

Case Number:	CM15-0207315		
Date Assigned:	10/26/2015	Date of Injury:	07/27/2015
Decision Date:	12/11/2015	UR Denial Date:	09/25/2015
Priority:	Standard	Application Received:	10/21/2015

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, District of Columbia, Maryland
 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:

This is a 34 year old male who sustained an industrial injury on 7-27-2015. A review of the medical records indicates that the injured worker is undergoing treatment for anxiety, bilateral carpal tunnel syndrome, radiculopathy, cervical sprain-strain, bilateral knee sprain-strain, lumbar sprain-strain and bilateral wrist sprain-strain. According to the progress report dated 9-16-2015, the injured worker complained of neck pain, low back pain, bilateral wrist pain and bilateral knee pain. The pain was rated 7 out of 10. Objective findings (9-16-2015) revealed decreased cervical spine range of motion with pain. There was tenderness to palpation and spasm of the bilateral upper trapezius muscles. There was decreased range of motion of the lumbar spine with pain and positive straight leg raise bilaterally. There was decreased range of motion of the bilateral wrists and hands with positive Phalen's test bilaterally, decreased median nerve sensation and tenderness to palpation. There was decreased range of motion of the bilateral knees with pain and tenderness to palpation. Treatment has included medications. Current medications (9-16-2015) included Pantoprazole, Cyclobenzaprine and Diclofenac; topical creams were ordered. The request for authorization was dated 9-16-2015. The original Utilization Review (UR) (9-25-2015) denied requests for HNPC1-Amitriptyline HCL 10%, Gabapentin 10%, Bupivacaine HCL 5%, Hyaluronic acid 0.2% in cream base, 240gms, 30 day supply and HMPC2-Flurbiprofen 20%, Baclofen 10%, Dexamethasone micro 0.2%, Hyaluronic acid 0.2% in cream base, 240gms, 30 day supply.

IMR ISSUES, DECISIONS AND RATIONALES

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

HNPC1-Amitriptyline HCL 10%, Gabapentin 10%, Bupivacaine HCL 5%, Hyaluronic acid 0.2% in cream base, 240 gms, 30 day supply: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Topical Analgesics.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Topical Analgesics.

Decision rationale: Per MTUS p113 with regard to topical gabapentin: "Not recommended. There is no peer-reviewed literature to support use." Note the statement on page 111: Any compounded product that contains at least one drug (or drug class) that is not recommended is not recommended. The MTUS is silent on the use of topical Bupivacaine, however, topical lidocaine is only recommended for neuropathic 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). There is no documentation that the injured worker has failed trial of these first-line therapies. Per the article "Topical Analgesics in the Management of Acute and Chronic Pain" published in Mayo Clinic Proceedings (Vol 88, Issue 2, p 195-205), "Studies in healthy volunteers demonstrated that topical amitriptyline at concentrations of 50 and 100 mmol/L produced a significant analgesic effect ($P < .05$) when compared with placebo and was associated with transient increases in tactile and mechanical nociceptive thresholds." Amitriptyline may be indicated. The CA MTUS, ODG, National Guidelines Clearinghouse, and ACOEM provide no evidence-based recommendations regarding the topical application of hyaluronic acid. It is the opinion of this IMR reviewer that a lack of endorsement, a lack of mention, inherently implies a lack of recommendation, or a status equivalent to "not recommended". Regarding the use of multiple medications, MTUS p60 states "Only one medication should be given at a time, and interventions that are active and passive should remain unchanged at the time of the medication change. A trial should be given for each individual medication. Analgesic medications should show effects within 1 to 3 days, and the analgesic effect of antidepressants should occur within 1 week. A record of pain and function with the medication should be recorded. (Mens, 2005) The recent AHRQ review of comparative effectiveness and safety of analgesics for osteoarthritis concluded that each of the analgesics was associated with a unique set of benefits and risks, and no currently available analgesic was identified as offering a clear overall advantage compared with the others." Therefore, it would be optimal to trial each medication individually. As gabapentin is not recommended, the compound is not recommended. The request is not medically necessary.

HMPC2-Flurbiprofen 20%, Baclofen 10%, Dexamethasone micro 0.2%, Hyaluronic acid 0.2% in cream base, 240 gms, 30 day supply: Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Topical Analgesics.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Medical Treatment 2009, Section(s): Topical Analgesics.

Decision rationale: Per MTUS with regard to Flurbiprofen (p112), "These medications may be useful for chronic musculoskeletal pain, but there are no long-term studies of their effectiveness or safety. (Mason, 2004) Indications: Osteoarthritis and tendinitis, in particular, that of the knee and elbow or other joints that are amenable to topical treatment: Recommended for short-term use (4-12 weeks). There is little evidence to utilize topical NSAIDs for treatment of osteoarthritis of the spine, hip or shoulder." Flurbiprofen may be indicated. Per MTUS p 112 "Indications: There are positive randomized studies with capsaicin cream in patients with osteoarthritis, fibromyalgia, and chronic non-specific back pain, but it should be considered experimental in very high doses. Although topical capsaicin has moderate to poor efficacy, it may be particularly useful (alone or in conjunction with other modalities) in patients whose pain has not been controlled successfully with conventional therapy." Per MTUS p113 with regard to topical baclofen, "Baclofen: Not recommended. There is currently one Phase III study of Baclofen- Amitriptyline-Ketamine gel in cancer patients for treatment of chemotherapy-induced peripheral neuropathy. There is no peer-reviewed literature to support the use of topical baclofen. Other muscle relaxants: There is no evidence for use of any other muscle relaxant as a topical product." Baclofen is not indicated. The CA MTUS, ODG, National Guidelines Clearinghouse, and ACOEM provide no evidence-based recommendations regarding the topical application of dexamethasone or hyaluronic acid. It is the opinion of this IMR reviewer that a lack of endorsement, a lack of mention, inherently implies a lack of recommendation, or a status equivalent to "not recommended". Since multiple agents are not medically indicated, then the overall product is not indicated per MTUS as outlined below. Regarding the use of multiple medications, MTUS p60 states "Only one medication should be given at a time, and interventions that are active and passive should remain unchanged at the time of the medication change. A trial should be given for each individual medication. Analgesic medications should show effects within 1 to 3 days, and the analgesic effect of antidepressants should occur within 1 week. A record of pain and function with the medication should be recorded. (Mens, 2005) The recent AHRQ review of comparative effectiveness and safety of analgesics for osteoarthritis concluded that each of the analgesics was associated with a unique set of benefits and risks, and no currently available analgesic was identified as offering a clear overall advantage compared with the others." Therefore, it would be optimal to trial each medication individually. Note the statement on page 111: Any compounded product that contains at least one drug (or drug class) that is not recommended is not recommended. As topical baclofen is not indicated, the requested compound is not medically necessary.