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## Live to SABR another day?

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Standard of care treatments should be considered and discussed - continuous and even intermittent androgen deprivation therapy (ADT)<sup>1-3</sup>.

For nodal recurrences, 3 different scenarios have been retrospectively evaluated (reviewed in ref<sup>4</sup>): stereotactic ablative radiotherapy (SABR), whole pelvic radiotherapy (WPRT) or salvage lymph node dissection (sLND). These treatments resulted in median progression-free survival (PFS) of 1–3 years, but was influenced by a heterogeneous use of ADT. The Belgian phase II, randomized STOMP trial without ADT used SABR and sLND, resulted in a 21 month freedom from palliative ADT without grade >1 toxicity (*J Clin Oncol in press*). WPRT with or without nodal boost probably prevents pelvic recurrences<sup>5</sup>, but it is unknown if it improves harder endpoints as compared to repeated SABR or sLND. Furthermore, it is unknown how these local therapies should be combined with systemic therapy.

This patient should be enrolled on studies examining the potential role of metastasis-directed therapy, such as the Baltimore ORIOLE trial<sup>6</sup>, British CORE (NCT02759783), Canadian PCS IX (NCT02685397), French STEREO-OS (NCT03143322), OLIGOPELVIS (NCT02274779/GETUG P07) and soon to be opened Belgian-Suisse STORM trial. The role of pelvic sLND for recurrent nodal PCa as assessed by 68Ga-PSMA PET-CT is being prospectively evaluated (NCT02974075).

If the patient has no access to a trial option, but wants a more aggressive treatment beyond ADT then SABR to all the suspicious nodes (e.g. 30 Gy in 3 fractions), comprehensive WPRT (e.g. 46–54 Gy from the aortic bifurcation down to distal iliacs with a nodal boost) or sLND is also not unreasonable.

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**CONFLICT OF INTEREST**

PTT co-owns the patent “Compounds and Methods of Use in Ablative Radiotherapy” (patent#: 9114158). PTT receives institutional research support from Medivation Inc-Astellas Pharma and has consulted for Dendreon Pharmaceuticals, Inc and RefleXion Medical.

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