

<b>Case Number:</b>	CM15-0047254		
<b>Date Assigned:</b>	03/19/2015	<b>Date of Injury:</b>	01/30/2013
<b>Decision Date:</b>	04/24/2015	<b>UR Denial Date:</b>	02/20/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	03/12/2015

### 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: New Jersey

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:

This 62 year old female sustained an industrial injury to the neck and right shoulder on 1/30/14. Previous treatment included magnetic resonance imaging, physical therapy and medications. In a PR-2 dated 1/31/15, the injured worker complained of pain to the right shoulder and cervical spine, rated 7-9/10 on the visual analog scale with numbness and tingling. The injured worker also complained of anxiety. Current diagnoses included cervical disc syndrome and right shoulder internal derangement. The treatment plan included medications (Tramadol and Lidoderm patches), right shoulder arthroscopy and a home interferential stimulator unit for a 60 day initial trial rental.

### IMR ISSUES, DECISIONS AND RATIONALES

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

**Home Interferential Stimulator 60 days trial:** Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Interferential Current Stimulation (ICS) Page(s): 120.

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Interferential Current Stimulation (ICS) Page(s): 118-120.

**Decision rationale:** The MTUS Chronic Pain Guidelines do not recommend interferential current stimulation (ICS) as an isolated intervention as there is no quality evidence. It may be considered as an adjunct if used in conjunction with recommended treatments, including return to work, exercise, and medications if these have not shown to provide significant improvements in function and pain relief, and has already been applied by the physician or physical therapist with evidence of effectiveness in the patient. Criteria for consideration would include if the patient's pain is ineffectively controlled due to diminished effectiveness of medications, pain is ineffectively controlled with medications due to side effects, if the patient has a history of substance abuse, if the patient has significant pain from postoperative conditions which limits the ability to perform exercise programs or physical therapy treatments, or if the patient was unresponsive to conservative measures (repositioning, heat/ice, etc.). A one month trial may be appropriate if one of these criteria are met as long as there is documented evidence of functional improvement and less pain and evidence of medication reduction during the trial period. Continuation of the ICS may only be continued if this documentation of effectiveness is provided. Also, a jacket for ICS should only be considered for those patients who cannot apply the pads alone or with the help of another available person, and this be documented. In the case of this worker, although the records provided for review suggest she is a candidate for an interferential unit trial, this request for an interferential stimulator was for a 60-day trial, which is longer than necessary to learn if it is helpful or not. Therefore, due to the fact that the request was more than a 30-day trial, it will be regarded as not medically necessary.