

<b>Case Number:</b>	CM15-0061720		
<b>Date Assigned:</b>	04/07/2015	<b>Date of Injury:</b>	10/14/2013
<b>Decision Date:</b>	05/06/2015	<b>UR Denial Date:</b>	03/02/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	04/01/2015

### HOW THE IMR FINAL DETERMINATION WAS MADE

MAXIMUS Federal Services sent the complete case file to an expert reviewer. He/she has no affiliation with the employer, employee, providers or the claims administrator. He/she has been in active clinical practice for more than five years and is currently working at least 24 hours a week in active practice. The expert reviewer was selected based on his/her clinical experience, education, background, and expertise in the same or similar specialties that evaluate and/or treat the medical condition and disputed items/Service. He/she is familiar with governing laws and regulations, including the strength of evidence hierarchy that applies to Independent Medical Review determinations.

The Expert Reviewer has the following credentials:  
 State(s) of Licensure: North Carolina  
 Certification(s)/Specialty: Family Practice

### CLINICAL CASE SUMMARY

The expert reviewer developed the following clinical case summary based on a review of the case file, including all medical records:

The injured worker is a 57 year old male, who sustained an industrial injury on 10/14/2013. He reported a drill hitting his right hand and his head hitting a pipe. Diagnoses have included right wrist tenosynovitis, sacroiliac sprain/strain, right carpal tunnel syndrome, myofascial pain syndrome of the right upper limb and low back pain. Treatment to date has included lumbar magnetic resonance imaging (MRI), splinting, physical therapy, right wrist injection and medication. According to the progress report dated 2/19/2015, the injured worker complained of moderate low back pain with deep aching and burning. He complained of moderate, intermittent right wrist pain. He also complained of occasional neck aching pains and pain that shoots to his right shoulder and arm. Range of motion of the lumbar spine was very stiff and guarded. There was tenderness to palpation of the paravertebral muscles. Straight leg raising was positive on the left side. Exam of the right wrist revealed swelling. Authorization was requested for a transcutaneous electrical nerve stimulation (TENS) unit and transcutaneous electrical nerve stimulation (TENS) electrodes purchase.

### IMR ISSUES, DECISIONS AND RATIONALES

The Final Determination was based on decisions for the disputed items/services set forth below:

**TENS (transcutaneous electrical nerve stimulation) Electrodes, Qty 2 - E0720, A4595:**  
 Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Transcutaneous electrotherapy Page(s): 114-116.

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines TENS Page(s): 114.

**Decision rationale:** The California chronic pain medical treatment guidelines section on transcutaneous electrical nerve stimulation states: TENS, chronic pain (transcutaneous electrical nerve stimulation) Not recommended as a primary treatment modality, but a one-month home-based TENS trial may be considered as a noninvasive conservative option, if used as an adjunct to a program of evidence-based functional restoration, for the conditions described below. While TENS may reflect the long-standing accepted standard of care within many medical communities, the results of studies are inconclusive; the published trials do not provide information on the stimulation parameters which are most likely to provide optimum pain relief, nor do they answer questions about long-term effectiveness. (Carroll-Cochrane, 2001) Several published evidence-based assessments of transcutaneous electrical nerve stimulation (TENS) have found that evidence is lacking concerning effectiveness. One problem with current studies is that many only evaluated single-dose treatment, which may not reflect the use of this modality in a clinical setting. Other problems include statistical methodology, small sample size, influence of placebo effect, and difficulty comparing the different outcomes that were measured. This treatment option is recommended as an adjunct to a program of evidence based functional restoration. However, it is recommended for a one-month trial to document subjective and objective gains from the treatment. There is no provided documentation of a one-month trial period with objective measurements of improvement. Therefore, criteria have not been met and the request is not medically necessary.

**TENS (transcutaneous electrical nerve stimulation) Unit - E0720, A4595, Purchase:** Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Transcutaneous electrotherapy Page(s): 114-116.

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines TENS Page(s): 114.

**Decision rationale:** The California chronic pain medical treatment guidelines section on transcutaneous electrical nerve stimulation states: TENS, chronic pain (transcutaneous electrical nerve stimulation) Not recommended as a primary treatment modality, but a one-month home-based TENS trial may be considered as a noninvasive conservative option, if used as an adjunct to a program of evidence-based functional restoration, for the conditions described below. While TENS may reflect the long-standing accepted standard of care within many medical communities, the results of studies are inconclusive; the published trials do not provide information on the stimulation parameters, which are most likely to provide optimum pain relief, nor do they answer questions about long-term effectiveness. (Carroll-Cochrane, 2001) Several published evidence-based assessments of transcutaneous electrical nerve stimulation (TENS) have found that evidence is lacking concerning effectiveness. One problem with current studies

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