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Multicentre evaluation of a nursing clinic for dyspnoea in patients with cancer of the lung

Submission date 19/08/2002	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 19/08/2002	**Overall study status** Completed	
Last Edited 26/03/2020	**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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Contact details
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
MPDU LU1

Study information

Scientific Title
Multicentre evaluation of a nursing clinic for dyspnoea in patients with cancer of the lung

Study objectives
Not provided at time of registration

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Lung (non-small cell), Lung (small cell)

Interventions
1. Control Group: Best supportive care plus breathing assessment
2. Intervention Group: Best supportive care, breathing assessment, breathing retraining and individual psychosocial support

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date

01/01/2000

Completion date
31/12/2000

Eligibility

Key inclusion criteria
1. Either sex any age
2. Small cell, non-small cell or mesothelioma of the lung
3. Completed first line therapy for the disease (surgery and/or chemotherapy and/or radiotherapy)
4. Change in breathing or a degree of breathlessness
5. Patients with metastatic disease are not excluded

Participant type(s)
Patient

Age group
Not Specified

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Patients who have radiotherapy, chemotherapy or surgery planned for the 8 weeks that are required for the study, or who have had radiotherapy, chemotherapy or surgery in the last 3 weeks are excluded

Date of first enrolment
01/01/2000

Date of final enrolment
31/12/2000

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
UKCCCR Register Co-ordinator
London

United Kingdom
NW1 2DA

Sponsor information

Organisation
Macmillan Cancer Relief (UK)

Sponsor details
89 Albert Embankment
London
United Kingdom
SE1 7YQ

Sponsor type
Charity

Website
http://www.macmillan.org.uk

ROR
https://ror.org/05vfhev56

Funder(s)

Funder type
Charity

Funder Name
Macmillan Cancer Relief (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	03/04/1999	26/03/2020	Yes	No