

| | | | |
|-----------------------|--------------|------------------------------|------------|
| Case Number: | CM15-0139755 | | |
| Date Assigned: | 07/29/2015 | Date of Injury: | 09/20/2002 |
| Decision Date: | 08/26/2015 | UR Denial Date: | 07/07/2015 |
| Priority: | Standard | Application Received: | 07/20/2015 |

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 55 year old male patient who sustained an industrial injury on September 20, 2002. A primary treating office visit dated August 09, 2012 reported the patient with chief complaint of having bilateral neck pain that radiates into bilateral shoulders and bilateral upper extremities with associated headaches. Current medications are: Androgel, Testim, Dilaudid, Viagra, Avinza, Cymbalta, Klonopin, Norco, Opana, and Senokot. The assessment found the patient with cervicgia, bilateral shoulder joint pain. Previous treatment to include: modified work duty, oral and topical medications, surgical intervention, and radiofrequency lesioning in January 2010 which offered a 90 % reduction in pain for over a three month time period. The urine drug screening results discussed at visit dated December 13/2012 resulted in findings positive for Amphetamines, Benzodiazepines, THC, Methamphetamines, Opiates and Oxycodone. A recent pain management follow up visit dated June 12, 2015 reported subjective complaint of having chronic neck pain. He states that he is experiencing some increased pain due to having increased activity with having to move his residence. He states that with his usual activity the current medication regimen offers good support for the pain and consists of: Morphine Sulphate 90mg TID, Valium, Cymbalta, and Senokot. The treating diagnoses were: post laminectomy syndrome, cervical; and neck pain. The patient is requesting increased medications. He is to remain permanent and stationary.

IMR ISSUES, DECISIONS AND RATIONALES

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

Retrospective request for one (1) prescription of Morphine Sulfate ER 30mg #180 (DOS: 6/12/2015): Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Morphine Sulfate ER, CR.

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

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 non-adherent) 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 non-opioid 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. There is no documented significant decrease in objective pain measures such as VAS scores for significant periods of time. There are no objective measures of improvement of function. Therefore all criteria for the ongoing use of opioids have not been met and the request is not medically necessary