

Case Number:	CM15-0051322		
Date Assigned:	03/24/2015	Date of Injury:	11/07/2012
Decision Date:	05/01/2015	UR Denial Date:	02/17/2015
Priority:	Standard	Application Received:	03/18/2015

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: North Carolina
 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 58 year old male who sustained an industrial injury on 11/7/12. The injured worker reported symptoms in the bilateral upper extremities. The injured worker was diagnosed as having right wrist sprain with mild capsulitis, status post right carpal tunnel release, right stenosing tenosynovitis, left stenosis tenosynovitis, right ulnar neuropathy, left ulnar neuropathy and left median neuropathy. Treatments to date have included splinting, anti-inflammatory medications, cold/heat packs, physical therapy, activity modifications, and a cortisone injection. Currently, the injured worker complains of pain in the bilateral wrists and right elbow. The plan of care was for a wrist brace, medication prescriptions and a follow up appointment at a later date.

IMR ISSUES, DECISIONS AND RATIONALES

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

45 Tylenol #4 ([REDACTED]): Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines opioids
Page(s): 76-84.

Decision rationale: The ACOEM chapter on low back complaints and treatment recommendations states: Lumbar supports have not been shown to have any lasting benefit beyond the acute phase of symptom relief. This patient has chronic ongoing low back complaints and is status post-lumbar laminectomy. Per the ACOEM, lumbar supports have no lasting benefit outside of the acute phase of injury. This patient is well past the acute phase of injury and there is no documentation of acute flare up of chronic low back pain. Therefore criteria for use of lumbar support per the ACOEM have not been met and the request is not medically necessary.

30 Prilosec 20mg ([REDACTED]): Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines NSAIDS
Page(s): 68.

Decision rationale: The California chronic pain medical treatment guidelines section on NSAID therapy and proton pump inhibitors (PPI) states: Recommend with precautions as indicated below. Clinicians should weight the indications for NSAIDs against both GI and cardiovascular risk factors. 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). Recent studies tend to show that H. Pylori does not act synergistically with NSAIDS to develop gastro duodenal lesions. Recommendations Patients with no risk factor and no cardiovascular disease: Non-selective NSAIDs OK (e.g., ibuprofen, naproxen, etc.) 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 mg 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). Patients at high risk for gastrointestinal events with no cardiovascular disease: A Cox-2 selective agent plus a PPI if absolutely necessary. There is no documentation provided that places this patient at intermediate or high risk that would justify the use of a PPI. There is no mention of current gastrointestinal or cardiovascular disease. For these reasons the criteria set forth above per the California MTUS for the use of this medication has not been met. Therefore the request is not medically necessary.

30 Robaxin 750mg ([REDACTED]): Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines muscle relaxants
Page(s): 63-65.

Decision rationale: California chronic pain medical treatment guidelines section on muscle relaxants states: Recommend non-sedating muscle relaxants with caution as a second-line option for short-term treatment of acute exacerbations in patients with chronic LBP. (Chou, 2007) (Mens, 2005) (Van Tulder, 1998) (van Tulder, 2003) (van Tulder, 2006) (Schnitzer, 2004) (See, 2008) Muscle relaxants may be effective in reducing pain and muscle tension, and increasing mobility. However, in most LBP cases, they show no benefit beyond NSAIDs in pain and overall improvement. Also there is no additional benefit shown in combination with NSAIDs. Efficacy appears to diminish over time, and prolonged use of some medications in this class may lead to dependence. (Homik, 2004) (Chou, 2004) This medication is not intended for long-term use per the California MTUS. The medication has not been prescribed for the flare-up of chronic low back pain. This is not an approved use for the medication. For these reasons, criteria for the use of this medication have not been met. Therefore the request is not medically necessary.