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Systemic Therapy for Advanced or Metastatic Prostate cancer: Evaluation of Drug Efficacy (STAMPEDE) investigators (2018). Radiotherapy to the primary tumour for newly diagnosed, metastatic prostate cancer (STAMPEDE): a randomised controlled phase 3 trial. Lancet, 392(10162), 2353-2366.
https://doi.org/10.1016/S0140-6736(18)32486-3

Published in:
Lancet

Document Version:
Publisher's PDF, also known as Version of record

Queen's University Belfast - Research Portal:
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Radiotherapy to the primary tumour for newly diagnosed, metastatic prostate cancer (STAMPEDE): a randomised controlled phase 3 trial

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Summary

Background Based on previous findings, we hypothesised that radiotherapy to the prostate would improve overall survival in men with metastatic prostate cancer, and that the benefit would be greatest in patients with a low metastatic burden. We aimed to compare standard of care for metastatic prostate cancer, with and without radiotherapy.

Methods We did a randomised controlled phase 3 trial at 117 hospitals in Switzerland and the UK. Eligible patients had newly diagnosed metastatic prostate cancer. We randomly allocated patients open-label in a 1:1 ratio to standard of care (control group) or standard of care and radiotherapy (radiotherapy group). Randomisation was stratified by hospital, age at randomisation, nodal involvement, WHO performance status, planned androgen deprivation therapy, planned docetaxel use (from December, 2015), and regular aspirin or non-steroidal anti-inflammatory drug use. Standard of care was lifelong androgen deprivation therapy, with up-front docetaxel permitted from December, 2015.

Men allocated radiotherapy received either a daily (55 Gy in 20 fractions over 4 weeks) or weekly (36 Gy in six fractions over 6 weeks) schedule that was nominated before randomisation. The primary outcome was overall survival, measured as the number of deaths; this analysis had 90% power with a one-sided α of 2.5% for a hazard ratio (HR) of 0.75. Secondary outcomes were failure-free survival, progression-free survival, metastatic progression-free survival, prostate cancer-specific survival, and symptomatic local event-free survival. Analyses used Cox proportional hazards and flexible parametric models, adjusted for stratification factors. The primary outcome analysis was by intention to treat. Two prespecified subgroup analyses tested the effects of prostate radiotherapy by baseline metastatic burden and flexible parametric models, adjusted for stratification factors. The primary outcome analysis was by intention to treat. Two prespecified subgroup analyses tested the effects of prostate radiotherapy by baseline metastatic burden and radiotherapy schedule. This trial is registered with ClinicalTrials.gov, number NCT00268476.

Findings Between Jan 22, 2013, and Sept 2, 2016, 2061 men underwent randomisation, 1029 were allocated the control and 1032 radiotherapy. Allocated groups were balanced, with a median age of 68 years (IQR 63–73) and median amount of prostate-specific antigen of 97 ng/mL (33–315). 367 (18%) patients received early docetaxel. 1082 (52%) and 1032 radiotherapy. Allocated groups were balanced, with a median age of 68 years (IQR 63–73) and median amount of prostate-specific antigen of 97 ng/mL (33–315). 367 (18%) patients received early docetaxel. 1082 (52%) and 1032 radiotherapy. Allocated groups were balanced, with a median age of 68 years (IQR 63–73) and median amount of prostate-specific antigen of 97 ng/mL (33–315).

Radiotherapy improved failure-free survival (HR 0.76, 95% CI 0.68–0.84; p<0.0001) but not overall survival (0.75). Secondary outcomes were failure-free survival, progression-free survival, metastatic progression-free survival, prostate cancer-specific survival, and symptomatic local event-free survival. Analyses used Cox proportional hazards and flexible parametric models, adjusted for stratification factors. The primary outcome analysis was by intention to treat. Two prespecified subgroup analyses tested the effects of prostate radiotherapy by baseline metastatic burden and radiotherapy schedule. This trial is registered with ClinicalTrials.gov, number NCT00268476.

Interpretation Radiotherapy to the prostate did not improve overall survival for unselected patients with newly diagnosed metastatic prostate cancer.

Funding Cancer Research UK, UK Medical Research Council, Swiss Group for Clinical Cancer Research, Astellas, Clovis Oncology, Janssen, Novartis, Pfizer, and Sanofi-Aventis.

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Introduction Patients with metastatic cancer typically receive systemic treatment, with local therapy reserved—if required—for symptom palliation. However, local treatment to the primary tumour might be more useful than previously appreciated. In animal models of cancer, primary tumours metastasise not merely by disseminating tumour cells into the circulation but also by priming the
About the role of prostate radiotherapy in metastatic prostate cancer. Our findings showed no overall survival benefit of radiotherapy to the prostate in men with newly diagnosed prostate cancer. However, a subgroup analysis supported the hypothesis of HORRAD, that prostate radiotherapy improves survival in men with low metastatic burden.

Methods
Study design and participants
We did a randomised controlled phase 3 trial at 117 hospitals in Switzerland and the UK. Eligible patients had prostate cancer that was newly diagnosed, with no previous radical treatment, and had metastatic disease confirmed on a bone scintigraphic scan and soft-tissue imaging done within 12 weeks of starting androgen deprivation therapy. All patients were intended for long-term androgen deprivation therapy and started treatment no earlier than 12 weeks before randomisation. There were no age restrictions; patients were required to have no contraindications to radiotherapy and no clinically significant cardiovascular history.

This trial was done in accordance with Good Clinical Practice guidelines and the Declaration of Helsinki and had relevant regulatory and ethics approvals. All patients gave written informed consent. The rationale and design have been described previously.13 Full details are in the protocol.

Randomisation and masking
Patients were randomised centrally using a computerised algorithm, which was developed and maintained by the Medical Research Council (MRC) Clinical Trials Unit at University College London. Minimisation with a random element of 20% was used, stratifying for hospital, age at randomisation (<70 years vs ≥70 years), nodal involvement (negative vs positive vs indeterminate), WHO performance status (0 vs 1 or 2), planned androgen deprivation therapy, and regular aspirin or non-steroidal anti-inflammatory drug use (yes or no). Planned docetaxel use was added as a stratification factor on Dec 17, 2015. Allocation was 1:1 to either standard of care (control) or standard of care and radiotherapy
(radiotherapy). Patients and clinical and study staff were aware of the treatment allocation for practical reasons, and the key efficacy outcome measures were objective.

**Procedures**

All patients received lifelong androgen deprivation therapy as either gonadotrophin-releasing hormone agonists or antagonists or orchidectomy. Docetaxel was permitted in addition to hormone therapy after its approval in the UK on Dec 17, 2015. Docetaxel, when used, was given as six 3-weekly cycles of 75 mg/m², with or without prednisolone 10 mg daily.

External-beam radiotherapy to the prostate was given as one of two schedules nominated before randomisation: either 36 Gy in six consecutive weekly fractions of 6 Gy, or 55 Gy in 20 daily fractions of 2.75 Gy over 4 weeks. Radiotherapy was given with the patient supine and with a full bladder and an empty rectum. The planning target volume consisted of the prostate only, with an 8 mm margin posteriorly and a 10 mm margin elsewhere.

Radiotherapy was to commence as soon as practicable after randomisation, and within 3–4 weeks after the last docetaxel dose.

Patients were followed up every 6 weeks until 6 months after randomisation, then every 12 weeks to 2 years, then annually thereafter. Nadir PSA was the lowest level of PSA reported within 24 weeks after randomisation. Toxic effects and symptoms were reported at regular follow-up visits or when an adverse event was categorised as serious. Adverse events were graded with the National Cancer Institute Common Terminology Criteria for Adverse Events (CTCAE) version 4.0. Adverse effects on the bowel and bladder during radiotherapy, and possible long-term effects of radiotherapy, were recorded separately in patients assigned standard of care and radiotherapy using the Radiation Therapy Oncology Group (RTOG) scale.

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For the protocol see http://www.stampedetrial.org

**Figure 1:** Trial profile
Metastatic burden at randomisation was assessed through whole-body scintigraphy and CT or MRI staging scans. Scans were centralised and reviewed by one of us (AA), with 10% independent review by a radiologist (HD). The metastatic burden was classified according to the definition used in the CHAARTED trial: high metastatic burden was defined as four or more bone metastases with one or more outside the vertebral bodies or pelvis, or visceral metastases, or both; all other assessable patients were considered to have low metastatic burden.

**Outcomes**

The primary efficacy outcome was overall survival, defined as time from randomisation to death from any cause. Failure-free survival was the primary activity outcome measure for interim analyses and was defined as time from randomisation to first evidence of at least one of: biochemical failure; progression either locally, in lymph nodes, or in distant metastases; or death from prostate cancer. Biochemical failure was based on a rise above the lowest PSA value reported within 24 weeks after enrolment of 50% and to at least 4 ng/mL; patients without a fall of 50% were considered to have biochemical failure at time zero. Secondary outcomes were progression-free survival (defined as failure-free survival but without biochemical events) and metastatic progression-free survival (defined as time from randomisation to new metastases or progression of existing metastases or death). Cause of death was determined by the site investigator, with some causes reclassified as prostate cancer according to predefined criteria that indicated prostate cancer to be the likely cause. Symptomatic local events were defined as any of the following: urinary tract infection, new urinary catheterisation, acute kidney injury, transurethral resection of the prostate, urinary-tract obstruction, ureteric stent, nephrostomy, colostomy, and surgery for bowel obstruction. Patients without the event of interest were censored at the time last known to be event-free.

**Statistical analysis**

This randomised comparison was incorporated within the Systemic Therapy for Advanced or Metastatic Prostate...
cancer: Evaluation of Drug Efficacy (STAMPEDE) multi-arm multistage (MAMS) platform protocol (appendix p 6). It was designed with a seamless phase 2/3 approach. The sample size was calculated using nstage and its predecessor programs in Stata, which enable design of MAMS trials. Assuming, for the control group, a median failure-free survival of roughly 1 year and median survival of about 3.5 years, we targeted a 25% relative improvement (HR 0.75) in both failure-free survival and overall survival for the group allocated radiotherapy to the prostate over the control group.

For the efficacy stage analysis of the pairwise comparison of standard of care and radiotherapy versus standard of care for overall survival, approximately 267 deaths in patients allocated to the control group were needed for 90% power and a one-sided α of 2.5%, accounting for three intermediate analyses of failure-free survival (analysed June, 2014, November, 2014, and May, 2015). For this comparison, the pairwise and family-wide error rates were judged very similar, because of the limited overlap in events with other reported comparisons from the protocol and the non-binding nature of the interim analyses.

The initial sample size target was 1250 patients. During the trial, weekly and daily radiotherapy schedules were nominated approximately equally. Therefore, the sample size was increased to roughly 1800, without reference to outcome data, to provide good power for failure-free survival in each radiotherapy schedule-defined subgroup when the comparison reached its target power overall, assuming that the effect of radiotherapy would be the same regardless of schedule. We predicted about 300 failure-free survival events in the control group on each schedule at the time of the main analysis, which would provide approximately 90% power with a one-sided α of 0.015 to detect an HR of 0.75. The effect of radiotherapy on survival within a nominated radiotherapy schedule would be investigated if there was both an effect on failure-free survival and 200 or more deaths in the control group were reported for that nominated schedule.

In May, 2018, based on accumulating external data and without reference to any data from this comparison in STAMPEDE, we prespecified that any effect from radiotherapy would be greatest in patients with a low baseline metastatic burden, we anticipated more than 90% power for failure-free survival events in the control group on each schedule at the time of the main analysis, which would provide approximately 90% power with a one-sided α of 0.015 to detect an HR of 0.75. The effect of radiotherapy on survival within a nominated radiotherapy schedule would be investigated if there was both an effect on failure-free survival and 200 or more deaths in the control group were reported for that nominated schedule.

Standard survival analysis methods were used to analyse time-to-event data in Stata version 15. A non-parametric stratified log-rank test was used to detect a difference in survival between treatment groups; this analysis was stratified across the minimisation factors used at randomisation (except hospital and planned androgen deprivation therapy) plus protocol-specific periods defined by other arms recruiting to STAMPEDE or changes to standard of care that could affect the population being randomised. Cox proportional hazards regression models adjusting for the same stratification factors and stratified by time were used to estimate relative treatment effects. An HR less than 1.00 favoured radiotherapy. Flexible parametric models were fitted with degrees of freedom (5,5) and adjusted for stratification factors and time. Medians and 3-year survival estimates are presented from the flexible parametric models fitted.
to the data; these are more reliable than reading the Kaplan-Meier curves; graphs show estimated survival over time from both. The proportional hazards assumption was tested; restricted mean survival time was emphasised in the presence of non-proportionality, using a t-star of 59 months as determined by the Royston and Parmar method.18 Cause-specific and Fine and Gray regression models19 were used for competing risk analysis of prostate cancer-specific, lymph node, and metastatic progression-free and symptomatic local event-free survival. All tests are presented as two-sided, with 95% CIs and relevant p values.

Subgroup analyses were prespecified for the nominated radiotherapy schedule (daily vs weekly) and for baseline metastatic burden (low vs high), when determinable. Exploratory interaction analyses considered the consistency of treatment effect within stratification factors, by time, by Gleason score, and by PSA before hormone therapy.

Median follow-up was ascertained by reverse-censoring on death. All patients were included in the primary efficacy analysis according to allocated treatment, and the analysis was done on an intention-to-treat basis. Adverse event data are shown for the safety population, which consisted of patients with at least one follow-up assessment analysed according to the treatment approach started; patients were excluded if they had no adverse event data. A sensitivity analysis was done on an intention-to-treat basis. Data for symptomatic local events are also presented. All other analyses are exploratory.

Accumulating interim data were reviewed by an Independent Data Monitoring Committee, guided by lack-of-benefit stopping guidelines.

This trial is registered at ClinicalTrials.gov (number NCT00268476) and ISRCTN.com (ISRCTN78818544).

Role of the funding source
MRC employees contributed to study design, data collection, data analysis, data interpretation, and writing of this report. CDB, MRS, APH, and AA had access to raw data. The corresponding author had final responsibility for the decision to submit for publication.

Results
Between Jan 22, 2013, and Sept 2, 2016, 2061 patients were randomly allocated either standard of care

### Table 2: Summary of estimated treatment effect for main outcome measures, for all patients and by metastatic burden

| Outcome Measure                                      | Adjusted hazard ratio (95% CI) | Survival at 3 years* | Restricted mean survival time (months)* |
|------------------------------------------------------|--------------------------------|----------------------|----------------------------------------|
|                                                      | Control                        | Radiotherapy         | Control                                | Radiotherapy | Difference (95% CI) |
| Overall survival                                     |                                |                      |                                        |              |                   |
| All patients                                         | 0·92 (0·80–1·06)               | 62%                  | 41·6                                   | 42·5         | 1·0 (–0·6 to 2·5)  |
| Low metastatic burden                                | 0·68 (0·52–0·90)               | 73%                  | 45·4                                   | 49·1         | 3·6 (1·0 to 6·2)   |
| High metastatic burden                               | 1·07 (0·90–1·28)               | 54%                  | 38·8                                   | 37·6         | –1·2 (–3·5 to 1·1) |
| Failure-free survival                                |                                |                      |                                        |              |                   |
| All patients                                         | 0·76 (0·68–0·84)               | 23%                  | 21·4                                   | 26·2         | 4·8 (2·8 to 6·7)   |
| Low metastatic burden                                | 0·59 (0·49–0·72)               | 33%                  | 27·4                                   | 36·1         | 8·6 (5·6 to 11·7)  |
| High metastatic burden                               | 0·88 (0·77–1·01)               | 17%                  | 17·3                                   | 18·8         | 1·5 (0·7 to 3·6)   |
| Progression-free survival                            |                                |                      |                                        |              |                   |
| All patients                                         | 0·96 (0·85–1·08)               | 44%                  | 32·4                                   | 33·1         | 0·7 (–0·9 to 2·3)  |
| Low metastatic burden                                | 0·78 (0·63–0·98)               | 58%                  | 39·4                                   | 42·9         | 3·5 (0·4 to 6·7)   |
| High metastatic burden                               | 1·09 (0·94–1·26)               | 35%                  | 28·0                                   | 26·2         | –1·8 (–4·3 to 0·8) |
| Metastatic progression-free survival                 |                                |                      |                                        |              |                   |
| All patients                                         | 0·97 (0·86–1·10)               | 47%                  | 33·9                                   | 34·4         | 0·4 (–1·5 to 2·4)  |
| Low metastatic burden                                | 0·80 (0·63–1·01)               | 62%                  | 41·1                                   | 44·2         | 3·1 (0·2 to 6·0)   |
| High metastatic burden                               | 1·10 (0·95–1·28)               | 37%                  | 29·3                                   | 27·3         | –2·0 (–4·7 to 0·7) |
| Prostate cancer-specific survival                    |                                |                      |                                        |              |                   |
| All patients†                                        | 0·93 (0·80–1·09)               | 66%                  | 43·9                                   | 44·6         | 0·7 (–1·1 to 2·5)  |
| Low metastatic burden                                | 0·65 (0·47–0·90)               | 79%                  | 48·6                                   | 51·8         | 3·3 (0·0 to 5·5)   |
| High metastatic burden                               | 1·10 (0·92–1·32)               | 58%                  | 40·6                                   | 39·0         | –1·6 (–3·9 to 0·7) |
| Symptomatic local event-free survival                |                                |                      |                                        |              |                   |
| All patients                                         | 1·07 (0·93–1·22)               | 57%                  | 38·2                                   | 37·2         | –1·1 (–3·1 to 0·9) |
| Low metastatic burden                                | 0·82 (0·64–1·05)               | 65%                  | 41·6                                   | 44·0         | 2·4 (–0·7 to 5·4)  |
| High metastatic burden                               | 1·23 (1·05–1·46)               | 50%                  | 35·8                                   | 32·2         | –3·6 (–6·2 to –1·0) |

Hazard ratio and restricted mean survival time differences are for radiotherapy relative to control. *Survival probabilities and restricted mean survival time estimates are taken from flexible parametric models (t-star, 59 months). †Competing risks analysis, sub-hazard ratio 0·94, 95% CI 0·81–1·10; p=0·431.
(control group, n=1029) or standard of care and radiotherapy (radiotherapy group, n=1032; figure 1). Groups were well balanced with respect to baseline characteristics (table 1). Median age was 68 years (IQR 63–73) and median PSA before androgen deprivation therapy was 97 ng/mL (33–315). 1630 (79%) patients had a Gleason score of 8–10. 1836 (89%) had bone metastases. 1466 (71%) had a WHO performance status of zero. Baseline characteristics of 1939 (94%) patients in whom metastatic disease burden could be determined are shown in the appendix (p 1).

Standard hormone therapy was luteinising hormone-releasing hormone analogues for 2046 (99%) men. Standard of care included docetaxel for 367 (18%) patients. Of 2061 patients undergoing random assignment, roughly half were nominated for each radiotherapy schedule, with 979 (48%) nominating the weekly schedule and 1082 (52%) the daily schedule. Of 968 patients assigned radiotherapy who started radiotherapy within 1 year after randomisation, 906 received their planned schedule and 62 received the alternative or another schedule. Two patients assigned radiotherapy received their planned schedule later than 1 year after randomisation, and 62 did not receive radiotherapy at all (mainly because of patient’s choice). In patients who started radiotherapy, median time to starting radiotherapy was 35 days (IQR 28–60) after randomisation, and 95 days (74–120) from starting hormone therapy (most patients started androgen deprivation therapy before randomisation; appendix p 2). Only 20 (2%) patients allocated to the control group received radiotherapy within 1 year of randomisation.

Median follow-up was 37 months (IQR 24–48). 391 patients assigned to the control group died (median survival 46 months [IQR 27–not reached]; 3-year survival 62%). Compared with controls, no survival advantage was noted with radiotherapy (stratified log-rank test \(p=0.451\); HR 0.92, 95% CI 0.80–1.06; \(p=0.266\)), with 370 deaths in the radiotherapy group (median survival 48 months [IQR 27–not reached]; 3-year survival 65%; figure 2A; table 2). There was no evidence of non-proportional hazards.

Figure 3 shows prespecified and exploratory subgroup analyses. In the analysis by metastatic burden, overall survival was improved in patients with low metastatic burden at baseline who were allocated radiotherapy (HR 0.68, 95% CI 0.52–0.90; \(p=0.007\); 3-year survival 73% with control vs 81% with radiotherapy; table 2), with no evidence of non-proportional hazards. We found some evidence of heterogeneity of treatment effect by metastatic burden (interaction \(p=0.0098\); figure 4). In patients with a high metastatic burden, there was no evidence of a treatment effect (HR 1.07, 95% CI 0.90–1.28; \(p=0.420\)). The appendix (p 7) shows further exploratory consistency-of-effect analyses.

758 failure-free survival events were reported in patients assigned to the control group, largely driven by rising PSA (appendix p 3); median failure-free survival was 13 months (IQR 6–33) and 3-year failure-free survival was 23%. In patients assigned to the radiotherapy group, 685 failure-free survival events were reported, with median failure-free survival of 17 months (IQR 8–53) and 3-year failure-free survival of 32%. Overall, failure-free survival was improved with radiotherapy (HR 0.76, 95% CI 0.68–0.84; \(p=0.0001\); figure 2B). There was some evidence of non-proportional hazards (\(p=0.066\)). An analysis of restricted mean survival time found mean

![Figure 3: Treatment effect on overall survival within selected baseline categories](https://example.com/figure3.png)

**HR**=hazard ratio. PSA before androgen deprivation therapy (continuous), \(p=0.029\); effect of adding radiotherapy is smaller with higher PSA. Patients with unknown T stage (TX), unknown N category (NX), or unknown Gleason sum score are not presented in the forest plot and do not contribute to interaction test results. Dotted line shows the overall hazard ratio. (A) Prespecified subgroup analyses. (B) Exploratory subgroup analyses.
failure-free survival, restricted to the first 59 months on trial, was 21.4 months in the control group, compared with 26.2 months with radiotherapy (difference 4.8 months, 95% CI 2.8–6.7; p<0.0001; table 2).

In the prespecified subgroup analysis by metastatic burden, failure-free survival was improved in patients with low metastatic burden at baseline who were allocated radiotherapy (HR 0.59, 95% CI 0.49–0.72; p<0.0001; table 2). Evidence of a differential treatment effect from radiotherapy compared with the high metastatic burden subgroup was also noted (interaction p=0.002; HR 0.88, 95% CI 0.71–1.01; p=0.059; figure 4). The appendix (p 8) shows prespecified and exploratory consistency of effect analyses.

643 (84%) of 761 deaths were attributed to prostate cancer (329 [84%] of 391 in the control group and 314 [85%] of 370 in the radiotherapy group). A adjusted competing risks regression for prostate cancer specific survival using the Fine and Gray method provided no evidence of an overall treatment effect (sub-HR 0.94, 95% CI 0.81–1.10; robust p=0.431; table 2). There was evidence of an effect in patients with low metastatic burden (sub-HR 0.65, 95% CI 0.47–0.90; robust p=0.010), but no evidence of a treatment effect was noted in patients with high metastatic burden (1.11, 0.92–1.33; robust p=0.279). A significant interaction was seen between treatment effect and metastatic burden (robustly estimated interaction p=0.007).

The appendix (pp 9, 10) shows the analysis of progression-free survival for all patients and by baseline metastatic burden. A treatment effect was only noted in patients with a low metastatic burden (HR 0.78, 95% CI 0.63–0.98; p=0.033; table 2).

One or more symptomatic local events were reported by 432 (42%) of 1029 patients allocated to the control group compared with 450 (44%) of 1032 patients assigned to the radiotherapy group. There was no evidence of a difference in time to first symptomatic local event by treatment allocation (HR 1.07, 95% CI 0.90–1.28; p=0.420).

Table 3 shows the number of patients reporting each type of symptomatic local event at least once.

Figure 4: Overall survival and failure-free survival by treatment and metastatic burden

HR=hazard ratio. Solid lines show the Kaplan-Meier analysis and dotted lines show the flexible parametric model.
analyses in 1082 patients who nominated the daily schedule before randomisation (55 Gy in 20 fractions over 4 weeks) found strong evidence of a failure-free survival advantage with radiotherapy compared with control (HR 0·69, 95% CI 0·59–0·80; p=0·0001). Among these 1082 patients, 212 deaths were reported in the control group and 188 in the radiotherapy group (stratified log-rank p=0·123; HR 0·86, 95% CI 0·71–1·05; p=0·128). There was insufficient evidence of a difference in failure-free survival in 979 patients who nominated the weekly radiotherapy schedule (36 Gy in six fractions over 6 weeks; HR 0·85, 95% CI 0·73–0·99; p=0·033) to report on survival.

Adverse effects on the RTOG scale during radiotherapy were modest, with 48 (5%) of 920 patients allocated radiotherapy who started radiotherapy and who completed at least one acute toxicity form reporting grade 3 or 4 adverse events; 43 (5%) patients reported their worst acute bladder toxic effect as grade 3 or 4, and eight (1%) reported their worst acute bowel toxic effect as grade 3 or 4 (table 4; appendix p 11); no grade 5 toxic effects were reported. The incidence of acute bladder and bowel effects (grade 1–4) was lower for those who nominated the weekly radiotherapy schedule than for those who nominated the daily schedule (282 [65%] bladder and 206 [47%] bowel vs 341 [71%] bladder and 297 [62%] bowel). Patients in both control and radiotherapy groups reported a low incidence of grade 3 and 4 RTOG late effects (one [1%] control vs 37 [4%] radiotherapy; table 5).

The proportion of patients in the safety population reporting at least one severe adverse event of CTCAE grade 3 or worse was similar in both study groups and was dominated by side-effects associated with long-term hormone therapy (398 [38%] of 1050 in the control group and 380 [39%] of 985 in the radiotherapy group; appendix pp 4, 12); with no evidence of a difference in time to first grade 3 or worse event (HR 1·01, 95% CI 0·97–1·05; p=0·128). In 2028 patients with adverse event data at approximately 6 months, the proportions reporting a grade 3 or worse adverse event were similar (225 [21%] of 1047 in the control group and 212 [22%] of 981 in the radiotherapy group). Of 1125 patients with adverse event data at 1 year, 63 (12%) of 531 patients in the control group and 78 (13%) of 594 in the radiotherapy group reported a grade 3 or worse adverse event. At 2 years, of 533 patients with data available, 37 (15%) of 240 in the control group and 37 (13%) of 293 in the radiotherapy group reported a grade 3 or worse event. The pattern and levels of adverse events were very similar when considering the intention-to-treat population (data not shown). No deaths were reported as related to the research treatment.

The appendix (pp 13, 14) shows time to first new treatment (data not shown). No deaths were reported as related to the research treatment. There were no reported grade 5 late radiotherapy toxic events. RTOG=Radiation Therapy Oncology Group.

Table 4: Worst reported acute radiotherapy bladder and bowel toxic effect (RTOG scale) in patients allocated radiotherapy

Table 5: Grade 3 or 4 worst late radiotherapy toxicity score (RTOG scale) in patients who received radiotherapy (for research or progression)
Radiotherapy therefore seems safe to conclude that radiotherapy would improve survival of men with a low metastatic burden, which represented 40% of the comparison population.

Our subgroup finding meets all criteria proposed by Sun and colleagues to assess credibility of subgroup effects;²⁰ low metastatic burden status was determined from scans taken before randomisation; the hypothesis—including the direction of the effect—was specified a priori; only a few hypothesised subgroup effects were tested; the interaction test suggested a low likelihood that the apparent subgroup effect could be accounted for by chance; the subgroup effect was independent of other assessed variables; the size of the subgroup effect was large (HR 0·68 for low metastatic burden and HR 1·07 high metastatic burden); and the subgroup effect was large (HR 0·68 for low metastatic burden). It also seems plausible that the effect of local radiotherapy would be diminished in patients with a greater burden of metastatic disease. There is, therefore, good reason to accept that prostate radiotherapy improves survival of men with a low metastatic burden and that it should now be a standard treatment.

Unlike many other new interventions for metastatic cancer, prostate radiotherapy does not require regulatory approval and is readily available at modest cost in most parts of the world.

When this comparison was designed, the standard radical radiotherapy dose schedule for localised prostate cancer was 74 Gy in 37 fractions over 7·5 weeks. It was felt that this schedule would be too burdensome for patients with metastatic disease, and the two more convenient schedules permitted in the trial were chosen based on a survey of investigators’ opinions. The trial has proven the principle that local radiotherapy can improve survival, but the optimum dose schedule and technique are uncertain. Radical radiotherapy for localised prostate cancer is now typically given to a dose of 60 Gy in 20 fractions over 4 weeks.²¹ With contemporary techniques for target delineation and treatment delivery, this schedule is well tolerated²² and might be expected to be at least as effective as the two schedules tested in the trial.

It is well known that prostate radiotherapy improves survival for men with locally advanced (T³–4 N0 M0) prostate cancer.²² We have now found that prostate radiotherapy also improves survival for men with a low metastatic burden (T³³ N³³ M¹³¹) prostate cancer. It therefore seems safe to conclude that radiotherapy would also improve survival for men with pelvic node-positive prostate cancer (T³³ N¹ M0³). This is important, because it is not feasible to do randomised trials specifically in men with non-metastatic node-positive prostate cancer and because such men often receive systemic treatment alone. In the current study, roughly 60% of patients were N1 in both the high and low metastatic burden subgroups.

The benefit we have shown for prostate radiotherapy in prostate cancer with a low metastatic burden also raises another question: would there be further benefit from additional radiotherapy to the oligometastases themselves?

Low metastatic burden disease is sometimes known as oligometastatic. Although this term is widely used, it is imprecise and potentially misleading because it implies only a small number of metastases. Patients with low metastatic burden disease, according to the CHAARTED definition, may have an unlimited number of metastases provided they are confined to lymph nodes and the axial skeleton.

Our data have several strengths to note. This is a large randomised dataset with broad engagement from more than 100 hospitals across Switzerland and the UK. By incorporating the comparison into the established STAMPEDE protocol, following peer-review and protocol amendment, we recruited to an enlarged target well ahead of schedule (2061 patients in 3·5 years rather than 1250 patients in 4 years).

Our data also have some limitations. First, the possible clinical relevance of metastatic burden in patients with prostate cancer only became widely apparent when the CHAARTED trial reported.²³ We determined metastatic

Figure 5: Time from randomisation to life-prolonging therapy.
Articles

Caution will be required in extrapolating these results to patients imaged with more sensitive techniques (eg, PSMA PET). For example, patients with low metastatic burden on conventional imaging should not be denied prostate radiotherapy because they have additional lesions identified on a PET scan.

In summary, radiotherapy to the prostate did not improve survival for unselected patients with newly diagnosed metastatic prostate cancer, but, in a pre-specified subgroup analysis, overall survival did improve in men with a low metastatic burden. Therefore, prostate radiotherapy should be a standard treatment option for men with a low metastatic burden. These findings also raise the possibility that local treatment to the primary tumour should be explored for patients with small-volume metastatic disease from other malignant diseases.

**Contributors**

CCP was comparison chief investigator. NDJ was protocol chief investigator. CCP, MRS, NDJ, MDM, DPD, NWc, and MKBP contributed to comparison design. CLA contributed to trial operations. APH, AA, and NWc collated and reviewed bone and CT scans. HD reviewed bone and CT scans. MRS, CDB, CCP, MRC, NWc, MKBP, NDJ, APH, and AA contributed to the analysis plan. CDB, MRS, APH, and AA did analyses. CCP, CDB, MRS, NDJ, MKBP, NWc, APH, and AA contributed to writing of the report. The Trial Management Group consisted of NDJ (chair), MDM, and NWc (vice chairs), and MKBP, MRS, DPD, NWc, AWR, REL, GNT, WC, SG, ZIM, SG, CCP, GA, DM, JMR, and RJF. All authors collated and interpreted data, and edited, reviewed, and approved the final report.

**Systemic Therapy for Advanced or Metastatic Prostate cancer: Evaluation of Drug Efficacy (STAMPEDE) investigators**

**Independent Oversight Committee Members:** Independent Data Monitoring Committee—John Yarnold (chair), Ronald de Wit, Bertrand Tombal (previous: Doug Altman, Reg Hall, Chris Williams); Independent members of Trial Steering Committee: Jonathan Ledermann (chair), Jan Erik Damber, Richard Emsley, Alan Horwich (previous: John Fitzpatrick, David Kirk, Jim Paul). Participating Site List (hospital [n patients during recruitment window: recruiting investigators])—Addenbrooke’s Hospital (7: D Mazhar); Ashford William Harvey Hospital (10: C Thomas, N Mithal, A Edwards); Aylesbury, Stoke Mandeville Hospital (4: T Pwint, P Camilleri); Ayr Hospital (15: H Glen, J Ansari); Barnet General Hospital (15: U McGovern, A Ichlholt); Basingstoke & North Hampshire Hospital (7: R Shaffer); Bath, Royal United Hospital (14: O Frim, M Beresford, P Kehagiology): Belfast City Hospital (66: J’O’Sullivan, D Mitchell, S Jain, P L Shum); Birmingham, City Hospital (14: D Ford); Birmingham, Good Hope Hospital (10: D Ford); Birmingham, Heartlands Hospital (11: A Zarkar); Birmingham, Queen Elizabeth (36: N James); Blackburn, East Lancashire Trust (52: O Parikh, N Charnley); Bolton, Royal Bolton Hospital (11: T Elliott); Boston, Pilgrim Hospital (18: M Panades, D Ballesteros-Quintal); Bournemouth, Royal Bournemouth Hospital (23: S Brock); Bradford Royal Infirmary (20: S Brown); Brighton, Royal Sussex County Hospital (17: A Robinson, G Plantanios); Bristol Haematology & Oncology Centre (58: A Bahl, C Herbert, S Massignon); Burton, Queen’s Hospital (40: M Smith-Howell, S Chettyawardana, P Pattu); Bury St Edmunds, West Suffolk Hospital (14: C Woodward, Y Rimmer); Cardiff, Velindre (86: J Yarnold); Crewe, Leighton Hospital (22: J Wylie); Cumbria, Cumberland Infirmary (8: A Kumar); Darlington Memorial Hospital (26: M Kajzzi, J Hardman, C Peedell); Derby, Royal Derby Hospital (43: P Chakrabotti); Devon, North Devon District Hospital
Declaration of interests

GA reports personal fees from Veridx, Novartis, Millennium Pharmaceuticals, Takeda, and Sanofi-Aventis, outside the submitted work; personal fees and non-financial support from Roche/Ventana, Astellas, Medivation, Pfizer, Abbott Laboratories, Essa Pharmaceuticals, Takeda, and Sanofi, outside the submitted work; and personal fees from Janssen, outside the submitted work; and personal fees from Innocrin Pharmaceuticals, Novartis, and Cell Search, outside the submitted work; and personal fees from MaxiVax SA, outside the submitted work.

SC reports personal fees from Janssen, Astellas, Takeda, and Sanofi, outside the submitted work; grants, personal fees, and non-financial support from Innocrin Pharmaceuticals, Novartis, and Cell Search, outside the submitted work; and personal fees from Innocrin Pharmaceuticals, Novartis, and Cell Search, outside the submitted work.

GC reports personal fees from Astellas, Takeda, Sandoz, Janssen, and the Institute of Cancer Research, during the conduct of the study; grants from Cancer Research UK (programme grant C46/A10588, C33589/A1972), during the conduct of the study; personal fees, speaker fees, and advisory board fees from Nectar Therapeutics and ProteoMediX, outside the submitted work; and personal fees from Bayer, outside the submitted work. DPD reports grants and other from the National Institute for Health Research Biomedical Research Centre at the Royal Marsden NHS Foundation Trust and the Institute of Cancer Research, during the conduct of the study; grants from Cancer Research UK (programme grant C46/A10588, C33589/A1972), during the conduct of the study; personal fees from Amgen, Astellas, Takeda, Sandoz, Janssen, and the Institute of Cancer Research (share of royalties for abiraterone), outside the submitted work; and has a patent (EP1933799B1) issued. SC reports personal fees and honoraria to her hospital from Bayer, outside the submitted work; and personal fees from Bayer, outside the submitted work. WC reports advisory board fees from Bayer and AstraZeneca, outside the submitted work. ABi reports advisory board fees from Bayer and AstraZeneca, outside the submitted work; honoraria to her hospital from Bayer, outside the submitted work; and personal fees and non-financial support from Roche/Ventana, Astellas, Medivation, Pfizer, Abbott Laboratories, Essa Pharmaceuticals, Takeda, and Sanofi, outside the submitted work; and personal fees from Janssen, outside the submitted work; and personal fees from Innocrin Pharmaceuticals, Novartis, and Cell Search, outside the submitted work; and personal fees from MaxiVax SA, outside the submitted work. CG reports grants from Clovis Oncology, outside the submitted work. NDJ reports advisory board fees from
from Sanofi and Novartis, outside the submitted work; and grants, personal fees, non-financial support, advisory board fees, speaker fees, and travel fees from Janssen, outside the submitted work. RJJ reports personal fees and non-financial support from Janssen, outside the submitted work; grants, personal fees, research funding, honoraria, speaker fees, and advisory board fees from Astellas, outside the submitted work; and personal fees and advisory board fees from Sanofi and Novartis, outside the submitted work. JFL reports personal fees, non-financial support, and travel fees from Janssen, outside the submitted work. ZIM reports personal fees, consultancy, advisory board fees, honoraria, and travel fees from Janssen and Sanofi, outside the submitted work; advisory board fees, honoraria, and travel fees from Astellas, outside the submitted work; and travel fees from Bayer, outside the submitted work. MDM reports personal fees, speaker fees, and advisory board fees from Sanofi, outside the submitted work; personal fees and speaker fees, outside the submitted work; and personal fees from Janssen and Bayer, outside the submitted work. JMO’s reports advisory board fees from Sanofi, outside the submitted work; personal fees, speaker fees, and advisory board fees from Janssen, outside the submitted work; and personal fees from Janssen and Bayer, outside the submitted work. CCP reports a research grant, personal fees, and advisory board fees from Bayer, outside the submitted work; advisory board fees from AAA, outside the submitted work; and speaker fees, advisory board fees, and research funding from Bayer, outside the submitted work. MKBP reports educational grants from Astellas, Clovis Oncology, Novartis, Pfizer, and Sanofi, outside the submitted work. DJS reports fees for meeting attendance from Astellas and Ipsen, outside the submitted work. NNS reports personal fees from Janssen, Pfizer, Sanofi, Merck Sharp & Dohme, and Roche, outside the submitted work. MRS reports grants and non-financial support from Astellas, Clovis Oncology, Novartis, Pfizer, and Sanofi, outside the submitted work; personal fees from Eli Lilly, outside the submitted work; and grants, personal fees and non-financial support from Janssen, outside the submitted work. AA, RA, CLA, ABA, CDB, OD, HD, CE, JG, MRG, APH, SJ, SK, REL, DM, RM, OAP, IDP, DMP, AWSR, GNT, ATTH, and JMR declare no competing interests.

Acknowledgments
This trial was funded by Cancer Research UK (CRUK/A12459), UK Medical Research Council (MRC_MC_UU_12023/25), and the Swiss Group for Clinical Cancer Research. Research support for the protocol was provided by Cancer Research UK (CRUK/A12459), UK Medical Research Council (MRC_MC_UU_12023/25), the Swiss Group for Clinical Cancer Research, Astellas, Clovis Oncology, Janssen, Novartis, Pfizer, and Sanofi-Aventis. This Article represents independent research part-funded by the National Institute for Health Research (NIHR) Biomedical Research Centre at the Royal Marsden NHS Foundation Trust and the Institute of Cancer Research. The views expressed in this Article are those of the authors and not necessarily those of the National Health Service, the NIHR, or the Department of Health. We thank Fiona Ingleby and Adrian Cook (Medical Research Council Clinical Trials Unit at University College London) for reviewing analyses.

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