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Patient Information

Name _____

DOB _____

Medical Necessity Information

1. **Diagnosis:** Primary Lymphedema (Congenital) Q82.0 Post-mastectomy Lymphedema I97.2
 Acquired Lymphedema I89.0 Secondary due to surgery I97.89

2. **Location of Lymphedema:**

	Swelling	Fibrosis	Pain		Swelling	Fibrosis	Pain		Swelling	Fibrosis	Pain
Feet	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Right Upper Leg	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Left Arm	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Left Upper Leg	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Right Lower Leg	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Right Arm	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Left Lower Leg	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Underarm/Axilla	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Hand(s)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

3. **Patient is currently experiencing the following related complications/impairments:**

- Unable to control swelling
 Fibrosis
 Pain
 Impaired ROM
 Compromised skin integrity
 Impaired mobility
 Contractures
 Scarring
 Infections

4. **What home treatments has patient been performing for the past 4 weeks?**

	Yes	No		Yes	No	
Compression garments and bandaging	<input type="checkbox"/>	<input type="checkbox"/>	Exercise	<input type="checkbox"/>	<input type="checkbox"/>	Comments: _____ _____ _____
Manual Lymph Drainage	<input type="checkbox"/>	<input type="checkbox"/>	Elevation	<input type="checkbox"/>	<input type="checkbox"/>	

Patient has the following barriers to using compression garments:

- None
 Adverse reaction
 Unable to don/doff
 Unable to afford
 Unable to accommodate size

Patient tried and failed* home treatments for at least 4 weeks? Yes No **Failure defined as significant symptoms remain or no significant improvement.*

5. **Patient tried and failed a basic pump?** *Basic pump defined as uncalibrated segmental pressure pump (E0651)* Yes No

If yes, after trials with the basic pump, the patient was unable to tolerate due to:

- Exacerbated symptoms
 Compromised skin integrity
 Pain
 Unable to accommodate size
 No clinical improvement

Name/Signature of Person Answering Questions (if other than physician)

Name _____

Signature _____

Date _____

Physician Certification and Signature

I certify that this patient is under my care and the above medical necessity information is true and accurate to the best of my knowledge.

Physician Name _____

Signature _____

Date _____

*Please refer to this guide when completing the
Additional Medical History Form — Lymphedema.
The requirements listed below are necessary
to obtain coverage by most insurers.*

Lymphedema

1. Diagnosis

The patient must have a diagnosis of Lymphedema which has failed* to resolve after a 4-week trial of conservative therapy. The types of Lymphedema approved for coverage by most commercial insurers are Congenital Lymphedema, Post-mastectomy Lymphedema, and Acquired Lymphedema Secondary to Surgery or Cancer.

**Failure is defined as significant symptoms remain or no significant improvement.*

2. Location of Lymphedema

At least one part of lower extremity (ie: upper/lower/foot) **OR** upper extremity (ie: upper/lower/hand/underarm/axilla), needs to have swelling, fibrosis, and/or pain documented.

3. Complications/Impairments

Document any of the complications that your patient is experiencing due to Lymphedema.

4. Home Treatments

Patients must have tried and failed* conservative therapy** for a minimum of 4 weeks.

**Failure is defined as significant symptoms remain or no significant improvement.*

***Conservative therapies must include: either compression garments or compression bandaging, exercise, elevation, and if appropriate, MLD.*

- Exercise includes ambulation and does not have to be a formal exercise program.

5. Basic pump

Some insurers require a treatment trial with a “basic pump” (an E0651) before they will cover an E0652 device. If your patient has tried and failed* a basic pump, please document that here.

**Failure is defined as significant symptoms remain or no significant improvement.*