

PNEUMATIC COMPRESSION DEVICE ORDER FORM

FAX ORDER TO 1.866.435.3949

REQUIRED ATTACHMENTS: PATIENT DEMOGRAPHICS/FACE SHEET AND COPY OF INSURANCE CARD

FACILITY INFORM	IATION									
FACILITY NAME: CONTACT:					PHONE:	PHONE:			EMAIL:	
			FAX							
REFERRAL SUBMITTED BY (if different from Facility Contact above or Prescriber below):										
PATIENT INFORM	ATION									
FIRST NAME: MIDDLE INITIAL: LAST NAME:					DATE OF BIRTH: (mm/dd/yy) / /		MED	MEDICARE ID (IF APPLICABLE):		
ADDRESS:			CITY:		STATE:	ZIP:		PHONE:		
LOCATION OF EDEMA:										
PRODUCT ORDER										
ENTRE® System (Basic Pump E0651) COMPLETE ALL SECTIONS BELOW					FLEXITOUCH® Plus (Advanced Pump E0652) STOP! FAX MEDICAL RECORDS AND THIS FORM (SKIP BELOW SECTIONS)					
PATIENT MEASUREMENTS FOR GARMENT SIZING										
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A BECEPS 	ARM	Left (cm)	Right (d	cm)		LEG	Lef	t (cm)	Right (cm)	
	Inseam				/	Inseam				
	Biceps				CALF	Thigh				
	Forearm					Calf				
					21-1 - 65					
ALL SECTIONS BELOW MUST BE COMPLETED BY A HEALTHCARE PROVIDER										
SECTION A: DIAGNOSIS INFORMATION										
 I89.0 Secondary Lymphedema due to										
SECTION B: MEDICAL NECESSITY AND COVERAGE CRITERIA INFORMATION										
ALL QUESTIONS MUST BE ANSWERED 1. □ Yes □ No Has patient tried and failed home treatments (appropriate compression garments/exercise/elevation/wound dressings, as appropriate) for at least 4-weeks (or 6 months for VLUs) and significant symptoms remain or with no significant improvement? 2. □ Yes □ No Have measurements been documented in the patient's medical record that confirm the persistence of lymphedema? 3. □ Yes □ No Is patient CURRENTLY experiencing any related complications/impairments/persisting symptoms? Check all that apply: □ Hyperkeratosis □ Hyperpigmentation □ Papillomatosis (warts, nodules, papules) □ Cellulitis 4. Date of last face to face encounter with prescriber (mm/dd/yy)://*Medicare requires a visit within the past 6 months.										
RX: PNEUMATIC COMPRESSION DEVICE AND GARMENTS										
DEVICE AND GARM		TION								
ENTRE System (E0651) ARM (E0668) Left Right FULL LEG (E0667) Left Right HALF LEG (E0669) Left Right										
TREATMENT PROTOCOL										
Duration per Extremity (hour):Frequency per Day: \Box 1 \Box 2 \Box Other: \Box 1x \Box 2x \Box Other:								_ength of Need (choose one):		
PRESCRIBER'S ORE	DER AND AT	TESTATION								
I am the treating physician or practitioner for the above-named patient. I have examined the patient, maintained oversight of their condition throughout treatment, and have determined that the patient has a medical necessity for a pneumatic compression device. I have received the list of contraindications listed in the User Guide, and the patient has no contraindications that would prohibit use of the prescribed equipment. The patient's medical record contains documentation showing the patient meets coverage criteria for a pneumatic compression device in accordance with applicable Medicare and other third-party payer coverage policies as indicated above. I will make such medical records available to Tactile Medical and third-party payer(s) upon request.										
PRESCRIBER NAME:			PRESCRIBER SIGNATURE:			DATE:			NPI:	
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TACTILE SYSTEMS TECHNOLOGY INC. | 3701 WAYZATA BLVD., SUITE 300, MINNEAPOLIS, MN 55416 | NPI: 1427131424 If you have questions regarding your patient's clinical case and Medicare criteria, please call our Medicare Hotline 1-855-319-9949