

Case Number:	CM14-0203268		
Date Assigned:	12/15/2014	Date of Injury:	06/17/2013
Decision Date:	02/06/2015	UR Denial Date:	11/26/2014
Priority:	Standard	Application Received:	12/04/2014

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. The expert reviewer is Board Certified in Neurology, has a subspecialty in Neuromuscular Medicine, and is licensed to practice in New Jersey. 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/services. He/she is familiar with governing laws and regulations, including the strength of evidence hierarchy that applies to Independent Medical Review determinations.

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 patient is a 29 year old woman who sustained a work-related injury on June 17, 2013 . Subsequently, the patient developed a chronic . According to a progress report dated on August 17 2013 , the patient was complaining of ongoing back pain which improved with the pain medications and topical analgesics. The pain severity was rated 6-8/10 . The patient physical examination demonstrated lumbar tenderness with reduced range of motion. The patient nor her examination was normal with negative straight leg raising. The patient was diagnosed with sprain neck, sprain of the lumbar spine, chronic pain syndrome and knee contusion. The provider requested authorization for the following therapies.

IMR ISSUES, DECISIONS AND RATIONALES

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

8 Acupuncture visits for the lumbar spine: Upheld

Claims Administrator guideline: Decision based on MTUS Acupuncture Treatment Guidelines.

MAXIMUS guideline: Decision based on MTUS Acupuncture Treatment Guidelines.

Decision rationale: According to MTUS guidelines, "Acupuncture is used as an option when pain medication is reduced or not tolerated, it may be used as an adjunct to physical rehabilitation and/or surgical intervention to hasten functional recovery. It is the insertion and

removal of filiform needles to stimulate acupoints (acupuncture points). Needles may be inserted, manipulated, and retained for a period of time. Acupuncture can be used to reduce pain, reduce inflammation, increase blood flow, increase range of motion, decrease the side effect of medication-induced nausea, promote relaxation in an anxious patient, and reduce muscle spasm." Furthermore and according to MTUS guidelines, "Acupuncture with electrical stimulation" is the use of electrical current (microamperage or milli-amperage) on the needles at the acupuncture site. It is used to increase effectiveness of the needles by continuous stimulation of the acupoint. Physiological effects (depending on location and settings) can include endorphin release for pain relief, reduction of inflammation, increased blood circulation, analgesia through interruption of pain stimulus, and muscle relaxation. It is indicated to treat chronic pain conditions, radiating pain along a nerve pathway, muscle spasm, inflammation, scar tissue pain, and pain located in multiple sites. The patient developed chronic back pain and musculoskeletal disorders. She is a candidate for treatment with acupuncture. However, the frequency of the treatment should be reduced from 8 to 3 or less sessions. More sessions will be considered when functional and objective improvements are documented. Therefore, the request for 8 acupuncture visits for the lumbar spine is not medically necessary.

TENS unit with supplies: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Transcutaneous electrotherapy Page(s): 114.

Decision rationale: According to MTUS guidelines, TENS is not recommended as primary treatment modality, but a one month based trial may be considered, if used as an adjunct to a functional restoration program. There is no evidence that a functional restoration program is planned for this patient. There is no recent documentation of recent flare of neuropathic pain. There is no strong evidence supporting the benefit of TENS for back disorders. Therefore, the request for TENS unit with supplies is not medically necessary.

Nucynta 50 mg QID #120: Upheld

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

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Criteria for use of opioids Page(s): 76-79.

Decision rationale: According to MTUS guidelines, ongoing use of opioids should follow specific rules:<(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: currentpain; 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 non-adherent) 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.>There is no clear evidence and documentation from the patient file, of a continuous need for Nucynta. There is no documentation of functional improvement with previous use of Nucynta. There is no documentation of compliance of the patient with his medications. Therefore, the prescription of Nucynta 50 mg QID #120 is not medically necessary.