

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

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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Louise Jones

Contact details
Marie Curie Palliative Care Research Unit
Royal Free and University College Medical School
Hampstead Campus
Rowland Hill Street
London
United Kingdom
NW3 2PF
-
l.jones@medsch.ucl.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Quality of life

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

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 measure

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

Secondary outcome measures

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

Overall study start date

15/01/2007

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

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Target number of participants

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**

England

United Kingdom

Study participating centre

Marie Curie Palliative Care Research Unit

London

United Kingdom

NW3 2PF

Sponsor information**Organisation**

University College London (UK)

Sponsor details

Gower Street

London

England

United Kingdom

WC1E 6BT

-
o.avwenagha@medsch.ucl.ac.uk

Sponsor type

University/education

Website

<http://www.ucl.ac.uk/>

ROR

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

Funder(s)

Funder type

Charity

Funder Name

Dimbleby Cancer Care (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**

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?
Results article	results	01/03/2011		Yes	No
Plain English results		20/07/2011	29/03/2022	No	Yes