ISRCTN56736625 https://doi.org/10.1186/ISRCTN56736625

A randomised comparison of ten versus three fractions of radiotherapy for palliation in patients unsuitable for radical radiotherapy or chemotherapy

Submission date
28/02/2001

Recruitment status
No longer recruiting

Registration date
28/02/2001

Overall study status
Completed

Last Edited
11/09/2007

Condition category
Cancer

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
BA09

Study information

Scientific Title

Study objectives
To compare the efficacy and side effects of two palliative radiotherapy schedules in patients with bladder cancer considered unsuitable for radical radiotherapy or chemotherapy.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Cancer

Interventions
Two palliative radiotherapy schedules in patients with bladder cancer considered unsuitable for radical radiotherapy or chemotherapy

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Change in bladder and bowel symptoms at three months, quality of life, response, survival.

Secondary outcome measures

Not provided at time of registration

Overall study start date
01/11/1992

Completion date
01/11/1997

Eligibility

Key inclusion criteria
Life expectancy of at least three months, no planned administration of cytotoxic chemotherapy

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
500

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/11/1992

Date of final enrolment
01/11/1997

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
MRC Clinical Trials Unit
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Medical Research Council (MRC) (UK)

Sponsor details
20 Park Crescent
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clinical.trial@headoffice.mrc.ac.uk

Sponsor type
Research council

Website
http://www.mrc.ac.uk

Funder(s)

Funder type
Research council

Funder Name
Medical Research Council (UK)

Alternative Name(s)
Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United Kingdom

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results:	01/05/2000		Yes	No