

Salvage radiotherapy after radical prostatectomy: Preserving patient quality of life with modern treatment techniques

Martin Korzeniowski, MD

Queen's University, Kingston, ON, Canada

Cite as: *Can Urol Assoc J* 2018;12(2):50-1 <http://dx.doi.org/10.5489/cuaj.5112>

Published online December 14, 2017

See related article on page 45

In this issue of *CUAJ*, Ajib et al present data showcasing oncological and functional outcomes after salvage radiation (SRT) for biochemical failure for prostate cancer treated with robot-assisted radical prostatectomy (RARP). The article is particularly interesting because it involves a cohort of men treated with modern radiotherapy and surgical techniques. The disciplines of urologic and radiation oncology have each benefited from technological innovations, and this has contributed to the improved functional outcomes seen in this study. Advances in surgical technique with the advent of RARP have shortened hospital stays, decreased perioperative complications, and improved urinary function. Similarly, radiation oncology has seen improvements in treatment planning and delivery that allow highly conformal treatments to be delivered with reduced genitourinary and gastrointestinal toxicity.¹

In their paper, Ajib et al report on the clinical outcomes for SRT after RARP from a tertiary academic centre in Canada. A prospectively collected database included clinically relevant outcomes for urinary and sexual function. Validated measures, such as the International Prostate Symptom Score (IPSS) and Sexual Health Inventory for Men (SHIM), as well as continence pad usage, were used to demonstrate the safety and tolerability of SRT following RARP from a patient-reported outcome perspective.

While followup is limited, the early oncological outcomes are encouraging, with good levels of prostate-specific antigen (PSA) control despite a mean PSA of 0.5 ng/ml at the time of treatment. Men with extracapsular extension or positive surgical margins had excellent biochemical control (PSA <0.2) at 24 months (80% and 81%, respectively) while those with seminal vesicle invasion did somewhat less well, as expected (55%). Despite the encouraging PSA results, some urologists may still be reluctant to refer patients for salvage

owing to the variable clinical course after biochemical failure. Some patients have true local failure (especially those with high-risk features) and benefit from salvage treatment, while others have occult metastatic disease and derive greater benefit from treatment with androgen deprivation therapy. To complicate matters further, both clinicians and patients have concerns that any degree of urinary incontinence after surgery may worsen after treatment.

In the current series, Ajib et al demonstrate that it is safe to treat patients with SRT after robotic prostatectomy: men experience excellent functional urinary outcomes in addition to promising PSA control. They report that urinary quality of life is effectively unchanged from baseline (mean IPSS 3.3) up to 24 months of followup (mean IPSS 3.6). Furthermore, complete urinary continence (defined as no pads) is maintained in the majority of men at 24 months (70% compared to baseline 78%; $p=NS$). Any incontinence was typically very mild, usually only requiring one pad per day. In older series, where men were treated with open radical prostatectomy and four-field radiotherapy, the rate of complete continence (Grade 0) was less than 40% after treatment.² Allowing for differences in patient selection and treatment planning volumes, Ajib et al provide us with indirect evidence that advances in both surgical technique and radiation delivery may be contributing to the meaningful improvements in functional outcomes we see in this modern cohort of patients.

In radiation oncology, we have long known that toxicity is related to volume of normal tissue irradiated. We have always been conscientious about normal tissue-sparing and have exercised all options available to minimize toxicity. With advances in diagnostic imaging, treatment planning, delivery, and image guidance, highly conformal treatment can be administered with a very high degree of precision. This has translated into meaningful reduction in the volume of normal tissue irradiated, especially in the bladder and rectum. Considering these technological advances and their impact on reducing acute toxicity, it really comes as no surprise that we see so little change in functional urinary outcomes after SRT in the current series. When offered after

RARP, a large proportion of men are salvaged with RT and experience minimal change in urinary function and bother

The timing of RT after biochemical failure is also a complex decision. Large studies have not yet demonstrated a clear benefit for adjuvant over early salvage treatment with radiation;³⁻⁵ however, clinical evidence consistently suggests that early treatment with lower PSAs (≤ 0.2 ng/ml) at the time of SRT is associated with improved treatment outcomes. Questions about the utility of adjuvant treatment remain. Does early salvage at low PSAs (< 0.2 ng/ml) provide the same benefits as adjuvant treatment? While we await the results of the National Cancer Institute of Canada's (NCIC CTG) Radiotherapy and Androgen Deprivation in Combination After Local Surgery trial (RADICALS, IISRCTN#40814031), we are reassured to know that men can be safely treated with SRT after RARP with minimal impact on urinary function and continence. There remains little to be gained by delaying salvage and potentially a great deal to lose.

Competing interests: The author reports no competing personal or financial interests.

References

1. Michalski JM, Yan Y, Watkins-Bruner D, et al. Preliminary toxicity analysis of 3DCRT vs. IMRT on the high-dose arm of the RTOG 0126 prostate cancer trial. *Int J Radiat Oncol Biol Phys* 2013;87:932-8. <https://doi.org/10.1016/j.ijrobp.2013.07.041>
2. Pearce M, Choo R, Danjoux C, et al. Prospective assessment of gastrointestinal and genitourinary toxicity of salvage radiotherapy for patients with prostate-specific antigen relapse or local recurrence after radical prostatectomy. *Int J Radiat Oncol Biol Phys* 2008;72:792-8. <https://doi.org/10.1016/j.ijrobp.2008.05.063>
3. Tendulkar RD, Agrawal S, Gao T, et al. Contemporary update of a multi-institutional predictive nomogram for salvage radiotherapy after radical prostatectomy. *J Clin Oncol* 2016. [Epub ahead of print]. <https://doi.org/10.1200/JCO.2016.67.9647>
4. Thompson IM, Tangen CM, Paradelo J, et al. Adjuvant radiotherapy for pathological T3N0M0 prostate cancer significantly reduces risk of metastases and improves survival: Long-term followup of a randomized clinical trial. *J Urol* 2017; 181:956-62. <https://doi.org/10.1016/j.juro.2008.11.032>
5. Wiegel T, Botte D, Steiner U, et al. Phase 3 postoperative adjuvant radiotherapy after radical prostatectomy compared with radical prostatectomy alone in pT3 prostate cancer with postoperative undetectable prostate-specific antigen: ARO 96-02/AUO AP 09/95. *J Clin Oncol* 2009;27:2924-30. <https://doi.org/10.1200/JCO.2008.18.9563>

Correspondence: Dr. Martin Korzeniewski, Queen's University, Kingston, ON, Canada; korzenim@KGH.KARI.NET

NOW AVAILABLE! / MAINTENANT DISPONIBLE!

2017 CUA GUIDELINES in French



Topics include:

- CUA recommendations on prostate cancer screening and early diagnosis
- CUA guideline on the management of cystic renal lesions
- CUA guideline on adult overactive bladder
- CUA-PUC Canadian guidelines for the diagnosis, management and follow-up of cryptorchidism

GUIDES DE PRATIQUE 2017 DE L'AUC en français

Thèmes abordés :

- Recommandations de l'AUC sur le dépistage et le diagnostic précoce du cancer de la prostate
- Guide de pratique sur la prise en charge des lésions kystiques aux reins
- Guide de pratique sur la vessie hyperactive chez l'adulte
- Guide de pratique AUC-UPC pour le diagnostic, la prise en charge et le suivi de la cryptorchidie au Canada

cua.org

cuaj.ca