# 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	<ul><li>Prospectively registered</li></ul>
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	<ul><li>Record updated in last year</li></ul>

## 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

## 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

**Study outputs** 

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet
Participant information sheet
11/11/2025 No Yes