

Advance care planning in advanced cancer - can it be achieved? A patient preference trial of a care planning discussion

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
08/01/2007	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
16/02/2007	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
29/03/2022	Cancer	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Advance care planning in advanced cancer - can it be achieved? A patient preference trial of a care planning discussion

Study objectives

1. What is the acceptability and feasibility of a patient preference randomised controlled trial of an intervention to facilitate planning for end-of-life care?
2. Which outcomes are appropriate and measurable to assess the effectiveness of this intervention?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Royal Free Hospital and Medical School Local Research Ethics Committee. Approved on 17.08.06, ref 06/Q501/93

Study design

Randomised controlled patient preference trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Cancer

Interventions

The intervention will consist of a one-to-one discussion with a trained mediator using a checklist of topic domains. Discussions will take place usually in participants' homes and last no more than one hour. Participants will be offered up to two further sessions with the mediator, to take place at a time of their choice. Discussions will be audio-taped and transcribed to examine the range and development of topics covered, how these change with time, and to ensure that the advisor has covered each domain as appropriate.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Change in scores on visual analogue scales measuring ability to discuss end-of-life planning with primary and secondary care professionals and close persons/members of the family

Key secondary outcome(s)

1. Completion of an advance directive
2. Psychological status
3. Satisfaction with care

Completion date

07/07/2008

Eligibility

Key inclusion criteria

1. Have completed a primary course of treatment and still have clinically detectable, progressive, active disease
2. Are considered by the referring health professional to be well enough to complete the advance care planning discussion
3. Are over 18 years of age
4. Are able to give informed consent
5. Do not have psychotic illness
6. Speak and understand English (able to complete the questionnaires/interview)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Not Specified

Total final enrolment

77

Key exclusion criteria

1. Have not completed a primary course of treatment and still have clinically detectable, progressive, active disease
2. Are not considered by the referring health professional to be well enough to complete the advance care planning discussion
3. Are under 18 years of age
4. Are unable to give informed consent
5. Have psychotic illness
6. Do not speak and understand English (able to complete the questionnaires/interview)

Date of first enrolment

15/01/2007

Date of final enrolment

07/07/2008

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Marie Curie Palliative Care Research Unit

London

United Kingdom

NW3 2PF

Sponsor information

Organisation

University College London (UK)

ROR

<https://ror.org/02jx3x895>

Funder(s)

Funder type

Charity

Funder Name

Dimbleby Cancer Care (United Kingdom)

Results and Publications

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<u>Results article</u>	results	01/03/2011		Yes	No
<u>Participant information sheet</u>	Participant information sheet	11/11/2025	11/11/2025	No	Yes
<u>Plain English results</u>		20/07/2011	29/03/2022	No	Yes