

Case Number:	CM15-0090999		
Date Assigned:	05/15/2015	Date of Injury:	08/08/2014
Decision Date:	07/03/2015	UR Denial Date:	04/27/2015
Priority:	Standard	Application Received:	05/11/2015

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 38 year old female injured worker suffered an industrial injury on 08/08/2014. The diagnoses included neck pain, cervical degenerative disc disease, cervical discogenic pain, low back pain, lumbar discogenic pain, chronic pain syndrome and myalgia. The diagnostics included cervical and lumbar magnetic resonance imaging. The injured worker had been treated with chiropractic therapy, medications, physical therapy. On 4/17/2015 the treating provider reported neck and low back and the pain was getting worse. She reported occasional numbness and weakness to the right foot with pain rated 9/10 without medications and 3 to 4/10 with medications. On exam there was tenderness over the cervical muscle with pain on range of motion. The lumbar spine had moderate tenderness and positive straight leg raise. The treatment plan included Prilosec, electromyographic studies/nerve conduction velocity studies for bilateral upper extremities.

IMR ISSUES, DECISIONS AND RATIONALES

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

Prilosec 20mg, QTY: 30: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines NSAIDs, GI symptoms & cardiovascular risk Page(s): 68.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines NSAIDs; GI risk Page(s): 68-69. Decision based on Non-MTUS Citation Official Disability Guidelines (ODG) Pain (Chronic), NSAIDs, GI symptoms & cardiovascular risk.

Decision rationale: MTUS states "Determine if the patient is at risk for gastrointestinal events: (1) age > 65 years; (2) history of peptic ulcer, GI bleeding or perforation; (3) concurrent use of ASA, corticosteroids, and/or an anticoagulant; or (4) high dose/multiple NSAID (e.g., NSAID + low-dose ASA)." And "Patients at intermediate risk for gastrointestinal events and no cardiovascular disease: (1) A non-selective NSAID with either a PPI (Proton Pump Inhibitor, for example, 20 mg omeprazole daily) or misoprostol (200 g four times daily) or (2) a Cox-2 selective agent. Long-term PPI use (> 1 year) has been shown to increase the risk of hip fracture (adjusted odds ratio 1.44)." The medical documents provided do not establish the patient has having documented GI bleeding/perforation/peptic ulcer or other GI risk factors as outlined in MTUS. Additionally, there is no evidence provided to indicate the patient suffers from dyspepsia because of the present medication regimen. As such, the request for Omeprazole 20mg #30 is not medically necessary.

EMG of the right upper extremity: Upheld

Claims Administrator guideline: Decision based on MTUS ACOEM Chapter 8 Neck and Upper Back Complaints Page(s): 178.

MAXIMUS guideline: Decision based on MTUS ACOEM Chapter 11 Forearm, Wrist, and Hand Complaints Page(s): 260-262. Decision based on Non-MTUS Citation Official Disability Guidelines (ODG) Pain, Electrodiagnostic testing (EMG/NCS).

Decision rationale: ACOEM States Appropriate electro diagnostic studies (EDS) may help differentiate between CTS and other conditions, such as cervical radiculopathy. These may include nerve conduction studies (NCS), or in more difficult cases, electromyography (EMG) may be helpful. ODG states Recommended needle EMG or NCS, depending on indications. Surface EMG is not recommended. Electromyography (EMG) and Nerve Conduction Studies (NCS) are generally accepted, well-established and widely used for localizing the source of the neurological symptoms and establishing the diagnosis of focal nerve entrapments, such as carpal tunnel syndrome or radiculopathy, which may contribute to or coexist with CRPS II (causalgia), when testing is performed by appropriately trained neurologists or physical medicine and rehabilitation physicians (improperly performed testing by other providers often gives inconclusive results). As CRPS II occurs after partial injury to a nerve, the diagnosis of the initial nerve injury can be made by electrodiagnostic studies. ODG further clarifies NCS is not recommended, but EMG is recommended as an option (needle, not surface) to obtain unequivocal evidence of radiculopathy, after 1-month conservative therapy, but EMG's are not necessary if radiculopathy is already clinically obvious. In this case, the physical exam clearly demonstrates radiculopathy, so the request is not medically necessary.

NCS of the right upper extremity: Upheld

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EMG of the left upper extremity: Upheld

Claims Administrator guideline: Decision based on MTUS ACOEM Chapter 8 Neck and Upper Back Complaints Page(s): 178.

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