

<b>Case Number:</b>	CM15-0198297		
<b>Date Assigned:</b>	10/13/2015	<b>Date of Injury:</b>	07/05/2006
<b>Decision Date:</b>	12/01/2015	<b>UR Denial Date:</b>	09/28/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	10/08/2015

### 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: Texas, California

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:

This is a 62 year old female patient, who sustained an industrial injury on 7-5-2006. The diagnoses include osteochondral loose body, degenerative joint disease, right greater than left, status post total knee replacement (2012), and low back pain secondary to antalgic gait. According to the progress report dated 9-15-2015, the patient had complaints of worse back from limping; left knee giving out more and more and getting worse. The physical examination of the knee revealed effusion, crepitus, patellofemoral grind, tenderness over the medial and lateral joint line, and restricted range of motion. Examination of the lumbar spine revealed tight hamstrings without radiculopathy. The current medications list includes Norco (since at least 4-22-2015). She has undergone right total knee replacement in 7/2012. She has had physical therapy for this injury. Work status is not indicated. The original utilization review (9-28-2015) partially approved a request for Norco #45 (original request was for #60).

### IMR ISSUES, DECISIONS AND RATIONALES

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

**Norco 5/325mg #60:** Upheld

**Claims Administrator guideline:** Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Opioids, criteria for use, Opioids, specific drug list, Weaning of Medications.

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

**Decision rationale:** Norco contains hydrocodone and acetaminophen. Hydrocodone is an opioid analgesic. According to the cited guidelines, "A therapeutic trial of opioids should not be employed until the patient has failed a trial of non-opioid analgesics. Before initiating therapy, the patient should set goals, and the continued use of opioids should be contingent on meeting these goals." The records provided do not specify that that patient has set goals regarding the use of opioid analgesic. The treatment failure with non-opioid analgesics is not specified in the records provided. Other criteria for ongoing management of opioids are: "The lowest possible dose should be prescribed to improve pain and function. Continuing review of overall situation with regard to non-opioid means of pain control. Ongoing review and documentation of pain relief, functional status, appropriate medication use, and side effects. Consider the use of a urine drug screen to assess for the use or the presence of illegal drugs." The records provided do not provide a documentation of response in regards to pain control and objective functional improvement to opioid analgesic for this patient. The continued review of the overall situation with regard to non-opioid means of pain control is not documented in the records provided. As recommended by the cited guidelines a documentation of pain relief, functional status, appropriate medication use, and side effects should be maintained for ongoing management of opioid analgesic, these are not specified in the records provided. The response to lower potency opioid like Tramadol for chronic pain is not specified in the records provided. A recent urine drug screen report is not specified in the records provided. This patient does not meet criteria for ongoing continued use of opioids analgesic. The medical necessity of Norco 5/325mg #60 is not established for this patient, based on the clinical information submitted for this review and the peer reviewed guidelines referenced. If this medication is discontinued, the medication should be tapered, according to the discretion of the treating provider, to prevent withdrawal symptoms.