

<b>Case Number:</b>	CM15-0098684		
<b>Date Assigned:</b>	06/01/2015	<b>Date of Injury:</b>	05/09/2001
<b>Decision Date:</b>	06/30/2015	<b>UR Denial Date:</b>	04/23/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	05/21/2015

### 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 40 year old female, who sustained an industrial injury on 5/9/01. Initial complaints were not reviewed. The injured worker was diagnosed as having lumbago; sciatica; pain in joint of left hip. Treatment to date has included medications as reported. No diagnostics were reported or submitted in the provider's medical documentation. Currently, the PR-2 notes dated 3/9/15 indicated the injured worker presents in the office as a follow-up on medications. The reports she is about the same. She is still having pain on the left side that radiates all the way down the back of the left leg at times along with left hip pain. In addition, she gets pain that shoots all the way across her low back at times to the right-hand side. She reports no pain in the right leg. The injured worker reports she has not been using Tramadol but just wanted "codon" at the time and having fair pain relief. The current medications are listed as: Ibuprofen 600mg, Montelukast Sodium 10mg, Pantoprazole Sodium 40mg, Ranitidine HCl 150mg, Ventolin HFA 108 (90 Base) Mcg/ACT inhalation Aerosol, Carisoprodol 350mg for spasm, Tramadol 50mg, Trazodone HCl 50mg, Hydrocodone-Acetaminophen 10/325mg. The physical examination reveals obese female in mild distress. She is tender in the left lumbosacral spine and sciatic notch areas with some tenderness radiating across to the right hand-side of her low back. Musculo-skeletal area is tender about the left hip joint. The provider's treatment plan includes: water aerobics at the gym, chiropractic care for flare-ups, and continue weight loss efforts. He wants to discontinue Carisoprodol 350mg (Soma) and Tramadol HCl 50 mg (Ultram). He is requesting Hydrocodone 10/325mg #60.

## IMR ISSUES, DECISIONS AND RATIONALES

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

**Hydrocodone 10/325 mg #60:** Upheld

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

**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 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. There is no documented significant improvement in VAS scores. There are also no objective measurements of improvement in function. Therefore criteria for the ongoing use of opioids have not been met and the request is not certified.

