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Helping senior doctors involved in cancer care improve their ability to break bad news, elicit and respond to patients' concerns

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 01/10/2012	**Condition category** Cancer	[_] Individual participant data [_] Record updated in last year

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

N0063044242

Study information

Scientific Title

Study objectives
To determine if training consultants and senior registrars specialising in cancer in breaking bad news, eliciting and responding to patients' concerns.

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

Health condition(s) or problem(s) studied
Miscellaneous cancers

Interventions
Randomised controlled trial of training consultants and senior registrars specialising in cancer in breaking bad news, eliciting and responding to patients' concerns.
Arm A = Training, Arm B = without Training.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Doctors: Objective ratings of recorded interviews with 2 simulated patients before training, after training and 3 months later. Objective ratings via radiomicrophone recordings of 10 consultations when breaking bad news before and after training. Subjective: Maslach Burnout Inventory, self efficacy, outcome expectancy before and after training and 3 months later.

Patients:
1. Pre-consultation: Hospital Anxiety and Depression Scale (HADS), Spielberger state anxiety
2. Post-consultation: Spielberger, concerns checklist
3. 1 and 3 months later: concerns checklist and HADS by post

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/1997

Completion date
31/12/2008

Eligibility

Key inclusion criteria
NHS Doctors

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/06/1997

Date of final enrolment
31/12/2008

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
CRC Psychological Medicine Group
Manchester
United Kingdom
M20 4BX

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Christie Hospital NHS Trust (UK)

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