

JONES MEDICAL EQUIPMENT

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ALL PHYSICIANS SCRIPTING FOR MEDICARE AND MEDICAID MUST BE PECOS CERTIFIED AND ENROLLED WITH GAMMIS. INFORMATION WILL BE VERIFIED DURING THE INTAKE PROCESS AND DENIED, IF YOU'RE NOT ENROLLED

Transcutaneous Electrical Nerve Stimulation Device and Supplies **TENS UNIT**

Patient Name _____ DOB _____

Insurance _____ MBI # _____

Order Date _____ End Date _____

Treating Provider _____ NPI _____

Phone # _____ Fax # _____

Equipment Ordered (Circle/Check All that Apply)

E0720- Tens, Two Lead, Localized Stimulation	QTY _____	
E0730- Tens, Four Lead, Larger area, Multiple Nerve Stimulator	QTY _____	
A4557- Lead Wires, Replacement	QTY _____	Weekly, Monthly, Yearly
A4595- Electrical stimulator supplies, 2 lead	QTY _____	Weekly, Monthly, Yearly
E0731- Garment for Tens Device	QTY _____	Back, Knee, Sock, Glove

Other or Specific Instructions by MD _____

To be completed by provider (must be signed by the ordering M.D.)

Prognosis (circle) Fair Good Poor Length of Need _____ (99= lifetime)

Date of Order (RX) _____ Date of Office Visit _____

DX _____ ICD-10 Code _____

Indication

Does the patient have acute post-operative pain? YES NO

Surgical Site _____ Surgery Date _____

Is the presumed etiology of the pain a type that is accepted as responding to TENS therapy? YES NO

Has the pain been present for at least 3 months? YES NO

Have other appropriate treatment modalities been tried and failed? YES NO

If yes, describe? _____

Provider Signature

Date

Individually Owned and Operated