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HB498E

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16102660D **HOUSE BILL NO. 498** 1 2 House Amendments in [] — January 26, 2016 3 A BILL to amend and reenact § 54.1-3303 of the Code of Virginia, relating to TPA-certified 4 optometrists; prescription of certain Schedule II controlled substances. 5 Patron Prior to Engrossment-Delegate Hodges 6 7 Referred to Committee on Health, Welfare and Institutions 8 9 Be it enacted by the General Assembly of Virginia: 1. That § 54.1-3303 of the Code of Virginia is amended and reenacted as follows: 10 § 54.1-3303. Prescriptions to be issued and drugs to be dispensed for medical or therapeutic 11 12 purposes only. 13 A. A prescription for a controlled substance may be issued only by a practitioner of medicine, 14 osteopathy, podiatry, dentistry or veterinary medicine who is authorized to prescribe controlled 15 substances, or by a licensed nurse practitioner pursuant to § 54.1-2957.01, a licensed physician assistant 16 pursuant to § 54.1-2952.1, or a TPA-certified optometrist pursuant to Article 5 (§ 54.1-3222 et seq.) of Chapter 32. The prescription shall be issued for a medicinal or therapeutic purpose and may be issued 17 18 only to persons or animals with whom the practitioner has a bona fide practitioner-patient relationship. 19 For purposes of this section, a bona fide practitioner-patient-pharmacist relationship is one in which a 20 practitioner prescribes, and a pharmacist dispenses, controlled substances in good faith to his patient for 21 a medicinal or therapeutic purpose within the course of his professional practice. In addition, a bona fide 22 practitioner-patient relationship means that the practitioner shall (i) ensure that a medical or drug history 23 is obtained; (ii) provide information to the patient about the benefits and risks of the drug being 24 prescribed; (iii) perform or have performed an appropriate examination of the patient, either physically 25 or by the use of instrumentation and diagnostic equipment through which images and medical records may be transmitted electronically; except for medical emergencies, the examination of the patient shall 26 27 have been performed by the practitioner himself, within the group in which he practices, or by a 28 consulting practitioner prior to issuing a prescription; and (iv) initiate additional interventions and 29 follow-up care, if necessary, especially if a prescribed drug may have serious side effects. 30 For the purpose of prescribing a Schedule VI controlled substance to a patient via telemedicine 31 services as defined in § 38.2-3418.16, a prescriber may establish a bona fide practitioner-patient relationship by an examination through face-to-face interactive, two-way, real-time communications 32 33 services or store-and-forward technologies when all of the following conditions are met: (a) the patient 34 has provided a medical history that is available for review by the prescriber; (b) the prescriber obtains 35 an updated medical history at the time of prescribing; (c) the prescriber makes a diagnosis at the time of 36 prescribing; (d) the prescriber conforms to the standard of care expected of in-person care as appropriate 37 to the patient's age and presenting condition, including when the standard of care requires the use of 38 diagnostic testing and performance of a physical examination, which may be carried out through the use 39 of peripheral devices appropriate to the patient's condition; (e) the prescriber is actively licensed in the 40 Commonwealth and authorized to prescribe; (f) if the patient is a member or enrollee of a health plan or carrier, the prescriber has been credentialed by the health plan or carrier as a participating provider and 41 42 the diagnosing and prescribing meets the qualifications for reimbursement by the health plan or carrier pursuant to § 38.2-3418.16; and (g) upon request, the prescriber provides patient records in a timely 43 manner in accordance with the provisions of § 32.1-127.1:03 and all other state and federal laws and 44 regulations. Nothing in this paragraph shall permit a prescriber to establish a bona fide 45 practitioner-patient relationship for the purpose of prescribing a Schedule VI controlled substance when 46 the standard of care dictates that an in-person physical examination is necessary for diagnosis. Nothing 47 in this paragraph shall apply to: (1) a prescriber providing on-call coverage per an agreement with 48 49 another prescriber or his prescriber's professional entity or employer; (2) a prescriber consulting with another prescriber regarding a patient's care; or (3) orders of prescribers for hospital out-patients or 50 51 in-patients. 52 Any practitioner who prescribes any controlled substance with the knowledge that the controlled 53 substance will be used otherwise than medicinally or for therapeutic purposes shall be subject to the criminal penalties provided in § 18.2-248 for violations of the provisions of law relating to the 54 55 distribution or possession of controlled substances. B. In order to determine whether a prescription that appears questionable to the pharmacist results 56 57 from a bona fide practitioner-patient relationship, the pharmacist shall contact the prescribing practitioner or his agent and verify the identity of the patient and name and quantity of the drug prescribed. The 58

59 person knowingly filling an invalid prescription shall be subject to the criminal penalties provided in § 18.2-248 for violations of the provisions of law relating to the sale, distribution or possession of

60 § 18.2-248 for violat 61 controlled substances.

No prescription shall be filled unless there is a bona fide practitioner-patient-pharmacist relationship.
 A prescription not issued in the usual course of treatment or for authorized research is not a valid prescription.

65 C. Notwithstanding any provision of law to the contrary and consistent with recommendations of the Centers for Disease Control and Prevention or the Department of Health, a practitioner may prescribe 66 Schedule VI antibiotics and antiviral agents to other persons in close contact with a diagnosed patient 67 when (i) the practitioner meets all requirements of a bona fide practitioner-patient relationship, as 68 69 defined in subsection A, with the diagnosed patient; (ii) in the practitioner's professional judgment, the practitioner deems there is urgency to begin treatment to prevent the transmission of a communicable 70 71 disease; (iii) the practitioner has met all requirements of a bona fide practitioner-patient relationship, as defined in subsection A, for the close contact except for the physical examination required in clause (iii) 72 of subsection A; and (iv) when such emergency treatment is necessary to prevent imminent risk of 73 74 death, life-threatening illness, or serious disability.

D. A pharmacist may dispense a controlled substance pursuant to a prescription of an out-of-state practitioner of medicine, osteopathy, podiatry, dentistry or veterinary medicine authorized to issue such prescription if the prescription complies with the requirements of this chapter and the Drug Control Act (§ 54.1-3400 et seq.).

79 E. A licensed nurse practitioner who is authorized to prescribe controlled substances pursuant to
80 § 54.1-2957.01 may issue prescriptions or provide manufacturers' professional samples for controlled
81 substances and devices as set forth in the Drug Control Act (§ 54.1-3400 et seq.) in good faith to his
82 patient for a medicinal or therapeutic purpose within the scope of his professional practice.

F. A licensed physician assistant who is authorized to prescribe controlled substances pursuant to
§ 54.1-2952.1 may issue prescriptions or provide manufacturers' professional samples for controlled
substances and devices as set forth in the Drug Control Act (§ 54.1-3400 et seq.) in good faith to his
patient for a medicinal or therapeutic purpose within the scope of his professional practice.

G. A TPA-certified optometrist who is authorized to prescribe controlled substances pursuant to 87 88 Article 5 (§ 54.1-3222 et seq.) of Chapter 32 may issue prescriptions in good faith or provide 89 manufacturers' professional samples to his patients for medicinal or therapeutic purposes within the 90 scope of his professional practice for the drugs specified on the TPA-Formulary, established pursuant to 91 § 54.1-3223, which shall be limited to (i) [analgesics included on] Schedule II controlled substances as 92 defined in § 54.1-3448 of the Drug Control Act (§ 54.1-3400 et seq.) consisting of hydrocodone in combination with acetaminophen; (ii) oral analgesics included in Schedules III through VI, as defined in 93 §§ 54.1-3450 and 54.1-3455 of the Drug Control Act (§ 54.1-3400 et seq.), which are appropriate to 94 95 relieve ocular pain, (iii); (iii) other oral Schedule VI controlled substances, as defined in § 54.1-3455 of the Drug Control Act, appropriate to treat diseases and abnormal conditions of the human eye and its 96 97 adnexa, (iii); (iv) topically applied Schedule VI drugs, as defined in § 54.1-3455 of the Drug Control 98 Act_i; and (iv) (v) intramuscular administration of epinephrine for treatment of emergency cases of 99 anaphylactic shock.

H. The requirement for a bona fide practitioner-patient relationship shall be deemed to be satisfied by
 a member or committee of a hospital's medical staff when approving a standing order or protocol for the
 administration of influenza vaccinations and pneumococcal vaccinations in a hospital in compliance with
 § 32.1-126.4.