

Case Number:	CM15-0111769		
Date Assigned:	06/22/2015	Date of Injury:	10/16/2008
Decision Date:	07/21/2015	UR Denial Date:	05/08/2015
Priority:	Standard	Application Received:	06/09/2015

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: 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 52 year old female patient who sustained an industrial injury on 10/16/2008. The accident was described as having had slipped and fallen over a puddle of water on the floor with resulting injury. A primary treating office visit dated 09/22/2014 reported the patient with subjective complaint of having severe right knee pain. Treatment to date included: medication, physical therapy session, injection, braces, activity modification and or rest; and underwent radiography and magnetic resonance imaging. The pain is localized to the right anterior knee graded as severe in nature. The patient is currently not working. Current medications are; Diclofenac Sodium, Norco 10/325mg, Topiramate, Carisoprodol, and Lorazepam. She is noted allergic: Naproxen and Celebrex. She is diagnosed with status post right total knee arthroplasty, and osteoarthritis left knee. There is recommendation to lose weight prior to undergoing a total knee arthroplasty. By 12/29/2014 noted the only change to case this month was that OxyContin was prescribed. There is recommendation to continue with weight loss and suggestion of undergoing a gastric banding procedure.

IMR ISSUES, DECISIONS AND RATIONALES

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

Weight loss program: Overturned

Claims Administrator guideline: The Claims Administrator did not cite any medical evidence for its decision.

MAXIMUS guideline: The Expert Reviewer did not base their decision on the MTUS. Decision based on Non-MTUS Citation NIH, weight loss.

Decision rationale: The California chronic pain medical treatment guidelines section on opioids states for ongoing management: On-Going Management. Actions Should Include: (a) Prescriptions from a single practitioner taken as directed, and all prescriptions from a single pharmacy. (b) The lowest possible dose should be prescribed to improve pain and function. (c) Office: Ongoing review and documentation of pain relief, functional status, appropriate medication use, and side effects. Pain assessment should include: current pain; the least reported pain over the period since last assessment; average pain; intensity of pain after taking the opioid; how long it takes for pain relief; and how long pain relief lasts. Satisfactory response to treatment may be indicated by the patient's decreased pain, increased level of function, or improved quality of life. Information from family members or other caregivers should be considered in determining the patient's response to treatment. The 4 A's for Ongoing Monitoring: Four domains have been proposed as most relevant for ongoing monitoring of chronic pain patients on opioids: pain relief, side effects, physical and psychosocial functioning, and the occurrence of any potentially aberrant (or nonadherent) drug-related behaviors. These domains have been summarized as the "4 A's" (analgesia, activities of daily living, adverse side effects, and aberrant drug taking behaviors). The monitoring of these outcomes over time should affect therapeutic decisions and provide a framework for documentation of the clinical use of these controlled drugs. (Passik, 2000) (d) Home: To aid in pain and functioning assessment, the patient should be requested to keep a pain diary that includes entries such as pain triggers, and incidence of end-of-dose pain. It should be emphasized that using this diary will help in tailoring the opioid dose. This should not be a requirement for pain management. (e) Use of drug screening or inpatient treatment with issues of abuse, addiction, or poor pain control. (f) Documentation of misuse of medications (doctor-shopping, uncontrolled drug escalation, drug diversion). (g) Continuing review of overall situation with regard to nonopioid means of pain control. (h) Consideration of a consultation with a multidisciplinary pain clinic if doses of opioids are required beyond what is usually required for the condition or pain does not improve on opioids in 3 months. Consider a psych consult if there is evidence of depression, anxiety or irritability. Consider an addiction medicine consult if there is evidence of substance misuse. When to Continue Opioids (a) If the patient has returned to work, (b) If the patient has improved functioning and pain. (Washington, 2002) (Colorado, 2002) (Ontario, 2000) (VA/DoD, 2003) (Maddox-AAPM/APS, 1997) (Wisconsin, 2004) (Warfield, 2004) The long-term use of this medication class is not recommended per the California MTUS unless there documented evidence of benefit with measurable outcome measures and improvement in function. The provided documentation for review meets these criteria and the request is medically necessary.

DME: Shower chair x 1: Upheld

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

MAXIMUS guideline: The Expert Reviewer did not base their decision on the MTUS. Decision based on Non-MTUS Citation ODG, durable medical equipment.

Decision rationale: The California MTUS and the ACOEM do not specifically address the requested item. Per the Official Disability Guidelines section on durable medical equipment, DME is primarily and customarily used to serve a medical purpose and generally not useful to a person in the absence of illness or injury. DME equipment is defined as equipment that can withstand repeated use i.e can be rented and used by successive patients, primarily serves a medical function and is appropriate for use in a patient's home. It does not serve a primary medical purpose that cannot be accomplished without it. Therefore criteria have not been met per the ODG and the request is not medically necessary.

DME: Scooter x 1: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines powered mobility devices Page(s): 99.

Decision rationale: The California MTUS section on powered mobility devices states: Not recommended if the functional mobility deficit can be sufficiently resolved by the prescription of a cane or walker, or the patient has sufficient upper extremity function to propel a manual wheelchair, or there is a caregiver who is available, willing, and able to provide assistance with a manual wheelchair. Early exercise, mobilization and independence should be encouraged at all steps of the injury recovery process, and if there is any mobility with canes or other assistive devices, a motorized scooter is not essential to care. The provided clinical documentation for review does not meet criteria as outlined above and the request is therefore not medically necessary.