



# E-STIM ORDER FORM

Commercial — Medicare — Medicaid — TRICARE — VA — Uninsured

## SECTION 1 — PATIENT & INSURANCE INFORMATION

Patient Name \_\_\_\_\_ DOB (MM/DD/YY) \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_ Gender \_\_\_\_\_  
Shipping Address \_\_\_\_\_ City \_\_\_\_\_ State \_\_\_\_\_ ZIP \_\_\_\_\_  
Primary Phone Number \_\_\_\_\_ Secondary Phone Number \_\_\_\_\_  
Email \_\_\_\_\_ Primary Language, if not English \_\_\_\_\_  
Primary Insurance \_\_\_\_\_ Member ID \_\_\_\_\_  
 If patient is not insured, check this box  
Secondary Insurance \_\_\_\_\_ Member ID \_\_\_\_\_

## SECTION 2 — CLINICAL REQUIREMENTS

\*Select at least one modality or check all that apply\*

**TENS, 4-Lead (E0730)**

Acute Postprocedural Pain (G89.18)

Chronic Postprocedural Pain (G89.28)

Chronic Pain Syndrome (G89.4)

Other: \_\_\_\_\_

**IFC (E1399)**

Rationale for IFC (required): \_\_\_\_\_

If Chronic Pain:

TENS Trial Conducted (Date Range: \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_ - \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_)

and  DID provide significant pain relief and improvement in function

DID NOT provide significant pain relief and improvement in function

No Trial Conducted

If Acute Post-Surgical, Date of Surgery: \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_

**NMES (E0745)**

Body Area	Left	Right
Shoulder	<input type="checkbox"/> M62.512	<input type="checkbox"/> M62.511
Upper Arm	<input type="checkbox"/> M62.522	<input type="checkbox"/> M62.521
Forearm	<input type="checkbox"/> M62.532	<input type="checkbox"/> M62.531
Hand	<input type="checkbox"/> M62.542	<input type="checkbox"/> M62.541
Thigh	<input type="checkbox"/> M62.552	<input type="checkbox"/> M62.551
Lower Leg	<input type="checkbox"/> M62.562	<input type="checkbox"/> M62.561
Ankle/Foot	<input type="checkbox"/> M65.572	<input type="checkbox"/> M65.571

Other Single Site (M62.58): \_\_\_\_\_ (Specify)

Other Diagnosis: \_\_\_\_\_ (Specify)

**Cause of Atrophy:**

Casting or Splinting of a Limb

Contracture Due to Scarring of Soft Tissue

ACL Repair/Knee Replacement Leading to Quad Atrophy

Other: \_\_\_\_\_

**Nerve Supply Intact?**  Yes  No

## SECTION 3 — THERAPY SCHEDULE

Length of Need: \_\_\_\_\_ month(s) (1-24)

**Prescribed Use**

Minutes Per Session: \_\_\_\_\_

Sessions Per Day: \_\_\_\_\_

Days Per Week: \_\_\_\_\_

Replace Electrodes:  Weekly  Bi-Weekly  Monthly

## SECTION 4 — SUPPLIES

**A. Initial Order includes:**

- Lead Wires (1 set)
- AC Adapter (1 unit)
- Batteries (4 each)
- Electrodes (16 - 2" round (4 x 4/pk))

**B. Optional Garment(s) (E0731):**

**Conductive**

- Elbow  Knee  Sock
- Glove, Small (7" x 3")
- Glove, Large (8.5" x 3.5")

**Non-Conductive**

- Care Vest: Chest circumference: \_\_\_\_\_ in.
- Back: Waist circumference: \_\_\_\_\_ in.
- Lumbar: Lumbar circumference: \_\_\_\_\_ in.
- Shoulder: Chest circumference: \_\_\_\_\_ in.
- Neck: Neck circumference: \_\_\_\_\_ in.

**Reason the Garment(s) Are Necessary:**

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## SECTION 5 — PRESCRIBER INFORMATION

Name: \_\_\_\_\_ License: \_\_\_\_\_ NPI: \_\_\_\_\_

Address: \_\_\_\_\_ City: \_\_\_\_\_

State: \_\_\_\_\_ ZIP: \_\_\_\_\_ Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

**Prescriber Attestation:** By signing below, I confirm that the patient is being treated by me. All the information contained on this form accurately reflects the patient's medical condition and the treatment regimen I have prescribed. The medical records for this patient substantiate the prescribed treatment frequency. The patient/caregiver is able to follow instructions for using an e-stim device and is able to use the ordered items. I certify my patient has no medical contraindications making this therapy inappropriate, e.g., pacemaker. For Medicare/Insurance requirements, I will maintain this signed original document in the patient's medical record file for post-payment review/audit purposes.

Prescriber Signature: \_\_\_\_\_

Date: \_\_\_\_\_

**FAX TO: +1 (866) 791-2026 OR EMAIL TO ORDERS@ZYNEX.COM**

**Required: Please attach medical records and copies of the front and back of all insurance cards with the order.**