

# Improving communication during recruitment to clinical trials of cancer therapy

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

**Protocol serial number**  
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

**Study type(s)**

Quality of life

**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(s)**

Added 06/08/09:

Quality of life

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

25/02/1997

# Eligibility

## Key inclusion criteria

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

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

All

## 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

United Kingdom

England

## 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)

# Funder(s)

## Funder type

Government

## Funder Name

NHS Cancer National Research and Development Programme (UK)

# Results and Publications

## 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              |
| <a href="#">Participant information sheet</a> | Participant information sheet | 11/11/2025   | 11/11/2025 | No             | Yes             |