

## **Medical Standards & Research Pre-approval Form**

This is a pre-requisite form provided upon request for the "Pre-Authorization Form for Bundled Lymphedema and Lipedema serivces for Thiqa".

Kindly fill in all the requested information given below. This is a mandatory step to proceed further. Failure to provide information relevant to approval will delay the processing of the applicant's request. The provider will be contacted in case further clarifications are required.

GE	GENERAL INFORMATION						
-	Member's Name:						
	☐ New ☐ Established						
-	Member Card #:						
-	Policy:						
-	Age:						
-	Gender:						
-	Date: / /						
D.D.							
PR	OVIDER INFORMATION						
-	Provider's Name:						
-	Ordering Clinician (ID # & Name):						
-	Performing Provider Name:						
-	Performing Clinician Speciality (ID # & Name):						
-	Referring Physician (ID # & Name):						
0.7							
SE	RVICE REQUESTED						
-	Principal/ Primary Diagnosis:						
-	ICD-10:						
-	Requested Procedural Service Code (SRVC):						
-	BMI:						
Clir	nical Assessment & Condition Type						
Cili	near Assessment a condition Type						
	☐ Lymphedema						
	□ Lipedema						
	□ Both						



	Onset and Duration				
Staging/Severity		Lymphedema			
	(ISL for Lymphedema / Clinical for Lipedema)	□ Stage I (Mild) □ Stage II (Moderate) □ Stage III (Severe)  Lipedema □ Type : (1-5) □ Stage: 1-4 (smooth/nodular/indurated skin)			
	Symptoms	□ Pain □ Swelling □ Bruising □ Fatigue □ Recurrent Infections □ Mobility Impairment □ Other:			
	Family History (for Lipedema)	□Yes □ No Details:			
Comorbidities		□ Obesity (BMI 30-34.9) □ Obesity (BMI >35) □ Diabetes □ Venous Insufficiency			
		□ □ Other:			
	Differential Diagnoses Ruled Out	□Yes □ No (e.g., Venous Disease)			
Quantitative Measures (lymphedema)		□ Volumetry differential (circumferential measurements and/or Perometry differential)>10% (if affected extremity dominant extremity) or >7% (affected extremity is nondominant extremity) OR			
		□ Circumference Difference ≥2 cm at one or more standardized measurement points (e.gthigh, knee, mid-calf, or ankle).			
		□ Bioimpedance (L-Dex) differential of at least 10 units			
		□ Body Water Balance Index (BWBI) ECW/TBW Ratio > 0.39			
	Quantitative Measures (lipedema)	Symmetrical and bilateral disproportionate fat distribution, often sparing the hands and feet, 8 leading to a characteristic (column-like) or (stovepipe) known as (cuff sign) or (Fat pad sign) appearance.			
		□ Chronic pain or discomfort, exacerbated by touch or pressure interfering in everyday activity.			
		□ Easy Bruising or Spontaneous Bruising with minimal or no trauma, due to the fragile nature of the capillaries in lipedema fat tissue.			
		□ Firm or nodular subcutaneous tissue with distinct "dimpled" texture or localized fibrosis or increased resistance compared to unaffected tissue.			
		□Stemmer's sign, non-pitting or minimally pitting edema.			
		□ Recurrent cellulitis and Skin ulceration.			

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	□Body Water Balance Index (BWBI) ECW/TBW Ratio (≤0.39).	
Exclusion-Presence of one of the	□ Venous disease (DVT, superior vena cava syndrome).	
following:	□ Congestive heart failure (CHF).	
	□ Medication-induced swelling.	
	□ Liver disease includes but is not limited to cirrhosis and hypoproteinemia.	
	□ Nephropathy includes end-stage renal disease.	
	□ Pregnancy.	
	□ Dye anaphylaxis.	
	□ Active infection of the affected extremity (cellulitis/erysipelas).	
	□ Active Cancer status	
	□ Morbid obesity	

DIAGNOSTIC TESTS (ATTACH REPORTS)
□ Lymphoscintigraphy/ICG Lymphography (for Lymphedema)
□ Ultrasound/Duplex (for Lipedema assessment)
□ Limb Circumference Measurements (Pre- and Post-Conservative)
□ MRI Findings
□ Photography
□ Other:

COMPLETE DECONGESTIVE THERAPY HISTORY (CDT) (REQUIRED PREREQUISITE)					
Therapy Type	Duration (Months)	Frequency	Outcome/Reason for Failure	Provider Name/Date	
Complete Decongestive Therapy (CDT: Manual Lymphatic Drainage + Compression)					
Exercise/Graded Activity Program					
Weight Management (if BMI >25)					
Compression Therapy					

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Psychosocial Support		
Other (e.g. Proumatic		
Other (e.g., Pneumatic Compression)		

SURGICAL INTERVENTION				
Primary Site/Limb Treated	Specify:			
Primary Limb Procedures Code ( <b>Include all</b> <b>services codes</b> )	□ 59-01 □ 59-02 □ 59-03 □ 59-04 □ 59-05 □ 59-06* □ 59-07 □ 59-08 □ 59-09			
Additional Site(s)/Limb(s) Treated	Specify:			
Additional Same Procedure Codes/Specify Quantity	☐ 59-01 QTY: ☐ 59-02 QTY: ☐ 59-03 QTY: ☐ 59-04 QTY: ☐ 59-05 QTY: ☐ 59-06* QTY: ☐ 59-07 QTY: ☐ 59-08 QTY: ☐ 59-09 QTY: ☐ 59-010 QTY:			
Additional Different Procedure Codes/ <b>Specify Quantity</b>	☐ 59-01 QTY: ☐ 59-02 QTY: ☐ 59-03 QTY: ☐ 59-04 QTY: ☐ 59-05 QTY: ☐ 59-06* QTY: ☐ 59-07 QTY: ☐ 59-08 QTY: ☐ 59-09 QTY: ☐ 59-10 QTY:			
Short Description				
Complexity (if 59-06-xx)*	□ 01 Low □ 02 Medium □ 03 High  Justification: (one or multiple complexity)			
Multiple Techniques	□ Yes □ No			
Bio-Bridge Implant (for 59-01/59-02)	☐ Yes ☐ No (Attach original invoice if applicable)			

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Expected Surgical Duration	Hours			
Anticipated Inpatient Stay	Days			
Post-Op Plan				
Risks/Expected Outcomes				
Supporting Documents	Checklist (Attach /	All)		
☐ Clinical notes/history/exam				
☐ Diagnostic Imaging/reports				
□ CDT records				
□ Patient Consent				
Surgeon Certification				
I certify that the proposed intervention meets medical necessity criteria per DOH standards and protocols, with conservative therapy failure documented. The bundle includes all pre-op, intra-op, and post-op services (excluding Bio-Bridge implant).				
Surgeon Signature	Date	Printed Name	DOH License #	

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