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 | Recruitment status | Prospectively registered |
|-------------------|----------------------|-----------------------------------------------|
| 12/09/2003 | No longer recruiting | Protocol |
| Registration date | Overall study status | Statistical analysis plan |
| 12/09/2003 | Completed | Results |
| Last Edited | Condition category | [] Individual participant data |
| 18/08/2015 | Circulatory System | 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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

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

Cardiovascular: Lymphoedema

Interventions

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

Intervention Type

Procedure/Surgery

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/02/2003

Completion date

15/04/2003

Eligibility

Key inclusion criteria

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

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

120

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

England

United Kingdom

Study participating centre

Bedford Hospital NHS Trust

Bedford United Kingdom MK42 9DJ

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

Bedford Hospital NHS Trust (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