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Improving communication during recruitment to clinical trials of cancer therapy

Submission date 23/01/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 23/01/2004	**Overall study status** Completed	
Last Edited 23/05/2012	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
NCP/D21

Study information

Scientific Title

Study objectives
Added 06/08/09:
Specific details about an individual patient's attitudes to clinical trials will be elicited so that the development and validation of the questionnaire will be completed. When the instrument has been refined satisfactorily it is hoped that clinicians will be provided with an easy means of identifying the potential areas of difficulty that individual patients might have with the concentp of trials and/or randomisation.

As of 06/08/09 this record has been extensively updated. All updates can be found under the relavent field with the above update date. Please also note that the start and end dates of this trial have been changed from 01/01/99 and 31/01/99 to 18/10/96 and 25/02/97, respectively. The initial dates were generated by the system.

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)
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
Miscellaneous cancers

Interventions
1. Standardised trial consent procedure.
2. Individualised trial consent procedure using a 'Profile of Patient Preferences'.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Added 06/08/09:
Quality of life

Secondary outcome measures
Not provided at time of registration

Overall study start date
18/10/1996

Completion date
25/02/1997

Eligibility

Key inclusion criteria
Patients aged over 16 with cancer; medical, clinical and surgical oncologists.

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
18/10/1996

Date of final enrolment
25/02/1997

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
CRC Psychosocial Oncology Group
London
United Kingdom
W1P 7PL

Sponsor information

Organisation
Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Cancer National Research and Development Programme (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/02/2001		Yes	No