

Case Number:	CM15-0048054		
Date Assigned:	04/15/2015	Date of Injury:	01/06/1994
Decision Date:	05/14/2015	UR Denial Date:	03/12/2015
Priority:	Standard	Application Received:	03/13/2015

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: Georgia

Certification(s)/Specialty: Anesthesiology, Pain Management

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 (IW) is a 57 year old female who sustained an industrial injury on 01/06/1994. She reported back pain. The injured worker was diagnosed as having post-laminectomy pain syndrome from a L5-S1 fusion and an additional L4-L5 fusions. Treatment to date has included serial left transforaminal epidural steroid injections most recently in 09/11/2014 with initial 100% relief in LLE. Six weeks later she is experiencing 50% reduction in pain. Her medications (listed as Vicodin, Ambien, and Lidoderm) are still working well. She is having difficulty getting her medications on time. Currently, the injured worker complains of hip pain, leg pain, and lower back pain. The current requests are for Percocet tablets and Lidoderm patches.

IMR ISSUES, DECISIONS AND RATIONALES

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

Percocet 5/325 mg #90: 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): 79.

Decision rationale: Percocet 5/325 mg #90 is not medically necessary. Per MTUS Page 79 of MTUS guidelines states that weaning of opioids are recommended if (a) there are no overall improvement in function, unless there are extenuating circumstances (b) continuing pain with evidence of intolerable adverse effects (c) decrease in functioning (d) resolution of pain (e) if serious non-adherence is occurring (f) the patient requests discontinuing. The claimant's medical records did not document that there was an overall improvement in function or a return to work with previous opioid therapy. The claimant has long-term use with this medication and there was a lack of improved function with this opioid; therefore the requested medication is not medically necessary.

Lidoderm patch #60: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Topical Analgesics.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Topical Analgesic Page(s): 111-112.

Decision rationale: Lidoderm patch #60 is not medically necessary. According to California MTUS, 2009, chronic pain, page 111 California MTUS guidelines does not cover topical analgesics that are largely experimental in use with a few randomized controlled trials to determine efficacy or safety. Any compounded product that contains at least one drug or drug class that is not recommended, is not recommended. Additionally, Per CA MTUS page 111 states that topical analgesics are "recommended for localized peripheral pain after there has been evidence of a trial of first-line therapy (anti-depressants or AED)." Only FDA-approved products are currently recommended. Non-neuropathic pain: Not recommended. The claimant was not diagnosed with neuropathic pain and there is no documentation of physical findings or diagnostic imaging confirming the diagnosis; therefore, the compounded mixture is not medically necessary.