MAXIMUS Federal Services

Case Number:	CM15-0138829		
Date Assigned:	07/28/2015	Date of Injury:	06/23/2004
Decision Date:	08/25/2015	UR Denial Date:	07/08/2015
Priority:	Standard	Application Received:	07/17/2015

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 39-year-old female, who sustained an industrial injury on June 23, 2004. She reported right ankle pain. Treatment to date has included medication, surgery, physical therapy, x-ray, CT scan, MRI, nerve block injections, aqua therapy, spinal cord stimulator implant, acupuncture, toxicology screen, steroid injection (right hip) and neurology consult. Currently, the injured worker complains of constant low back pain that radiates to her hips and buttocks. She reports a numbness sensation at her top back area and lower extremity that goes to her toes. The pain is described as burning, aching and stabbing and is severe. The pain is exacerbated by standing, walking, sitting, bending and twisting. She also reports constant right lower extremity pain that radiates to her hip, knee, ankle and foot and is associated with swelling, described as aching and burning that is severe. There is a sensation of numbness in the leg, ankle and foot on the right side and also in the left knee. The pain is exacerbated by walking and activity. She reports difficulty engaging in activities of daily living and sleep disturbance. The injured worker is diagnosed with reflex sympathetic dystrophy and post left knee surgery. Her work status is permanent and stationary; however the injured worker is retired. A note dated February 23, 2015 states the injured worker initially experienced efficacy from the spinal cord stimulator, but eventually lost efficacy. A progress note dated April 22, 2015 states the injured worker received very temporary relief from the nerve block (one day). A note dated July 2, 2015 states the injured workers pain is rated at 5-7 on 10 with medication and 8-9 on 10 without medication and she is unable to bear weight. A prescription for Tramadol 50 mg #120 with 3 refills is requested to alleviate pain.

IMR ISSUES, DECISIONS AND RATIONALES

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

Tramadol 50mg, quantity: 120 with 3 refills: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Opioids Page(s): 93-94, 76-80.

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

Decision rationale: 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 dairy 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 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.