MAXIMUS Federal Services

Case Number:	CM15-0174164		
Date Assigned:	09/16/2015	Date of Injury:	10/05/1999
Decision Date:	10/16/2015	UR Denial Date:	08/12/2015
Priority:	Standard	Application Received:	09/03/2015

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 48 year old female, who sustained an industrial injury on 10-05-1999, reported as due to repetitive stress on her low back. The injured worker was diagnosed as having lumbago. Treatment to date has included diagnostics, Functional Restoration Program, epidural steroid injection, lumbar facet blocks, physical therapy, and medications. Currently (5-19-2015), the injured worker complains of "epidurals wearing off". Last epidural was documented 1-13-2015, noting 100% relief of back and leg pain for 2 month and 75% overall improvement for 5 months. Objective findings noted "emotionally worse as pain increases". "Classic radicular findings" were documented on a diagram containing the upper and lower extremities. Her work status was permanent and stationary and she was not working. Current medications were documented as "same see previous". Medication logs, undated, were noted. The use of Suboxone 2-0.5mg, 2 films daily to total 4mg daily was noted on 4-02-2014 in the document labeled Outpatient Drug Optimization and Dextoxification Services. A progress report (7-21-2014) noted the use of Suboxone 12-3mg film, 1 three to four times daily, taken starting 7-09-2014. Per the request for authorization (8/06/2015), the treatment plan included Suboxone 2-0.5mg #180, non-certified by Utilization Review on 8-12-2015. Her current Suboxone dose-use was unclear, as a progress report (7-22-2015) noted persisting and increasing back pain with radiation to the legs and feet. Pain was rated 10 out of 10. Her medications were documented as Suboxone (12-3mg film, 1 sublingual three to four times daily, taken starting 7-09-2014 and active. Other medications on 7-22-2015 were noted as Effexor XR, Advil, Aleve, and Gabapentin, noting contraindication for anti-inflammatories and gastrointestinal history.

IMR ISSUES, DECISIONS AND RATIONALES

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

Suboxone 2/.5 MG #180: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Opioids for chronic pain.

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 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 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

or how suboxone improves activities. The work status is not mentioned. Therefore not all criteria for the ongoing use of opioids have been met and the request is not medically necessary.