

<b>Case Number:</b>	CM15-0073995		
<b>Date Assigned:</b>	04/24/2015	<b>Date of Injury:</b>	01/25/2014
<b>Decision Date:</b>	05/26/2015	<b>UR Denial Date:</b>	04/02/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	04/17/2015

### 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: Indiana

Certification(s)/Specialty: Preventive Medicine, Occupational Medicine

### 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 41 year old female, who sustained an industrial injury on 01/25/2014. According to the only progress report submitted for review and dated March 13, 2015, the injured worker complained of low back pain. She reported that Diclofenac gave her headaches and constipation. Medications included Diclofenac and Norco. Diagnoses included displaced lumbar intervertebral disc without myelopathy and unspecified thoracic/lumbar neuritis. Treatment plan included L4-S1 ESFI x 2. A urine toxicology screen was performed to monitor medication compliance. Currently under review is the request for L4-S1 Lumbar ESFI quantity 2, post-injection physical therapy 3 times a week lumbar spine quantity 9 and urinalysis/ pregnancy test.

### IMR ISSUES, DECISIONS AND RATIONALES

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

**L4-S1 Lumbar ESFI Qty 2:** Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines.

**MAXIMUS guideline:** Decision based on MTUS ACOEM Chapter 12 Low Back Complaints Page(s): 287-315, Chronic Pain Treatment Guidelines epidural steroid injections Page(s): 46. Decision based on Non-MTUS Citation Official Disability Guidelines (ODG) Low Back - Lumbar & Thoracic (Acute & Chronic), Epidural steroid injections (ESIs), therapeutic.

**Decision rationale:** MTUS Chronic pain medical treatment guidelines state that epidural steroid injections are "Recommended as an option for treatment of radicular pain (defined as pain in dermatomal distribution with corroborative findings of radiculopathy) . . . Epidural steroid injection can offer short term pain relief and use should be in conjunction with other rehab efforts, including continuing a home exercise program." There were no medical documents provided to conclude that other rehab efforts or home exercise program is ongoing. Additionally, no objective findings were documented to specify the dermatomal distribution of pain. MTUS further defines the criteria for epidural steroid injections to include: 1) Radiculopathy must be documented by physical examination and corroborated by imaging studies and/or electrodiagnostic testing. 2) Initially unresponsive to conservative treatment (exercises, physical methods, NSAIDs and muscle relaxants). 3) Injections should be performed using fluoroscopy (live x-ray) for guidance. 4) If used for diagnostic purposes, a maximum of two injections should be performed. A second block is not recommended if there is inadequate response to the first block. Diagnostic blocks should be at an interval of at least one to two weeks between injections. 5) No more than two nerve root levels should be injected using transforaminal blocks. 6) No more than one interlaminar level should be injected at one session. 7) In the therapeutic phase, repeat blocks should be based on continued objective documented pain and functional improvement, including at least 50% pain relief with associated reduction of medication use for six to eight weeks, with a general recommendation of no more than 4 blocks per region per year. (Manchikanti, 2003) (CMS, 2004) (Boswell, 2007) 8) Current research does not support a "series-of-three" injections in either the diagnostic or therapeutic phase. We recommend no more than 2 ESI injections. The patient is taking multiple medications, but the progress reports do not document how long the patient has been on these medications and the "unresponsiveness" to the medications. Additionally, treatment notes do not indicate if other conservative treatments were tried and failed (exercises, physical therapy, etc). As such, the request is not medically necessary.

**Post-Injection Physical Therapy 3 Times A Week for 3 Weeks Lumbar Spine Qty 9:**  
Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines.

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Physical Medicine Page(s): 98-99. Decision based on Non-MTUS Citation Official Disability Guidelines (ODG) Low Back - Lumbar & Thoracic (Acute & Chronic), Physical Therapy.

**Decision rationale:** California MTUS guidelines refer to physical medicine guidelines for physical therapy and recommends as follows: "Allow for fading of treatment frequency (from up to 3 visits per week to 1 or less), plus active self-directed home Physical Medicine." Additionally, ACOEM guidelines advise against passive modalities by a therapist unless

exercises are to be carried out at home by patient. ODG quantifies its recommendations with 10 visits over 8 weeks for lumbar sprains/strains and 9 visits over 8 weeks for unspecified backache/lumbago. ODG further states that a "six-visit clinical trial" of physical therapy with documented objective and subjective improvements should occur initially before additional sessions are to be warranted. The request is for post-op ESI, which was denied and then found by IMR to be not necessary. Therefore, the related physical therapy is not medically necessary.

**Urinalysis/Pregnancy Test:** Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines.

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines Opioids; Substance Abuse Page(s): 74-109. Decision based on Non-MTUS Citation University of Michigan Health System Guidelines for Clinical Care: Managing Chronic Non-terminal Pain, Including Prescribing Controlled Substances (May 2009), pg 32 Established Patients Using a Controlled Substance.

**Decision rationale:** MTUS states that use of urine drug screening for illegal drugs should be considered before therapeutic trial of opioids are initiated. Additionally, use of drug screening or inpatient treatment with issues of abuse, addiction, or poor pain control. Documentation of misuse of medications (doctor-shopping, uncontrolled drug escalation, drug diversion) would indicate need for urine drug screening. There is insufficient documentation provided to suggest issues of abuse, addiction, or poor pain control by the treating physician. University of Michigan Health System Guidelines for Clinical Care: Managing Chronic Non-terminal Pain, Including Prescribing Controlled Substances (May 2009) recommends for stable patients without red flags "twice yearly urine drug screening for all chronic non-malignant pain patients receiving opioids once during January-June and another July-December." The request is for pre-ESI testing, which was denied and then found by IMR to be not necessary. Therefore, the related UA/pregnancy test is not medically necessary.