A study to examine the impact of doctor training in how to respond to patient reported symptoms and health related quality of life

Submission date	Recruitment status No longer recruiting	Prospectively registered		
14/06/2011		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
28/09/2011	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
26/10/2022	Cancer			

Plain English summary of protocol

https://www.cancerresearchuk.org/about-cancer/find-a-clinical-trial/a-study-looking-impact-doctor-training-how-to-respond-to-patient-questionnaire-about-quality-of-life-QuEST-T2

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

C7775/A7424

Study information

Scientific Title

A pilot quasi-experimental study to examine the impact of doctor training in how to respond to patient reported symptoms and health related quality of life

Study objectives

Training oncologists to use HRQoL measures during routine chemotherapy review consultations will improve patient-centred communication. The data from this pilot study will provide an estimate of the impact of oncologist training and contribute to sample size calculations for a future randomised controlled trial.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Leeds Central Reseach Ethics Committee, 10/02/2011 ref: 11/H1313/2

Study design

Pilot Quasi-experimental study

Primary study design

Interventional

Secondary study design

Non randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Screening

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

Advanced stage breast, colorectalcancer, gynaecological cancers

Interventions

The study utilises a quasi-experimental design (before and after). 30 patients will be invited to take part before the participating oncologists attend the training programme and further 30 patients will be recruited after the training. Patients will be asked to complete a health related quality of life (HRQoL) questionnaire prior to routine chemotherapy review consultations. The participating oncologists (n = 3) will receive the results of the questionnaire and asked to incorporate these into the consultations. The before after study design will be used to assess

the impact of a doctor training session to aid the facilitation of the HRQoL questionnaire into consultations.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Doctor-patient communication before and after the doctor training will be compared by:

- 1. Analysing audio-recordings of the consultations using content analysis and the measure of patient centred communication to determine the number of issues raised and the quality of discussions
- 2. Patient and oncologists views will be assessed using the Patient Perception of Patient Centredness questionnaire and the corresponding measure for doctors

Secondary outcome measures

Interviews with participating oncologists at the end of the study will explore their experience of the intervention before and after the training and opinions of the training programme to obtain suggestions for improving its content.

Overall study start date

04/07/2011

Completion date

28/10/2011

Eligibility

Key inclusion criteria

- 1. Are attending oncology clinics at St James' Institute of Oncology
- 2. Are receiving anti cancer chemotherapy for advanced stage of disease (breast, colorectal and gynaecological cancer)
- 3. Are able to give informed consent
- 4. Are able to read and understand English

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

60

Total final enrolment

Key exclusion criteria

- 1. Are below 18 years of age
- 2. Show overt exhibition of psychopathology or serious cognitive dysfunction which would impede them being able to take part in the study
- 3. Are deemed too ill by the oncology staff

Date of first enrolment

04/07/2011

Date of final enrolment

28/10/2011

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Level 4, Bexley Wing

Leeds United Kingdom LS9 7TF

Sponsor information

Organisation

University of Leeds (UK)

Sponsor details

School of Medicine
The Worsley Medical & Dental Building
Clarendon Way
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England
United Kingdom
LS2 9NL
+44 113 343 0900
R.E.DeSouza@leeds.ac.uk

Sponsor type

University/education

Website

http://www.leeds.ac.uk/

ROR

https://ror.org/024mrxd33

Funder(s)

Funder type

Charity

Funder Name

Cancer Research UK (CRUK) (UK) ref: C7775/A7424

Alternative Name(s)

CR_UK, Cancer Research UK - London, CRUK

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

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

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	01/07/2015		Yes	No
Plain English results			26/10/2022	No	Yes