

<b>Case Number:</b>	CM15-0080476		
<b>Date Assigned:</b>	05/29/2015	<b>Date of Injury:</b>	10/23/2000
<b>Decision Date:</b>	06/25/2015	<b>UR Denial Date:</b>	04/15/2015
<b>Priority:</b>	Standard	<b>Application Received:</b>	04/27/2015

### 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: Arizona, California

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 is a 49-year-old male who sustained an industrial injury on 10/23/00. Initial complaints and diagnoses are not available. Treatments to date include medications. Diagnostic studies are not addressed. Current complaints include chronic low back pain and numbness and tingling down the left leg. Current diagnoses include chronic low back pain. In a progress note dated 03/17/15 the treating provider reports the plan of care as a MRI of the lumbar spine, and medications including Tylenol, Soma, Prilosec, and Oxycodone. The requested treatments include Soma.

### IMR ISSUES, DECISIONS AND RATIONALES

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

**Soma 350mg #30:** Upheld

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

**MAXIMUS guideline:** Decision based on MTUS Chronic Pain Treatment Guidelines SOMA Page(s): 29.

**Decision rationale:** According to the MTUS guidelines, SOMA is not recommended. Soma is a commonly prescribed, centrally acting skeletal muscle relaxant whose primary active metabolite is meprobamate (a schedule-IV controlled substance). Abuse has been noted for sedative and relaxant effects. As a combination with hydrocodone, an effect that some abusers claim is similar to heroin. In this case, it was combined with Oxycodone and Oxycontin which increases side effect risks and abuse potential. The use of SOMA is not medically necessary.