STATE OF NEW YORK

4043

2023-2024 Regular Sessions

IN SENATE

February 2, 2023

Introduced by Sens. RIVERA, GOUNARDES, KRUEGER, SALAZAR -- read twice and ordered printed, and when printed to be committed to the Committee on Higher Education

AN ACT to amend the education law, in relation to including nurse practitioners as a provider of services for purposes of 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 making the authorization for pharmacists to perform collaborative drug therapy management permanent

The People of the State of New York, represented in Senate and Assembly, do enact as follows:

Section 1. Section 6801-a of the education law, as amended by chapter 1 2 238 of the laws of 2015, is amended to read as follows: [demonstration 3 § 6801-a. Collaborative drug therapy management 4 **program**]. 1. As used in this section, the following terms shall have 5 the following meanings: a. "Board" shall mean the state board of pharmacy as established by 6 7 section sixty-eight hundred four of this article. 8 b. "Clinical services" shall mean the collection and interpretation of 9 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. c. "Collaborative drug therapy management" shall mean the performance 12 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 or nurse practitioner for a specific disease or associated 16 disease states, in accordance with a written agreement or protocol with 17 a voluntarily participating physician or nurse practitioner and in 18 accordance with the policies, procedures, and protocols of the facility.

EXPLANATION--Matter in <u>italics</u> (underscored) is new; matter in brackets [-] is old law to be omitted.

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Such agreement or protocol as entered into by the physician or nurse 1 practitioner and a pharmacist, may include, and shall be limited to: 2 3 (i) adjusting or managing a drug regimen of a patient, pursuant to a 4 patient specific order or protocol made by the patient's physician or 5 nurse practitioner, which may include adjusting drug strength, frequency 6 of administration or route of administration. Adjusting the drug regimen 7 shall not include substituting or selecting a different drug which 8 differs from that initially prescribed by the patient's physician or 9 nurse practitioner unless such substitution is expressly authorized in 10 the written order or protocol. The pharmacist shall be required to imme-11 diately document in the patient record changes made to the patient's 12 drug therapy and shall use any reasonable means or method established by facility to notify the patient's other treating physicians or nurse 13 the 14 practitioners with whom he or she does not have a written agreement or 15 protocol regarding such changes. The patient's physician or nurse prac-16 **<u>titioner</u>** may prohibit, by written instruction, any adjustment or change 17 in the patient's drug regimen by the pharmacist; 18 (ii) evaluating and, only if specifically authorized by the protocol 19 and only to the extent necessary to discharge the responsibilities set 20 forth in this section, ordering disease state laboratory tests related 21 to the drug therapy management for the specific disease or disease state 22 specified within the written agreement or protocol; and 23 (iii) only if specifically authorized by the written agreement or 24 protocol and only to the extent necessary to discharge the responsibil-25 ities set forth in this section, ordering or performing routine patient monitoring functions as may be necessary in the drug therapy management, 26 27 including the collecting and reviewing of patient histories, and order-28 ing or checking patient vital signs, including pulse, temperature, blood 29 pressure and respiration. d. "Facility" shall mean: (i) a [teaching hospital or general] hospi-30 tal, [including any diagnostic center, treatment center, or hospital-31 32 **baged outpatient department**] as defined in **subdivision one of** section 33 twenty-eight hundred one of the public health law; or (ii) a nursing home with an on-site pharmacy staffed by a licensed pharmacist; 34 35 provided, however, for the purposes of this section the term "facility" 36 shall not include dental clinics, dental dispensaries, residential 37 health care facilities and rehabilitation centers. 38 [For the purposes of this section, a "teaching hospital" shall mean a 39 hospital licensed pursuant to article twenty-eight of the public health law that is eligible to receive direct or indirect graduate medical 40 education payments pursuant to article twenty-eight of the public health 41 42 law.] In addition, a facility may also include up to fifteen community-43 practice sites, selected by the department in consultation with the 44 department of health, where pharmacists and physicians or nurse practi-45 tioners may propose to enter into collaborative arrangements, pursuant 46 to the provisions of this section. Such sites shall be selected based 47 upon a review of applications submitted to the department by such phar-48 macists and physicians or nurse practitioners, which demonstrate that 49 the applicants can satisfy the requirements of this section. e. "Physician" or "nurse practitioner" shall mean the physician or 50 51 nurse practitioner selected by or assigned to a patient, who has primary 52 responsibility for the treatment and care of the patient for the disease 53 and associated disease states that are the subject of the collaborative 54 drug therapy management.

55 f. "Written agreement or protocol" shall mean a written document, 56 pursuant to and consistent with any applicable state or federal require-

ments, that addresses a specific disease or associated disease states 1 and that describes the nature and scope of collaborative drug therapy 2 3 management to be undertaken by the pharmacists, in collaboration with 4 the participating physician or nurse practitioner in accordance with the 5 provisions of this section. б 2. a. A pharmacist who meets the experience requirements of paragraph 7 b of this subdivision and who is either employed by or otherwise affil-8 iated with a facility or is participating with a community-practice site 9 selected pursuant to paragraph d of subdivision one of this section 10 shall be permitted to enter into a written agreement or protocol with a 11 physician <u>or nurse practitioner</u> authorizing collaborative drug therapy 12 management, subject to the limitations set forth in this section, within 13 the scope of such employment [or], affiliation or participation. 14 b. A participating pharmacist must: 15 (i)(A) have been awarded either a master of science in clinical phar-16 macy or a doctor of pharmacy degree; 17 (B) maintain a current unrestricted license; and 18 (C) have a minimum of two years experience, of which at least one year 19 of such experience shall include clinical experience in a health facili-20 ty, which involves consultation with physicians or nurse practitioners 21 with respect to drug therapy and may include a residency at a facility 22 involving such consultation; or 23 (ii)(A) have been awarded a bachelor of science in pharmacy; 24 (B) maintain a current unrestricted license; and 25 (C) within the last seven years, have a minimum of three years experi-26 ence, of which at least one year of such experience shall include clin-27 ical experience in a health facility, which involves consultation with 28 physicians or nurse practitioners with respect to drug therapy and may 29 include a residency at a facility involving such consultation; and 30 (iii) meet any additional education, experience, or other requirements 31 set forth by the department in consultation with the board. 32 c. Notwithstanding any provision of law, nothing in this section shall 33 prohibit a licensed pharmacist from engaging in clinical services asso-34 ciated with collaborative drug therapy management, in order to gain 35 experience necessary to qualify under clause (C) of subparagraph (i) or 36 (ii) of paragraph b of this subdivision, provided that such practice is 37 under the supervision of a pharmacist that currently meets the refer-38 enced requirement, and that such practice is authorized under the writ-39 ten agreement or protocol with the physician or nurse practitioner. d. Notwithstanding any provision of this section, nothing herein shall 40 authorize the pharmacist to diagnose disease. In the event that a treat-41 42 ing physician or nurse practitioner may disagree with the exercise of 43 professional judgment by a pharmacist, the judgment of the treating 44 physician or nurse practitioner shall prevail. 45 3. The physician or nurse practitioner who is a party to a written 46 agreement or protocol authorizing collaborative drug therapy management 47 shall be employed by or otherwise affiliated with the same facility with 48 which the pharmacist is also employed or affiliated. 49 4. The existence of a written agreement or protocol on collaborative 50 drug therapy management and the patient's right to choose to not partic-51 ipate in collaborative drug therapy management shall be disclosed to any 52 patient who is eligible to receive collaborative drug therapy manage-53 ment. Collaborative drug therapy management shall not be utilized unless 54 the patient or the patient's authorized representative consents, in writing, to such management. If the patient or the patient's authorized 55 representative consents, it shall be noted on the patient's medical 56

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record. If the patient or the patient's authorized representative who 1 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 or nurse practitioner 7 and any other treating physician or nurse practitioner or healthcare 8 provider. 9 5. Participation in a written agreement or protocol authorizing colla-10 borative drug therapy management shall be voluntary, and no patient, 11 physician or nurse practitioner, pharmacist, or facility shall be 12 required to participate. 6. Nothing in this section shall be deemed to limit the scope of prac-13 14 tice of pharmacy nor be deemed to limit the authority of pharmacists and 15 physicians or nurse practitioners to engage in medication management 16 prior to the effective date of this section and to the extent authorized 17 by law. 18 Section 5 of chapter 21 of the laws of 2011 amending the educa-§ 2. 19 tion law relating to authorizing pharmacists to perform collaborative 20 drug therapy management with physicians in certain settings, as amended 21 by section 5 of part CC of chapter 57 of the laws of 2022, is amended to 22 read as follows: 23 § 5. This act shall take effect on the one hundred twentieth day after it shall have become a law[, provided, however, that the provisions of 24 25 sections two, three, and four of this act shall expire and be deemed repealed July 1, 2024]; provided, however, that the amendments to subdi-26 27 vision 1 of section 6801 of the education law made by section one of 28 this act shall be subject to the expiration and reversion of such subdivision pursuant to section 8 of chapter 563 of the laws of 2008, when 29 30 upon such date the provisions of section one-a of this act shall take

31 effect; provided, further, that effective immediately, the addition, 32 amendment and/or repeal of any rule or regulation necessary for the 33 implementation of this act on its effective date are authorized and 34 directed to be made and completed on or before such effective date. 35 § 3. This act shall take effect on the one hundred twentieth day after

it shall have become a law. Effective immediately, the addition, amendment and/or repeal of any rule or regulation necessary for the implementation of this act on its effective date are authorized to be made and completed on or before such effective date.