

A pragmatic, randomised controlled trial of reflexology, relaxation and guided imagery, and self-initiated support in patients with lung cancer

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/10/2012	Condition category Cancer	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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

N0084122777

Study information

Scientific Title

Study objectives

Do reflexology, relaxation and guided imagery enhance Quality of Life in men and women with lung cancer, and what are their psychoneuroimmunological effects?

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)

Other

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

Lung cancer

Interventions

RCT with 3 arms:

1. Self initiated support in the Oncology Health Centre
2. Arm 1 plus relaxation and guided imagery
3. Arm 1 plus reflexology

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Added 11/08/09:
Quality of life

Secondary outcome measures

Not provided at time of registration

Overall study start date

06/03/2003

Completion date

13/01/2006

Reason abandoned (if study stopped)

Lack of funding

Eligibility

Key inclusion criteria

180 patients will be recruited.

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

180

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

06/03/2003

Date of final enrolment

13/01/2006

Locations

Countries of recruitment

United Kingdom

Study participating centre
The Institute of Rehabilitation
Hull
United Kingdom
HU3 2 PG

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
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
Oncology Health Centre (UK)

Funder Name
The North and South Bank Research and Development Consortium (UK) (NHS R&D Support Funding)

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