

5805--A

2015-2016 Regular Sessions

I N   A S S E M B L Y

March 5, 2015

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Introduced by M. of A. McDONALD, BROOK-KRASNY, STIRPE, STECK, WEPRIN, LIFTON, ORTIZ, CLARK, FAHY, LUPINACCI, WALTER, BRINDISI, KEARNS, OTIS, DUPREY, GOTTFRIED, LAVINE, SIMANOWITZ, LUPARDO, BUTLER, GARBARINO, PEOPLES-STOKES -- Multi-Sponsored by -- M. of A. COOK, DiPIETRO, FITZPATRICK, GALEF, HAWLEY, HOOPER, McLAUGHLIN, SCHIMMINGER, THIELE, TITONE -- read once and referred to the Committee on Higher Education -- committee discharged, bill amended, ordered reprinted as amended and recommitted to said committee

AN ACT to amend the education law, in relation to authorizing pharmacists to perform collaborative drug therapy management, and to amend chapter 21 of the laws of 2011 amending the education law relating to authorizing pharmacists to perform collaborative drug therapy management with physicians in certain settings, in relation to extending such provisions

THE PEOPLE OF THE STATE OF NEW YORK, REPRESENTED IN SENATE AND ASSEMBLY, DO ENACT AS FOLLOWS:

1     Section 1. Section 6801-a of the education law, as added by chapter 21  
2     of the laws of 2011, is amended to read as follows:  
3     S    6801-a.    Collaborative drug therapy management demonstration  
4     program. 1. As used in this section, the following terms shall have the  
5     following meanings:  
6     a. "BOARD" SHALL MEAN THE STATE BOARD OF PHARMACY AS ESTABLISHED BY  
7     SECTION SIXTY-EIGHT HUNDRED FOUR OF THIS ARTICLE.  
8     B. "CLINICAL SERVICES" SHALL MEAN THE COLLECTION AND INTERPRETATION  
9     OF PATIENT DATA FOR THE PURPOSE OF INITIATING, MODIFYING AND MONITORING  
10    DRUG THERAPY WITH ASSOCIATED ACCOUNTABILITY AND RESPONSIBILITY FOR  
11    OUTCOMES IN A DIRECT PATIENT CARE SETTING.  
12    C. "Collaborative drug therapy management" shall mean the performance  
13    of CLINICAL services by a pharmacist relating to the review, evaluation  
14    and management of drug therapy to a patient, who is being treated by a  
15    physician for a specific disease or ASSOCIATED disease [state] STATES,  
16    in accordance with a written agreement or protocol with a voluntarily  
17    participating physician and in accordance with the policies, procedures,  
18    and protocols of the facility. Such agreement or protocol as entered

EXPLANATION--Matter in *ITALICS* (underscored) is new; matter in brackets [ ] is old law to be omitted.

LBD06620-12-5

into by the physician and a pharmacist, may include, and shall be limited to:

(i) adjusting or managing a drug regimen of a patient, pursuant to a patient specific [written] order or protocol made by the patient's physician, which may include adjusting drug strength, frequency of administration or route of administration. Adjusting the drug regimen shall not include substituting or selecting a different drug which differs from that initially prescribed by the patient's physician unless such substitution is expressly authorized in the written order or protocol. The pharmacist shall be required to immediately [enter into] DOCUMENT IN the patient record [any change or] changes made to the patient's drug therapy and shall use any reasonable means or method established by the facility [or the department] to notify [any of] the patient's other treating physicians with whom he or she does not have a written agreement or protocol regarding such changes. The patient's physician may prohibit, by written instruction, any adjustment or change in the patient's drug regimen by the pharmacist;

(ii) evaluating and, only if specifically authorized by the protocol and only to the extent necessary to discharge the responsibilities set forth in this section, ordering [clinical] DISEASE STATE laboratory tests related to the drug therapy management for the specific disease or disease state specified within the WRITTEN AGREEMENT OR protocol; and

(iii) only if specifically authorized by the WRITTEN AGREEMENT OR protocol and only to the extent necessary to discharge the responsibilities set forth in this section, ordering or performing routine patient monitoring functions as may be necessary in the drug therapy management, including the collecting and reviewing of patient histories, and ordering or checking patient vital signs, including pulse, temperature, blood pressure and respiration.

[b. "Written agreement or protocol" shall mean a written document, pursuant to and consistent with any applicable state or federal requirements, that addresses a specific disease or disease state and that describes the nature and scope of collaborative drug therapy management to be undertaken by the pharmacist, in collaboration with the participating physician, in accordance with the provisions of this section.

c. "Physician" shall mean the physician, selected by or assigned to a patient, who has primary responsibility for the treatment and care of the patient for the disease or disease state that is the subject of the collaborative drug therapy management.]

d. "Facility" shall mean: (I) a teaching hospital OR GENERAL HOSPITAL, including any diagnostic center, treatment center, or hospital-based outpatient department AS DEFINED IN SECTION TWENTY-EIGHT HUNDRED ONE OF THE PUBLIC HEALTH LAW; OR (II) A NURSING HOME WITH AN ON-SITE PHARMACY STAFFED BY A LICENSED PHARMACIST; PROVIDED, however, for the purposes of this section THE TERM "FACILITY" SHALL NOT INCLUDE DENTAL CLINICS, DENTAL DISPENSARIES, residential health care facilities and [nursing homes shall be excluded] REHABILITATION CENTERS.

For the purposes of this section, a "teaching hospital" shall mean a hospital licensed pursuant to article twenty-eight of the public health law that is eligible to receive direct or indirect graduate medical education payments pursuant to article twenty-eight of the public health law.

E. "PHYSICIAN" SHALL MEAN THE PHYSICIAN SELECTED BY OR ASSIGNED TO A PATIENT, WHO HAS PRIMARY RESPONSIBILITY FOR THE TREATMENT AND CARE OF THE PATIENT FOR THE DISEASE AND ASSOCIATED DISEASE STATES THAT ARE THE SUBJECT OF THE COLLABORATIVE DRUG THERAPY MANAGEMENT.

F. "WRITTEN AGREEMENT OR PROTOCOL" SHALL MEAN A WRITTEN DOCUMENT, PURSUANT TO AND CONSISTENT WITH ANY APPLICABLE STATE OR FEDERAL REQUIREMENTS, THAT ADDRESSES A SPECIFIC DISEASE OR ASSOCIATED DISEASE STATES AND THAT DESCRIBES THE NATURE AND SCOPE OF COLLABORATIVE DRUG THERAPY MANAGEMENT TO BE UNDERTAKEN BY THE PHARMACISTS, IN COLLABORATION WITH THE PARTICIPATING PHYSICIAN IN ACCORDANCE WITH THE PROVISIONS OF THIS SECTION.

2. a. A pharmacist who meets the experience requirements of paragraph b of this subdivision and who is employed by or otherwise affiliated with a facility shall be permitted to enter into a written agreement or protocol with a physician authorizing collaborative drug therapy management, subject to the limitations set forth in this section, within the scope of such employment or affiliation.

b. A participating pharmacist must:

(i)(A) have been awarded either a master of science in clinical pharmacy or a doctor of pharmacy degree;

(B) maintain a current unrestricted license; and

(C) have a minimum of two years experience, of which at least one year of such experience shall include clinical experience in a health facility, which involves consultation with physicians with respect to drug therapy and may include a residency at a facility involving such consultation; or

(ii)(A) have been awarded a bachelor of science in pharmacy;

(B) maintain a current unrestricted license; and

(C) within the last seven years, have a minimum of three years experience, of which at least one year of such experience shall include clinical experience in a health facility, which involves consultation with physicians with respect to drug therapy and may include a residency at a facility involving such consultation; AND

(III) MEET ANY ADDITIONAL EDUCATION, EXPERIENCE, OR OTHER REQUIREMENTS SET FORTH BY THE DEPARTMENT IN CONSULTATION WITH THE BOARD.

c. NOTWITHSTANDING ANY PROVISION OF LAW, NOTHING IN THIS SECTION SHALL PROHIBIT A LICENSED PHARMACIST FROM ENGAGING IN CLINICAL SERVICES ASSOCIATED WITH COLLABORATIVE DRUG THERAPY MANAGEMENT, IN ORDER TO GAIN EXPERIENCE NECESSARY TO QUALIFY UNDER CLAUSE (C) OF SUBPARAGRAPH (I) OR (II) OF PARAGRAPH B OF THIS SUBDIVISION, PROVIDED THAT SUCH PRACTICE IS UNDER THE SUPERVISION OF A PHARMACIST THAT CURRENTLY MEETS THE REFERENCED REQUIREMENT, AND THAT SUCH PRACTICE IS AUTHORIZED UNDER THE WRITTEN AGREEMENT OR PROTOCOL WITH THE PHYSICIAN.

D. Notwithstanding any provision of this section, nothing herein shall authorize the pharmacist to diagnose disease. In the event that a treating physician may disagree with the exercise of professional judgment by [the] A pharmacist, the judgment of the treating physician shall prevail.

3. The physician who is a party to a written agreement or protocol authorizing collaborative drug therapy management shall be employed by or otherwise affiliated with the same facility with which the pharmacist is also employed or affiliated.

4. The existence of a written agreement or protocol on collaborative drug therapy management and the patient's right to choose to not participate in collaborative drug therapy management shall be disclosed to any patient who is eligible to receive collaborative drug therapy management. Collaborative drug therapy management shall not be utilized unless the patient or the patient's authorized representative consents, in writing, to such management. If the patient or the patient's authorized representative consents, it shall be noted on the patient's medical

1 record. If the patient or the patient's authorized representative who  
2 consented to collaborative drug therapy management chooses to no longer  
3 participate in such management, at any time, it shall be noted on the  
4 patient's medical record. In addition, the existence of the written  
5 agreement or protocol and the patient's consent to such management shall  
6 be disclosed to the patient's primary physician and any other treating  
7 physician or healthcare provider.

8 5. Participation in a written agreement or protocol authorizing colla-  
9 borative drug therapy management shall be voluntary, and no patient,  
10 physician, pharmacist, or facility shall be required to participate.

11 6. Nothing in this section shall be deemed to limit the scope of prac-  
12 tice of pharmacy nor be deemed to limit the authority of pharmacists and  
13 physicians to engage in medication management prior to the effective  
14 date of this section and to the extent authorized by law.

15 S 2. The department of education, in consultation with the department  
16 of health, shall prepare or shall arrange for the preparation of a  
17 report on the impact of collaborative drug therapy management (CDTM) in  
18 New York state. The report shall be submitted to the speaker of the  
19 assembly and the temporary president of the senate and the chairs of the  
20 senate and assembly higher education committees at least four months  
21 prior to the expiration of this act. The report shall review the extent  
22 to which the continued application and expansion of CDTM contributed to  
23 the following: (i) patient health-related outcomes; (ii) quality of care  
24 for patients; (iii) reduced risk of medication error; (iv) health care  
25 expenditures. The report may make recommendations regarding the exten-  
26 sion, alteration and/or expansion of these provisions, which shall  
27 include recommendations for addressing any barriers to further implemen-  
28 tation of CDTM, and make any other recommendations related to the imple-  
29 mentation of CDTM pursuant to this act.

30 S 3. Section 5 of chapter 21 of the laws of 2011, amending the educa-  
31 tion law relating to authorizing pharmacists to perform collaborative  
32 drug therapy management with physicians in certain settings, as amended  
33 by chapter 125 of the laws of 2014, is amended to read as follows:

34 S 5. This act shall take effect on the one hundred twentieth day after  
35 it shall have become a law and shall expire [4] 7 years after such  
36 effective date when upon such date the provisions of this act shall be  
37 deemed repealed; provided, however, that the amendments to subdivision 1  
38 of section 6801 of the education law made by section one of this act  
39 shall be subject to the expiration and reversion of such subdivision  
40 pursuant to section 8 of chapter 563 of the laws of 2008, when upon such  
41 date the provisions of section one-a of this act shall take effect;  
42 provided, further, that effective immediately, the addition, amendment  
43 and/or repeal of any rule or regulation necessary for the implementation  
44 of this act on its effective date is authorized and directed to be made  
45 and completed on or before such effective date.

46 S 4. This act shall take effect immediately, provided that section one  
47 of this act shall take effect on the ninetieth day after it shall have  
48 become a law; provided that the amendments to section 6801-a of the  
49 education law, made by section one of this act, shall not affect the  
50 repeal of such section and shall be deemed repealed therewith; provided,  
51 further, that, effective immediately, the addition, amendment and/or  
52 repeal of any rule or regulation necessary for the implementation of  
53 this act on its effective date is authorized and directed to be made and  
54 completed on or before such effective date.