

Case Number:	CM15-0106394		
Date Assigned:	06/10/2015	Date of Injury:	05/22/2007
Decision Date:	07/13/2015	UR Denial Date:	05/15/2015
Priority:	Standard	Application Received:	06/02/2015

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: California

Certification(s)/Specialty: Physical Medicine & Rehabilitation

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 54 year old female, who sustained an industrial injury on 5/22/2007. Diagnoses include lumbar facet syndrome, spinal/lumbar degenerative disc disease, cervical radiculopathy, cervical disc disorder and head and neck symptoms NEC. Treatment to date has included diagnostics, medications including Soma, Lyrica, methadone, Norco, and topical creams, physical therapy, epidural steroid injections, functional restoration program, and surgical intervention (lumbosacral discectomy and fusion undated). Per the Primary Treating Physician's Progress Report dated 5/04/2015, the injured worker reported lower backache and bilateral lower extremity pain. Pain with medications is rated as 4.5/10 and without medications is rated as 9/10. Physical examination of the lumbar spine revealed restricted ranges of motion and tenderness to palpation on both sides of the paravertebral muscles. Lumbar facet loading was positive on both sides. The plan of care included medications and authorization was requested for Lyrica and MS Contin.

IMR ISSUES, DECISIONS AND RATIONALES

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

Morphine Sulfate Contin 15mg quantity 45: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Morphine Sulfate extended release.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Opioids, page(s) 74-96.

Decision rationale: Pain symptoms and clinical findings remain unchanged for this chronic injury. Submitted documents show no evidence that the treating physician is prescribing opioids in accordance to change in pain relief, functional goals with demonstrated improvement in daily activities, decreased in medical utilization or returned to work status. There is no evidence presented of random drug testing or utilization of pain contract to adequately monitor for narcotic safety, efficacy, and compliance. The MTUS provides requirements of the treating physician to assess and document for functional improvement with treatment intervention and maintenance of function that would otherwise deteriorate if not supported. From the submitted reports, there is no demonstrated evidence of specific functional benefit derived from the continuing use of opioids with persistent severe pain for this chronic injury of 2007. In addition, submitted reports have not adequately demonstrated the specific indication to support for chronic opioid use without acute flare-up, new injuries, or progressive clinical deficits to support for chronic opioids outside recommendations of the guidelines. The Morphine Sulfate Contin 15mg quantity 45 is not medically necessary and appropriate.