

A randomised controlled trial of occupational therapy in oncology: a pilot study

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 24/05/2012	Condition category 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
N0192119065

Study information

Scientific Title

Study objectives

To identify whether occupational therapy intervention improves the mood, fatigue management and activities of daily living (ADL) performance of outpatient oncology patients.

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)

Not Specified

Health condition(s) or problem(s) studied

Cancer: All

Interventions

Intervention arms: Patients will receive a minimum of five sessions.

Session 1 - initial interview, general assessment and baseline assessments.

Session 2 - fatigue management/energy conservation session and equipment provision.

Session 3 - anxiety management/relaxation session.

Session 4 -fatigue/anxiety management session - check understanding and use of principals.

Session 5 - review progress.

Further sessions will be conducted if considered appropriate, up to a maximum of seven.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

We aim to follow up two groups of patients at 6 and 12 weeks after randomisation and compare outcome.

Key secondary outcome(s))

Not provided at time of registration

Completion date

09/05/2003

Eligibility**Key inclusion criteria**

All patients will be seen on their second or third clinic appointment.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Living outside the hospital catchment area or refusal to consent.

Date of first enrolment

17/09/2002

Date of final enrolment

09/05/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Ageing and Disability Research Unit

Nottingham

United Kingdom

NG7 2UH

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Queen's Medical Centre University Hospital NHS Trust - UK (NHS R&D Support Funding)

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?
Results article	results	01/03/2006		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes