

Case Number:	CM15-0133270		
Date Assigned:	07/21/2015	Date of Injury:	07/25/2013
Decision Date:	08/17/2015	UR Denial Date:	06/27/2015
Priority:	Standard	Application Received:	07/09/2015

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: New Jersey, Alabama, California
 Certification(s)/Specialty: Neurology, Neuromuscular Medicine

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 49-year-old female who sustained an industrial injury on 07/25/2013 due to a bus collision. Diagnoses include patellar tendinitis; facet arthropathy versus sacroiliac joint; and cervical strain. Treatment to date has included medications, physical therapy, cold application and splinting/bracing. According to the progress notes dated 5/26/15, the IW reported continued neck pain rated 6/10 with associated bilateral arm pain, greater on the right; and low back pain rated 7/10 with associated right leg numbness and tingling. She also complained of buttock pain and neck stiffness. She reported Baclofen was helping with spasms, but her pain was recently worse due to the weather. On examination, the right patellar tendon, right upper trapezius and cervical paraspinal muscles were tender to palpation. Finkelstein's test was positive (left arm). Range of motion (ROM) of the cervical spine was: flexion 30, extension 30, right bending 10, left bending 30 and bilateral rotation 60. ROM of the lumbar spine was: flexion 80, extension 20, lateral bending 20, bilaterally and rotation 25 bilaterally. Hand grip strength was 12/10 on the right and 26/22 on the left. Medications were listed as Naproxen 500mg twice daily (#60), Baclofen 10mg and Gabapentin 100mg twice daily. A request was made for Naproxen 500mg, quantity unspecified.

IMR ISSUES, DECISIONS AND RATIONALES

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

Naproxen (no dosage or quantity): Upheld

Claims Administrator guideline: Decision based on MTUS Chronic Pain Treatment Guidelines Page(s): 22, 68.

MAXIMUS guideline: Decision based on MTUS Chronic Pain Treatment Guidelines non selective NSAIDS Page(s): 72.

Decision rationale: Naproxen (Naprosyn): delayed release (EC-Naprosyn), as Sodium salt (Anaprox, Anaprox DS, Aleve [otc]) Generic available; extended-release (Naprelan): 375 mg. Different dose strengths and formulations of the drug are not necessarily bioequivalent. Dosing Information: Osteoarthritis or ankylosing spondylitis: Dividing the daily dose into 3 doses versus 2 doses for immediate-release and delayed-release formulations generally does not affect response. Morning and evening doses do not have to be equal in size. The dose may be increased to 1500 mg/day of naproxyn for limited periods when a higher level of analgesic/anti-inflammatory activity is required (for up to 6 months). Naprosyn or naproxyn: 250-500 mg PO twice daily. Anaprox: 275-550 mg PO twice daily. (total dose may be increased to 1650 mg a day for limited periods). EC-Naprosyn: 375 mg or 500 mg twice daily. The tablet should not be broken, crushed or chewed to maintain integrity of the enteric coating. Naprelan: Two 375 mg tablets (750 mg) PO once daily or two 500 mg tablets (1000 mg) once daily. If required (and a lower dose was tolerated) Naprelan can be increased to 1500 mg once daily for limited periods (when higheranalgesia is required). Pain: Naprosyn or naproxyn: 250-500 mg PO twice daily. The maximum dose on day one should not exceed 1250 mg and 1000 mg on subsequent days. Anaprox: 275-550 mg PO twice daily. The maximum dose on day one should not exceed 1375 mg and 1100 mg on subsequent days. Extended-release Naprelan: Not recommended due to delay in absorption. (Naprelan Package Insert) There is no documentation of the rationale behind the long-term use of Naproxen. NSAID should be used for the shortest duration and the lowest dose. There is no documentation from the patient file that the provider titrated Naproxen to the lowest effective dose and used it for the shortest period possible. Naproxen was used without clear documentation of its efficacy. Furthermore, there is no documentation that the provider followed the patient for NSAID adverse reactions that are not limited to GI side effect, but also may affect the renal function. The request is not medically necessary.