

Case Number:	CM15-0047120		
Date Assigned:	03/19/2015	Date of Injury:	02/12/2001
Decision Date:	04/20/2015	UR Denial Date:	02/12/2015
Priority:	Standard	Application Received:	03/12/2015

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 (IW) is a 53 year old female who sustained an industrial injury on 02/12/2001. She reported back pain. The injured worker was diagnosed as having post laminectomy syndrome to the cervical, thoracic and lumbar spine. Treatments to date have included multiple spine surgery and placement of a spinal cord stimulator. Currently, the injured worker complains of back pain radiating to both lower extremities, worse on the right than the left with associated numbness, tingling and weakness. The spinal cord stimulator is not connected post thoracic spine surgery. She has trialed multiple non-narcotic drugs, but requires high dose opioid therapy to maintain function. A 02/04/2015 visit is planned to the specialist to discuss reconnecting the spinal stimulator. Requests for authorization were made for Fentanyl Patch 100 MCG/HR #15, Lidocaine 5 Percent Patch #90 with 5 Refills, Zolpidem ER 12.5 MG #30 with 3 Refills, and Gabapentin 800 MG.

IMR ISSUES, DECISIONS AND RATIONALES

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

Lidocaine 5 Percent Patch #90 with 5 Refills: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines topical analgesics Page(s): 111.

Decision rationale: The California chronic pain medical treatment guidelines section on topical lidocaine states: Lidocaine Indication: Neuropathic pain Recommended for localized peripheral pain after there has been evidence of a trial of first-line therapy (tri-cyclic or SNRI anti-depressants or an AED such as gabapentin or Lyrica). Topical lidocaine, in the formulation of a dermal patch (Lidoderm) has been designated for orphan status by the FDA for neuropathic pain. Lidoderm is also used off-label for diabetic neuropathy. No other commercially approved topical formulations of lidocaine (whether creams, lotions or gels) are indicated for neuropathic pain. Non-dermal patch formulations are generally indicated as local anesthetics and anti-pruritics. Further research is needed to recommend this treatment for chronic neuropathic pain disorders other than post-herpetic neuralgia. Formulations that do not involve a dermal-patch system are generally indicated as local anesthetics and anti-pruritics. In February 2007 the FDA notified consumers and healthcare professionals of the potential hazards of the use of topical lidocaine. Those at particular risk were individuals that applied large amounts of this substance over large areas, left the products on for long periods of time, or used the agent with occlusive dressings. Systemic exposure was highly variable among patients. Only FDA-approved products are currently recommended. (Argoff, 2006) (Dworkin, 2007) (Khaliq-Cochrane, 2007) (Knotkova, 2007) (Lexi-Comp, 2008) Non-neuropathic pain: Not recommended. There is only one trial that tested 4% lidocaine for treatment of chronic muscle pain. The results showed there was no superiority over placebo. (Scudds, 1995) This medication is recommended for localized peripheral pain. The patient has had a trial of the Gabapentin. The patient has not had failure of other first line neuropathic pain medications. Therefore, criteria as set forth by the California MTUS as outlined above have not been met and the request is not certified.