

Case Number:	CM15-0117512		
Date Assigned:	06/25/2015	Date of Injury:	10/31/2005
Decision Date:	07/24/2015	UR Denial Date:	05/19/2015
Priority:	Standard	Application Received:	06/18/2015

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: California
 Certification(s)/Specialty: Physical Medicine & Rehabilitation

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 61 year old female who sustained an industrial injury on 10/31/05. Initial complaints and diagnoses are not available. Treatments to date include medications, bilateral cubital tunnel releases, carpal tunnel releases, and ulnar nerve decompressions at the wrists. Diagnostic studies are not addressed. Current complaints include pain in the neck which radiates to the arm, and pain in the hands. Current diagnoses include cervical radiculopathy, bilateral carpometacarpal arthrosis, trapezial/paracervical/parascapular strain, bilateral forearm tendinitis, left thoracic outlet syndrome, and left lateral epicondylitis. In a progress note dated 05/07/15 the treating provider reports the plan of care as physical therapy, continued nonsteroidal anti-inflammatory and stomach protective medications, as well as cervical epidural steroid injection series and follow-up evaluation and treatment of the cervical spine. The requested treatments include physical therapy to the forearm and Prilosec.

IMR ISSUES, DECISIONS AND RATIONALES

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

12 physical therapy sessions for the forearm: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Physical Therapy (PT).

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Physical Therapy, pages 98-99.

Decision rationale: Physical therapy is considered medically necessary when the services require the judgment, knowledge, and skills of a qualified physical therapist due to the complexity and sophistication of the therapy and the physical condition of the patient. However, there is no clear measurable evidence of progress with the PT treatment already rendered including milestones of increased ROM, strength, and functional capacity. Review of submitted physician reports show no evidence of functional benefit, unchanged chronic symptom complaints, clinical findings, and functional status. There is no evidence documenting functional baseline with clear goals to be reached and the patient striving to reach those goals. The Chronic Pain Guidelines allow for visits of physical therapy with fading of treatment to an independent self-directed home program. It appears the employee has received significant therapy sessions without demonstrated evidence of functional improvement to allow for additional therapy treatments. There is no report of acute flare-up, new injuries, or change in symptom or clinical findings to support for formal PT in a patient that has been instructed on a home exercise program for this chronic injury. Submitted reports have not adequately demonstrated the indication to support further physical therapy when prior treatment rendered has not resulted in any functional benefit. The 12 physical therapy sessions for the forearm is not medically necessary and appropriate.

Prilosec 20mg #60: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines NSAIDs, GI Symptoms and Cardiovascular risk, Pages 68-69. Decision based on Non-MTUS Citation ODG, Pain Chapter, Proton Pump Inhibitors (Updated 6/15/15).

Decision rationale: Proton pump inhibitor (PPI) medication is for treatment of the problems associated with active gastric ulcers, erosive esophagitis, Barrett's esophagitis, or in patients with pathologic hypersecretion diseases. Although preventive treatment is effective for the mentioned diagnosis, studies suggest; however, nearly half of PPI prescriptions are used for unapproved or no indications. Per MTUS Chronic Pain Treatment Guidelines, the patient does not meet criteria for Omeprazole (Prilosec) namely reserved for patients with history of prior GI bleeding, the elderly (over 65 years), diabetics, and chronic cigarette smokers. Long term use of PPIs have potential increased risks of B12 deficiency; iron deficiency; hypomagnesemia; susceptibility to pneumonia, enteric infections, fractures, hypergastrinemia and cancer, and cardiovascular effects of myocardial infarction (MI). In the elderly, studies have demonstrated increased risk for Clostridium difficile infection, bone loss, and fractures from long-term use of PPIs. Given treatment criteria outweighing risk factors, if a PPI is to be used, omeprazole (Prilosec), lansoprazole (Prevacid), and esomeprazole (Nexium) are to be considered over second-line therapy of other PPIs such as pantoprazole (Protonix), dexlansoprazole (Dexilant), and rabeprazole (Aciphex). Submitted reports have not described or provided any GI diagnosis that meets the criteria to indicate medical treatment. Review of the records show no documentation of any history, symptoms, or GI diagnosis to warrant this medication. The Prilosec 20mg #60 is not medically necessary and appropriate.

