

DOCUMENTATION CHECKLIST FOR TRANSCUTANEOUS ELECTRICAL NERVE STIMULATORS (TENS)

Policy References:

- [Local Coverage Determination \(L33802\)](#)
- [Policy Article \(A52520\)](#)

Documentation References: [Standard Documentation Requirements Policy Article \(A55426\)](#)

The supplier must be able to provide all of these items on request:

[Standard Written Order \(SWO\)](#)

[Beneficiary Authorization](#)

[Proof of Delivery \(POD\)](#)

[Refill Requirements](#)

[Continued Need](#)

[Continued Use](#)

Medical records from treating practitioner as noted below

Medical records should contain:

TENS Unit (E0720, E0730)

Physician ordering the TENS unit and related supplies must be the treating physician for the disease or condition justifying the need for the TENS unit.

TENS is covered for the treatment of beneficiaries with chronic, intractable pain, or acute post-operative pain when one of the following coverage criteria are met:

Acute Post-Operative Pain

Limited to 30 days from the day of surgery

Payment made only as a rental

Documentation must include:

Date of the surgery

Nature of the surgery

Location and severity of the pain; **or**

Chronic Pain Other than Low Back Pain

Presumed etiology of the pain must be a type that is accepted as responding to TENS therapy; **and**

Pain must have been present for at least three months; **and**

Other appropriate treatment modalities must have been tried and failed.

Information in the medical record must describe:

Location of the pain

Severity of the pain

Duration of time the beneficiary has had the pain (pain must be present for at least three months)

Presumed etiology of the pain

Prior treatment and results of that treatment

Reevaluation of the beneficiary at the end of the trial period indicating how often the beneficiary used the TENS unit, typical duration of use each time, results (effectiveness of therapy)

Chronic Low Back Pain (CLBP)

TENS therapy for Chronic Low Back Pain (CLBP) will be denied as not reasonable and necessary.

General Requirements for Chronic Pain

Must be used on a trial basis for a minimum of 30 days, but not to exceed two months:

Trial period will be paid as a rental; **and**

Trial period must be monitored by the physician to determine the effectiveness of the TENS unit in modulating the pain.

For coverage as a purchase, the physician must determine that the beneficiary is likely to derive significant therapeutic benefit from continuous use over a long period of time.

If ordered for use with four leads, the medical record must document why two leads are insufficient to meet the beneficiary's needs.

General Requirements for CLBP

If ordered for use with four leads, the medical record must document why two leads are insufficient to meet the beneficiary's needs.

Conductive Garment (E0731)

Had been prescribed by the treating physician for use in delivering covered TENS treatment;
and

Beneficiary meets one of the covered medical indications:

Beneficiary cannot manage without the conduct garment because:

There is such a large area or so many sites to be stimulated; **and**

Stimulation would have to be delivered so frequently that it is not feasible to use conventional electrodes, adhesive tape, and lead wires; **or**

Beneficiary cannot manage without the conduct garment for the treatment of chronic intractable pain because the areas or sites to be stimulated are inaccessible with the use of conventional electrodes, adhesive tapes and lead wires; **or**

Beneficiary has a documented medical condition, such as skin problems, that precludes the application of conventional electrodes, adhesive tapes and lead wires; **or**

Beneficiary requires electrical stimulation beneath a cast to treat chronic intractable pain
Conductive garment is only covered during the trial rental period if:

Beneficiary has documented skin problem prior to the start of the trial period; **and**

TENS is reasonable and necessary for the beneficiary.