



<b>ISSUE DATE</b> October 22, 2012	<b>EFFECTIVE DATE</b> November 13, 2012	<b>NUMBER</b> *See below
<b>SUBJECT</b>  Prior Authorization of Botulinum Toxins – Pharmacy Services	<b>BY</b>   Vincent D. Gordon, Deputy Secretary Office of Medical Assistance Programs	

**PURPOSE:**

The purpose of this bulletin is to issue updated handbook pages that include instructions on how to request prior authorization of prescriptions for Botulinum Toxins that require prior authorization, including the type of medical information needed to evaluate requests for medical necessity.

**SCOPE:**

This bulletin applies to all licensed pharmacies and prescribers enrolled in the Medical Assistance (MA) Program and providing services in the fee-for-service (FFS) delivery system, including pharmacy services to residents of long term care facilities.

**BACKGROUND:**

The Department of Public Welfare’s (Department) Drug Utilization Review (DUR) Board meets semi-annually to review provider prescribing and dispensing practices for efficacy, safety, and quality and to recommend interventions for prescribers and pharmacists through the Department’s Prospective Drug Use Review (ProDUR) and Retrospective Drug Use Review (RetroDUR) programs.

**DISCUSSION:**

During the September 19, 2012 meeting, the DUR Board recommended that the Department revise the guidelines to determine medical necessity of Botulinum Toxins when prescribed for the treatment of urinary incontinence due to detrusor over activity associated with a neurologic condition. The guidelines to determine medical necessity of Botulinum

*01-12-53	09-12-50	27-12-46	33-12-48
02-12-44	11-12-43	30-12-46	
03-12-44	14-12-44	31-12-52	
08-12-49	24-12-45	32-12-44	

**COMMENTS AND QUESTIONS REGARDING THIS BULLETIN SHOULD BE DIRECTED TO:**

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Toxins, as recommended by the DUR Board, were subject to public review and comment, and subsequently approved for implementation by the Department. The requirements for prior authorization and clinical review guidelines to determine the medical necessity of Botulinum Toxins are included in the attached updated provider handbook pages.

**PROCEDURE:**

The procedures for prescribers to request prior authorization of Botulinum Toxins are located in SECTION I of the Prior Authorization of Pharmaceutical Services Handbook. The Department will take into account the elements specified in the clinical review guidelines (which are included in the provider handbook pages in the SECTION II chapter related to Botulinum Toxins) in reviewing the prior authorization request to determine medical necessity.

As set forth in 55 Pa. Code § 1101.67(a), the procedures described in the handbook pages must be followed to ensure appropriate and timely processing of prior authorization requests for drugs that require prior authorization.

**ATTACHMENTS:**

[Prior Authorization of Pharmaceutical Services Handbook - Updated pages](#)  
[SECTION II](#)  
[Botulinum Toxins](#)