

Case Number:	CM15-0091505		
Date Assigned:	05/15/2015	Date of Injury:	11/24/2010
Decision Date:	06/17/2015	UR Denial Date:	04/27/2015
Priority:	Standard	Application Received:	05/12/2015

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: Arizona, California

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 48-year-old male, who sustained an industrial injury on 11/24/10. He reported hearing a crack noise in back along with pain. The injured worker was diagnosed as having lumbosacral disc protrusion, lumbar sacral radiculopathy, cervical spine mild spinal stenosis and status post lumbar surgery. Treatment to date has included oral medications, physical therapy, chiropractic treatment and activity restrictions. It is noted (MRI) magnetic resonance imaging of cervical spine revealed C6-7 central and foraminal stenosis. Currently, the injured worker complains of back and buttocks pain with pain to shoulder and traveling down to right arm. He is currently off work and will return to work with modified duties. Physical exam noted radicular pain along C6-7 distribution and decreased sensation to lateral forearm and deltoid patch. The treatment plan included cervical epidural steroid injections.

IMR ISSUES, DECISIONS AND RATIONALES

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

Epidural steroid injection at C5-6: Overturned

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Epidural Steroid Injections (ESIs).

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines epidural
Page(s): 47.

Decision rationale: According to the guidelines, the criteria for the use of Epidural steroid injections: Note: The purpose of ESI is to reduce pain and inflammation, restoring range of motion and thereby facilitating progress in more active treatment programs, and avoiding surgery, but this treatment alone offers no significant long-term functional benefit. 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. In this case, the claimant has decreased sensation in the deltoid region. There are abnormalities in the MRI, which can cause radicular symptoms. The request for an ESI is appropriate and medically necessary.