ISRCTN43339432 https://doi.org/10.1186/ISRCTN43339432

Concern and continuity in the care of cancer patients and their carers: development of the intervention

Submission date
13/09/2005

Recruitment status
No longer recruiting

Registration date
17/10/2005

Overall study status
Completed

Last Edited
10/05/2012

Condition category
Cancer

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

Plain English summary of protocol
http://cancerhelp.cancerresearchuk.org/trials/a-study-looking-at-the-continuity-of-care-of-patients-with-cancer

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
SDO/13/2001 (e)

Study information

Scientific Title

Study objectives
That the development and testing of a complex intervention will lead to improved continuity of care in cancer for patients and those close to them by the delivery of the planned intervention.

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)
Other

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
Continuity of care in cancer

Interventions
A feasibility (phase II) randomised trial in which we are developing and testing a new intervention - this will be a form of care record for teams in which aspects of continuity of care for each patient will be assessed.
ECAP - engagement and continuity action plan: a continuity of care risk assessment completed by secondary care workers at multi-disciplinary meeting.

Intervention Type
Other

Phase
Phase II

Primary outcome measure
1. Acceptability
2. Practicality and feasibility of ECAP
3. Improved patient experience of engagement with care

Secondary outcome measures
1. Economic benefits
2. Patient and nominated friend/family member well being

Overall study start date
01/04/2006

Completion date
30/09/2007

Eligibility

Key inclusion criteria
1. Aged over 18 years
2. Able to give informed consent
3. Patients diagnosed with breast, lung or colorectal cancer, or their nominated relative or friend

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Both

Target number of participants
80 patients with up to two nominated relatives or friends

Key exclusion criteria
1. Unable to give informed consent
2. Aged under 18 years

Date of first enrolment
01/04/2006

Date of final enrolment
30/09/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Dept Mental Health Sciences
London
United Kingdom
NW3 2PF

Sponsor information

Organisation
University College London (UK)

Sponsor details
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Sponsor type
University/education

ROR
https://ror.org/02jx3x895

Funder(s)

Funder type
Government

Funder Name
NHS Service and Delivery Organisation (UK) [ref: SDO/13/2001 (e)]

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	27/01/2009		Yes	No