

THE GENERAL ASSEMBLY OF PENNSYLVANIA

HOUSE BILL

No. 1508 Session of 2017

INTRODUCED BY NESBIT, JUNE 12, 2017

REFERRED TO COMMITTEE ON HEALTH, JUNE 12, 2017

AN ACT

1 Amending the act of September 27, 1961 (P.L.1700, No.699),
 2 entitled "An act relating to the regulation of the practice
 3 of pharmacy, including the sales, use and distribution of
 4 drugs and devices at retail; and amending, revising,
 5 consolidating and repealing certain laws relating thereto,"
 6 providing for professional prescription, administration and
 7 dispensing.

8 The General Assembly of the Commonwealth of Pennsylvania
 9 hereby enacts as follows:

10 Section 1. The act of September 27, 1961 (P.L.1700, No.699),
 11 known as the Pharmacy Act, is amended by adding a section to
 12 read:

13 Section 9.5. Professional Prescription, Administration and
 14 Dispensing.--(a) Except when dispensed or administered directly
 15 to the patient by a practitioner or a practitioner's authorized
 16 agent, other than a pharmacist, to an ultimate user, no
 17 prescription shall be dispensed without an electronic
 18 prescription of a practitioner, except in emergency situations,
 19 as prescribed by the secretary by regulation.

20 (b) No prescription for a controlled substance in Schedule
 21 II of the Controlled Substance, Drug, Device and Cosmetic Act,

1 may be refilled.

2 (c) No prescription for a controlled substance in Schedule
3 III, IV and V of the Controlled Substance, Drug, Device and
4 Cosmetic Act may be filled or refilled more than six months
5 after the date of the prescription or be refilled more than five
6 times after the date of the prescription unless renewed by the
7 practitioner.

8 (d) This section shall not apply to prescriptions which are
9 issued:

10 (1) By a veterinarian.

11 (2) In a circumstance when electronic prescribing is not
12 available due to temporary technological or electrical failure.

13 (3) By a practitioner to be dispensed by a pharmacy located
14 outside this Commonwealth.

15 (4) By a practitioner treating a patient in an emergency
16 department or other health care facility and under a
17 circumstance when, notwithstanding the practitioner's present
18 ability to make an electronic prescription as required by this
19 subsection, the practitioner reasonably determines that it would
20 be impractical for the patient to obtain substances prescribed
21 by electronic prescription in a timely manner and that the delay
22 would adversely impact the patient's medical condition.

23 (5) By a practitioner without Internet access or an
24 electronic health record system. For purposes of this
25 subparagraph:

26 (i) The department shall:

27 (A) Within 90 days of the effective date of this
28 subparagraph, create a standardized form for practitioners to
29 submit to the department indicating that the practitioner will
30 not be electronically prescribing controlled substances in

1 Schedule II, III, IV or V of the Controlled Substance, Drug,
2 Device and Cosmetic Act due to the lack of Internet access or an
3 electronic health record system.

4 (B) Provide notice of the form under clause (A) through the
5 Pennsylvania Bulletin and post the form on the department's
6 publicly accessible Internet website.

7 (ii) A practitioner shall be required to provide notice to
8 the department whenever the exception in this subparagraph no
9 longer applies to the practitioner.

10 (iii) A practitioner who provides services in an emergency
11 department or other health care facility shall not be required
12 to submit the form under subparagraph (i) (A).

13 (e) The department shall promulgate regulations within two
14 years of the effective date of this section relating to the
15 exceptions provided for in subsection (d).

16 (f) As used in this section, the following words and phrases
17 shall have the meanings given to them in this subsection unless
18 the context clearly indicates otherwise:

19 "Controlled Substance, Drug, Device and Cosmetic Act" means
20 the act of April 14, 1972 (P.L.233, No.64), known as "The
21 Controlled Substance, Drug, Device and Cosmetic Act."

22 "Cosmetic" means a substance, excluding soap, which is
23 intended:

24 (1) to be rubbed, poured, sprinkled or sprayed on,
25 introduced into or otherwise applied to the human body or other
26 animal body or any part thereof for cleansing, beautifying,
27 promoting attractiveness or altering the appearance; or

28 (2) for use as a component of a substance under paragraph
29 (1).

30 "Department" means the Department of Health of the

1 Commonwealth.

2 "Practitioner" means any of the following:

3 (1) A physician, osteopath, dentist, veterinarian,
4 pharmacist, podiatrist, nurse, scientific investigator or other
5 person licensed, registered or otherwise permitted to
6 distribute, dispense, conduct research with respect to or to
7 administer a controlled substance, other drug or device in the
8 course of professional practice or research in this
9 Commonwealth.

10 (2) A pharmacy, hospital, clinic or other institution
11 licensed, registered or otherwise permitted to distribute,
12 dispense, conduct research with respect to or to administer a
13 controlled substance, other drug or device in the course of
14 professional practice or research in this Commonwealth.

15 "Secretary" means the Secretary of Health of the
16 Commonwealth.

17 "Ultimate user" means an individual who lawfully possesses a
18 controlled substance, other drug, device or cosmetic for the
19 individual's own use or for the use of a member of the
20 individual's household or for administering to an animal in the
21 individual's care.

22 Section 2. This act shall take effect in 60 days.