)
No	22 (74%)

PSA pre-SBRT (ng/ml)

Median 0.30 [range 0.18-1.89]

ADT

Yes	4 (13%)
No	26 (87%)

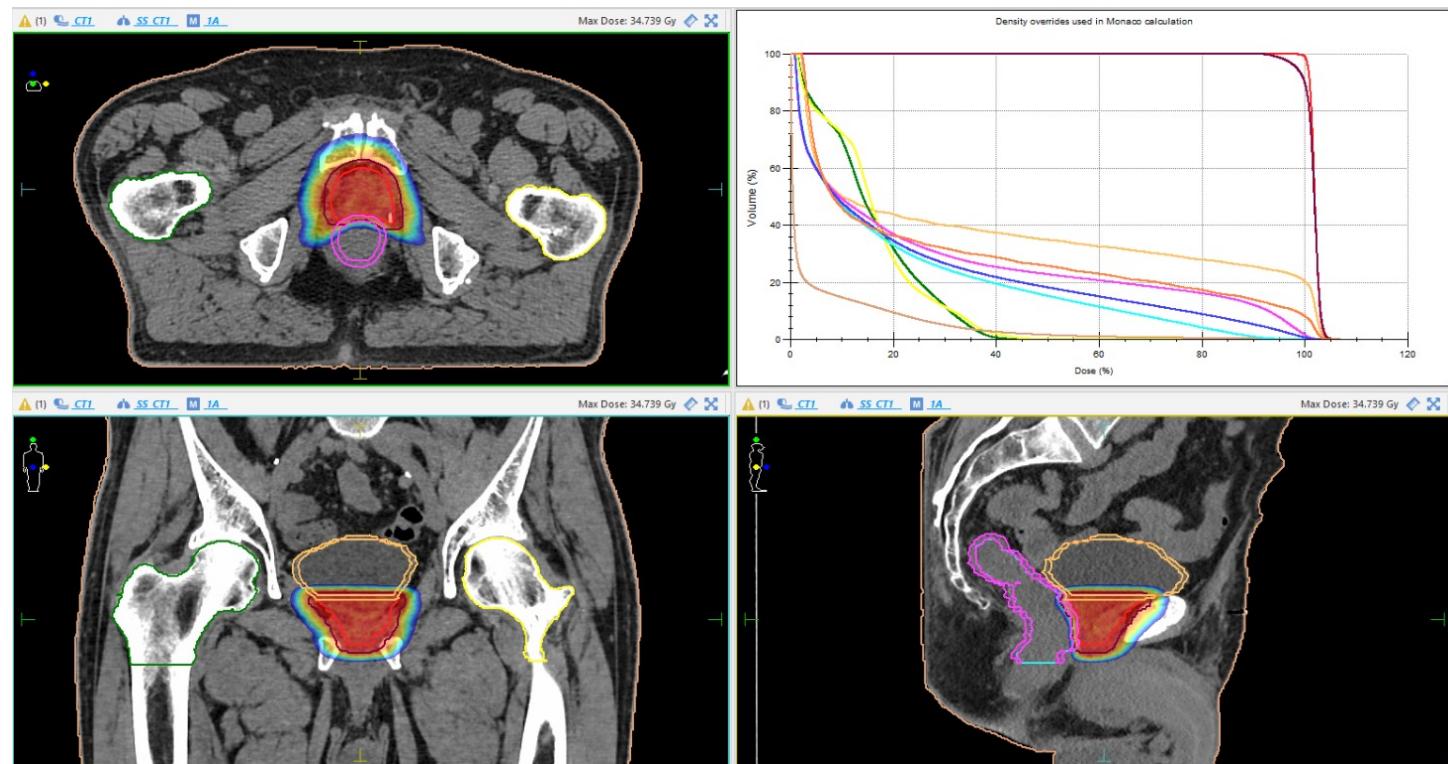


Anatomical reproducibility

- ❖ Organ motion mitigation was obtained before simulation and each treatment fraction with:
 - A rectal microenema
 - 500 ml of still water
- ❖ CTV was delineated according to the GFRU guideline

Treatment Planning

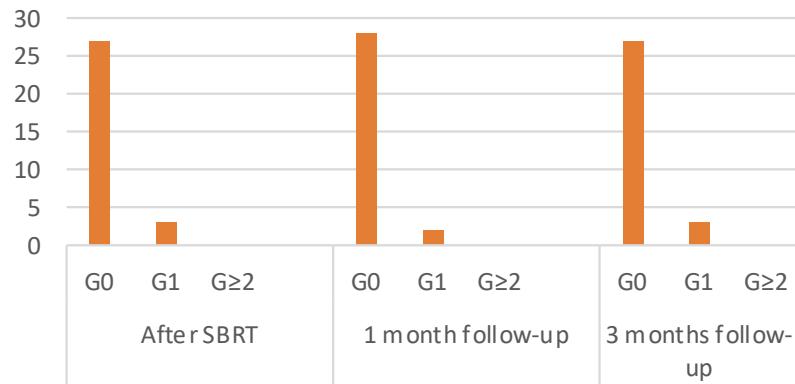
- ❖ 32.5 Gy in 5 fractions (EQD_{2[1.5]}= 74.2 Gy)
- ❖ Treatment was delivered on Linac platform with a Volumetric Modulated Arc Therapy (VMAT)
- ❖ Median CTV (cc) 29,39 [range 4.40-149.00]
- ❖ Median PTV (cc) 72 [range 14.8-250.2]



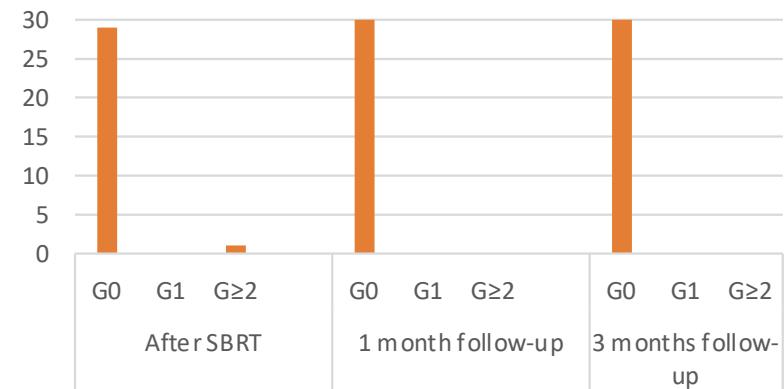


Early Toxicity (CTCAE v.5)

Genitourinary (GU) toxicity



Gastrointestinal (GI) toxicity





QoL (EPIC-CP) and Clinical Outcomes

	Baseline	Mean \pm SD score	
		End of treatment	3 months
Urinary incontinence	1.6 \pm 0.1	1.8 \pm 0.1	1.6 \pm 0.1
Urinary irritation/obstruction	1.2 \pm 0.1	1.8 \pm 0.2	1.6 \pm 0.1
Bowel symptoms	0.7 \pm 0.1	1.8 \pm 0.2	0.8 \pm 0.1
Sexual dysfunctions	5.3 \pm 0.2	5.3 \pm 0.2	5.8 \pm 0.2
Hormonal symptoms	1.1 \pm 0.1	1.3 \pm 0.1	1.6 \pm 0.1

Three months after SBRT all but two patients, who progressed distantly, were found biochemically controlled with a median post-treatment PSA level of 0.07 ng/ml (range 0-0.48 ng/ml)



Take Home Message

- ❖ Post-prostatectomy SBRT for biochemical and/or clinical relapse is a convenient and safe treatment
- ❖ No increase in short term toxicity nor significant decline in QoL was observed
- ❖ Long term results are needed to confirm these findings

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