

Case Number:	CM15-0219792		
Date Assigned:	11/12/2015	Date of Injury:	06/21/2007
Decision Date:	12/22/2015	UR Denial Date:	10/30/2015
Priority:	Standard	Application Received:	11/09/2015

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, Oregon, Washington
 Certification(s)/Specialty: Orthopedic Surgery

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 46 year old female who sustained an industrial injury 06-21-07. A review of the medical records reveals the injured worker is undergoing treatment for lumbar intervertebral disc displacement. Medical records (10-12-15) reveal the injured worker complains of low back pain, which is not rated. She reports 50% improvement in pain with acupuncture. The physical exam (10-12-15) reveals a generic exam, noting the injured worker is morbidly obese with no atrophy in the extremities. There is no documentation of an examination of the spine. Prior treatment includes 12 sessions of acupuncture, a lumbar epidural steroid injection, and medications including hydrocodone, and Topamax, Flexeril, Nabumetone-relafen. The original utilization review (10-30-15) non certified the request for Flexeril 5mg #60. The documentation supports that he injured worker has been on Flexeril since at least 08-14-15.

IMR ISSUES, DECISIONS AND RATIONALES

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

Flexeril 5mg quantity 60: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Cyclobenzaprine (Flexeril).

MAXIMUS guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Cyclobenzaprine (Flexeril).

Decision rationale: According to the CA MTUS, Chronic Pain Medical Treatment Guidelines, Cyclobenzaprine, pages 41-42 "Recommended as an option, using a short course of therapy. Cyclobenzaprine (Flexeril) is more effective than placebo in the management of back pain; the effect is modest and comes at the price of greater adverse effects. The effect is greatest in the first 4 days of treatment, suggesting that shorter courses may be better. (Browning, 2001) Treatment should be brief. There is also a post-op use. The addition of cyclobenzaprine to other agents is not recommended." CA MTUS Chronic Pain Medical Treatment Guidelines, pages 64-65, reports that muscle relaxants are recommended to decrease muscle spasm in condition such as low back pain although it appears that these medications are often used for the treatment of musculoskeletal conditions whether spasm is present or not. The mechanism of action for most of these agents is not known. CA MTUS Chronic Pain Medical Treatment Guidelines, page 41 and 42, report that Cyclobenzaprine, is recommended as an option, using a short course of therapy. See Medications for chronic pain for other preferred options. Cyclobenzaprine (Flexeril) is more effective than placebo in the management of back pain; the effect is modest and comes at the price of greater adverse effects. The effect is greatest in the first 4 days of treatment, suggesting that shorter courses may be better. (Browning, 2001) Treatment should be brief. This medication is not recommended to be used for longer than 2-3 weeks and is typically used postoperatively. The addition of cyclobenzaprine to other agents is not recommended. In this case there is no evidence of muscle spasms on review of the medical records from 10/12/15. There is no evidence of functional improvement, a quantitative assessment on how this medication helps, percentage of relief lasts, increase in function, or increase in activity. Therefore chronic usage is not supported by the guidelines. Per CA MTUS guidelines there is no indication for the prolonged use of a muscle relaxant. Thus the prescription is not medically necessary and the recommendation is for non-certification.