

Lymph node transfer for breast cancer related lymphoedema

Submission date 13/12/2017	Recruitment status Recruiting	☐ Prospectively registered ☐ Protocol
Registration date 12/02/2018	Overall study status Ongoing	☐ Statistical analysis plan ☐ Results
Last Edited 12/12/2024	Condition category Circulatory System	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol

Background and study aims

Lymphedema is a collection of fluid that causes swelling in the arms and legs. Lymphedema of the arm is often due to removal of the lymph glands in the armpit (axillary clearance) during breast cancer treatment. Every year in Sweden more than 8000 women are diagnosed with breast cancer of which more than half undergo surgery of their lymph glands in addition to their breast surgery. Today, about 5000-6000 people in Sweden suffer from lymphedema of the arm due to breast cancer treatment, with around 800 new cases diagnosed annually. Lymphedema of the arm can lead to disabilities like limited range of motion, a feeling of heaviness or pressure, pain, increased infection risks, difficulty wearing clothes, poor cosmetic appearance and psychological effects. Lymphedema can be treated in different ways. Conservative treatment entails different forms of compression and manual therapy. In addition to this, general recommendations include weight loss and range of motion (physiotherapy) training. Surgical options have changed from offering the patient resection surgery (where large portions of the swelling is removed) to offering either liposuction or microsurgical methods. The goal for these microsurgical methods is to reduce the lymphedema by improving the function of the lymphatic system. Two principal methods are used today: autologous lymph node transplantation (ALNT) and lymphatic-venous anastomoses (LVA). In both of these methods superficial veins and lymph nodes are anastomosed (connected) with the aid of a microscope which is why they are called microsurgical. The aim of this study is to assess the effect of lymph node transfer in patients with lymphedema of the arm after breast cancer treatment.

Who can participate?

Women eligible for breast reconstruction who have lymphedema of the arm on the same side as the breast reconstruction

What does the study involve?

Participants are randomly allocated to undergo breast reconstruction and tissue transfer to the armpit either with or without lymph nodes. Arm volume, patient satisfaction and lymph flow are assessed before and 3, 6, 12 and 24 months after surgery.

What are the possible benefits and risks of participating?

The outcome of the different treatments is uncertain. The transferred lymph nodes are taken

from the groin and in order to prevent donor-site problems a lymphoscintigraphy scan is performed before the operation.

Where is the study run from?

1. Uppsala University Hospital (Sweden)
2. Maastricht University (Netherlands)

When is the study starting and how long is it expected to run for?

May 2017 to May 2027

Who is funding the study?

1. Uppsala-Örebro Regionen (Sweden)
2. Uppsala University Hospital (Sweden)

Who is the main contact?

1. Mrs Åsa Wiberg (public)
asa.wiberg@akademiska.se
2. Dr Maria Mani (scientific)
maria.rydevik.mani@akademiska.se

Contact information

Type(s)

Public

Contact name

Mrs Åsa Wiberg

Contact details

Department of Plastic and Reconstructive Surgery
Uppsala University Hospital
Uppsala
Sweden
75185
+46 (0)186115427
asa.wiberg@akademiska.se

Type(s)

Scientific

Contact name

Dr Maria Mani

Contact details

Department of Plastic and Reconstructive Surgery
Uppsala University Hospital
Uppsala
Sweden
75185
+46 (0)186110471
maria.rydevik.mani@akademiska.se

Additional identifiers

Protocol serial number

LN001

Study information

Scientific Title

Randomised trial on lymph node transfer for treatment of breast cancer related lymphoedema

Acronym

LyNT

Study objectives

Hypothesis:

Microsurgical techniques can contribute to reducing lymphedema and effect patient symptoms in a positive manner. Lymph therapeutic treatment can improve the effect of surgical treatment and should be seen as an adjunct treatment in a standardized protocol.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Regional Ethical Board, Uppsala Sweden, 08/02/2017, ref: Dnr 2016/470

Primary study design

Interventional

Study design

Prospective single-blinded randomised controlled study

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Lymphedema of the arm secondary to breast cancer treatment

Interventions

Lymphnodetransfer as a complement to scar release and free flap (DIEP) reconstruction of the breast

Patients with lymphedema referred to the clinic are informed of the present study. If they give their consent to participate they are included and follow the usual treatment plan for microsurgical reconstructive surgery with either lymphatic-venous anastomoses (LVA) or autologous lymph node transplantation (ALNT) according to the method that is the most optimal for each individual according to the pre-operative investigations. If they are candidates for breast reconstruction with ALNT they will be randomized into 1 of 2 groups:

1. Breast reconstruction with tissue from their abdomen combined with fibrotic tissue release in the armpit and tissue transfer to the armpit
2. Breast reconstruction with tissue from the abdomen and combined lymph node transfer

Pre-operative investigations are performed according to routine clinical practice and postoperative follow up follows the study protocol (appendix 12). Study protocol is filled out during clinical visits and supplementary information is obtained from patient journals and the quality registry when needed (appendix 13). Data is gathered and processed without patient identification and follows routine guidelines for data and registry handling.

Investigations include history, clinical examination and photo documentation, volume measurement using measuring tape, water displacement test, microwave analysis (measure water content in tissue and describes the inner tissue composition) and 3D measurement. Lymph flow is measured using magnetic resonant imaging, lymphoscintigraphy and Photodynamic Eye (see below) (Ogata et al 2007, Mihara et al). Participants will also answer several questionnaires (appendix 5 a-x). Assessment will be done pre-operatively and 1, 6, 12, 24 and 36 months post-operatively.

Surgery is performed according to methods described below:

1. Lymph node transfer means lymph nodes are transferred with their nourishing vessels to the area affected by lymphedema. The vessels are anastomosed using so called microsurgical technique. The lymph nodes can be part of the tissue/flap that is moved from the abdomen during breast reconstruction with a DIEP flap or as its own flap (Becker et al 2006). A perioperative lymphoscintigraphy combined with Photodynamic Eye (PDE) is performed in order to ensure that the lymph nodes that are transferred aren't the lymph nodes draining the inferior limb and cause iatrogenic lymphedema (Ogata et al 2007, Mihara et al). Lymphoscintigraphy is performed for the leg and fluorescens marking from the abdomen. The lymph nodes that signal in the scintigraphy detector are not included in the flap (Dayan et al 2015). In order to distinguish the effects of the lymph node transfer from the fibrotic tissue release the lymph node transfer group is randomized to only fibrotic tissue release and flap or fibrotic tissue release and flap + lymph node transfer.
2. Lymphaticovenous anastomosis (LVA) is performed by identifying superficial lymph channels and vessels on the affected extremity. These anastomoses are done under a microscope. With the aid of pre-operative markings and peri-operative investigations with PDE etc the vessels to be anastomosed are identified. Usually 2-5 anastomoses are done in one extremity (Koshima et al 2000)

Intervention Type

Procedure/Surgery

Primary outcome(s)

Measured at baseline (preoperatively), 3, 6 and 12 months:

1. Volume of the extremity, measured using water displacement test, 3D camera, tape measurement
2. Patient satisfaction, measured using LyQli and ICL- lymph questionnaire

There will also be a long term follow-up at 24 months (not included in the study endpoint)

Key secondary outcome(s)

Lymph flow, measured using magnetic resonant imaging and scintigraphy at baseline (preoperatively), 3, 6 and 12 months

There will also be a long term follow-up at 24 months (not included in the study endpoint)

Completion date

01/05/2027

Eligibility

Key inclusion criteria

Patients eligible to breast reconstruction with a DIEP and concomitant lymphedema of the arm

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Same criteria as for the DIEP reconstruction

Date of first enrolment

05/05/2017

Date of final enrolment

31/12/2026

Locations

Countries of recruitment

Netherlands

Sweden

Study participating centre

Uppsala University Hospital

Department of Plastic and Reconstructive Surgery

Uppsala

Sweden

75185

Study participating centre

Maastricht University

Maastricht

Netherlands
6211 LK

Sponsor information

Organisation

Uppsala University Hospital

ROR

<https://ror.org/01apvbh93>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Uppsala-Örebro Regionen

Funder Name

Akademiska Sjukhuset

Alternative Name(s)

Uppsala University Hospital

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

Sweden

Results and Publications

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary

Data sharing statement to be made available at a later date