ISRCTN13391637 https://doi.org/10.1186/ISRCTN13391637

To evaluate the effects of Manual Lymphatic Drainage on limb volume and quality of life when used in addition to standard lymphoedema therapy.

Submission date
12/09/2003

Registration date
12/09/2003

Last Edited
18/08/2015

Recruitment status
No longer recruiting

Overall study status
Completed

Condition category
Circulatory System

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[_] Results
[_] Individual participant data
[_] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mrs Gloria Shingler

Contact details
Breast Cancer Specialist Nurse Counsellor
Bedford Hospital NHS Trust
South Wing
Kempston
Bedford
United Kingdom
MK42 9DJ
+44 (0)1234 355122 ext 2041
abc@email.com

Additional identifiers

Protocol serial number
N0040122626

Study information

Scientific Title
To evaluate the effects of Manual Lymphatic Drainage on limb volume and quality of life when used in addition to standard lymphoedema therapy.

Study objectives
To evaluate quality of life of patients with upper limb lymphoedema secondary to breast cancer, when Manual Lympathetic Drainage message is added to the standard Lymphoedema therapy using a quantitative validated questionnaire SF36

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)
Treatment

Health condition(s) or problem(s) studied
Cardiovascular: Lymphoedema

Interventions
Manual Lymphatic Drainage and standard lymphoedema therapy vs standard lymphoedema therapy only

Intervention Type
Procedure/Surgery

Primary outcome(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

Completion date
15/04/2003

Eligibility

Key inclusion criteria
A clinic list of 120 patients with 3-6 monthly appointments

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/02/2003

Date of final enrolment
15/04/2003

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Bedford Hospital NHS Trust
Bedford
United Kingdom
MK42 9DJ

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
Bedford Hospital NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration