

1 **Table 1.** Data collection schedule

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	Registration (within 4 weeks prior to registra tion)	Day 1	4 week s	8 week s	12 week s	24 week s	36 week s	48 week s	Every 12 weeks from 48 to 132 weeks	End of protocol treatment	Cancel protocol treatment	LPI from 3 years after last case registra tion
Assessment of eligibility criteria	•											
Obtain consent	•											
Registration	•											
Patient characteristics [#]	•											
Laboratory test	•		•	•	•	•	•	•	•	•	•	
PSA	•			•	•	•	•	•	•	•	•	
Medication						•						
Adverse events [§]						•				•	•	
Electrocardiogram	•				•	•		•	•	•	•	
CT bone scintigraphy [‡]	•				•	•		•	•	•	•	

Confirmation of disease progression					•	•	•	•	•	•	•	
Simultaneous survival check												•
Prohibited concomitant medications and therapies		•								•		

3 #Sex, date of birth, age, medical history, comorbidity, primary disease, TNM classification, number of days after surgery, details of
4 adjuvant therapy.

5 §Grade 3 or 4 (according to Common Terminology Criteria for Adverse Events version 4.0).

6 ‡Magnetic resonance imaging may be used instead of CT, with or without contrast media, but each patient should be assessed and
7 judged using one method only.

8 CT, computed tomography; LPI, last patient in; PSA, prostate-specific antigen