

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	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 18/08/2015	Condition category Circulatory System	☐ 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

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Contact details

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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 Lymphatic 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

Primary study design

Interventional

Study design

Randomised controlled trial

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