

# Improving communication during recruitment to clinical trials of cancer therapy

<b>Submission date</b> 23/01/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 23/01/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 23/05/2012	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data

**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**  
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 concept of trials and/or randomisation.

As of 06/08/09 this record has been extensively updated. All updates can be found under the relevant 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?
<a href="#">Results article</a>	results	01/02/2001		Yes	No