

Case Number:	CM15-0071050		
Date Assigned:	04/21/2015	Date of Injury:	02/18/2014
Decision Date:	07/02/2015	UR Denial Date:	03/30/2015
Priority:	Standard	Application Received:	04/14/2015

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, Georgia

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 42 year old male who sustained an industrial injury on 2/18/14. The injured worker reported symptoms in the back. The injured worker was diagnosed as having low back pain, lumbar degenerative disc disease, lumbar spondylosis, chronic pain syndrome and myalgia. Treatments to date have included chiropractic treatments, muscle relaxants, and non-steroidal anti-inflammatory drugs. Currently, the injured worker complains of back pain. The plan of care was for epidural steroid injection, 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:

Lumbar transforaminal epidural steroid injection at right S1: Overturned

Claims Administrator guideline: The Claims Administrator did not base their decision on the MTUS. Decision based on Non-MTUS Citation Official Disability Guidelines.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Section 2 Page(s): 46.

Decision rationale: CA MTUS guidelines state that epidural steroid injections are an option for the treatment of radicular pain with guidelines recommending no more than 2 epidural steroid injections to for diagnostic purposes. Criteria for ESI includes radiculopathy documented by physical examination and corroborated by imaging and documentation of trial of conservative therapies including NSAIDs, physical therapy, exercise. Repeat epidural blocks should be used only when a 50 % reduction in pain accompanied by reduced medication usage for 6-8 weeks. In this case, there is documented failure of conservative therapy and documentation by physical examination of radiculopathy, which is consistent with imaging findings. Epidural steroid injection is medically necessary.

Ultram 150mg #60: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Section 2 Page(s): 74-89.

Decision rationale: CA MTUS allows for the use of opioid medication, such as Ultram, for the management of chronic pain and outlines clearly the documentation that would support the need for ongoing use of an opioid. These steps include documenting pain and functional improvement using validated measures at 6 months intervals, documenting the presence or absence of any adverse effects, documenting the efficacy of any other treatments and of any other medications used in pain treatment. The medical record in this case does not use any validated method of recording the response of pain to the opioid medication or of documenting any functional improvement. Therefore, the record does not support medical necessity of ongoing opioid therapy with Ultram. The request is not medically necessary.

Flexeril 7.5mg #60: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Section 2 Page(s): 63-66.

Decision rationale: The CA MTUS allows for the use, with caution, of non-sedating muscle relaxers as second line treatment for acute exacerbations of chronic low back pain. While they may be effective in reducing pain and muscle tension, most studies show no benefits beyond NSAIDs in pain relief. Efficacy diminishes over time and prolonged use may lead to dependency. There is no recommendation for ongoing use in chronic pain. The medical record in this case does not document an acute exacerbation and the request is for ongoing regular daily use of Flexeril. This is not medically necessary and the original UR decision is upheld.

Anaprox 550mg #60: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Section 2 Page(s): 67-68.

Decision rationale: CA MTUS guidelines are clear that NSAIDs should be used at the lowest possible dose for the shortest period possible. There is specific caution that NSAIDS have been shown to slow healing in all soft tissue including muscle, ligaments, tendons and cartilage. The request for Anaprox 550 mg #60 does not meet the criteria of providing lowest dose of NSAID for the shortest time possible, as this dose is the maximum dose allowable. There is no documentation of response to this dose or of any trials of lower doses of Anaprox. Anaprox 550 mg #60 is not medically necessary.

Prilosec 20mg #60: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Section 2 Page(s): 68.

Decision rationale: CA MTUS guidelines state that a proton pump inhibitor should be considered for administration with anti-inflammatory medication if there is a high risk for gastro- intestinal events. In this case, the medical record does not document any history to indicate a moderate or high risk for gastrointestinal events and Prilosec therefore is not medically necessary.