

SUPPLEMENTARY FILE A

<i>Study/treatment group</i>	<i>N</i>	<i>Years of inclusion</i>	<i>Study specific inclusion criteria</i>
Feasibility study	30	2013-2017	<p><i>Patient characteristics</i></p> <p>Age \geq 18 years; Karnofsky Performance Status \geq 70; Fit for spinal anaesthesia; International Prostate Symptom Score $<$ 15.</p> <p><i>Tumour characteristics</i></p> <p>Tumour location technically feasible for brachytherapy; Tumour visible MRI and PSMA/choline-PET/CT; Biopsy proven local recurrence; PSA level \leq 10 ng/mL; PSA doubling time (PSADT) \geq 12 months; Tumour stage (MRI) \leq T2c; Recurrence \geq 2 years after primary radiotherapy; No lymph node and/or distant metastases.</p>
PRECISE study	42	2018-2019	<p><i>Patient characteristics</i></p> <p>See 'Feasibility study'.</p> <p><i>Tumour characteristics</i></p> <p>Tumour location technically feasible for brachytherapy; Concordance between PSMA-PET/CT and mp-MRI; PSA level \leq 20 ng/mL; PSA doubling time (PSADT) \geq 9 months; Tumour stage (MRI) \leq T3b; Recurrence \geq 2 years after primary radiotherapy; No lymph node and/or distant metastases.</p>
Non-study	78	2015-2020	<p><i>Patient characteristics</i></p> <p>See 'Feasibility study'.</p> <p><i>Tumour characteristics</i></p> <p>No specific criteria (i.e., those not eligible for treatment within either the feasibility of the PRECISE study, for example due to higher pre-salvage PSA and/or shorter PSA doubling time).</p>