SB1844

LRB10214943KMF20298b

102ND GENERAL ASSEMBLY

State of Illinois

2021 and 2022

SB1844

Introduced 2/26/2021, by Sen. Mattie Hunter

SYNOPSIS AS INTRODUCED:

720 ILCS 570/316

Amends the Illinois Controlled Substances Act. Provides that specified requirements also apply to opioid treatment programs that are licensed or certified by the Department of Human Services's Division of Substance Use Prevention and Recovery and are authorized by the federal Drug Enforcement Administration to prescribe Schedule II, III, IV, or V controlled substances for the treatment of opioid use disorders. Requires opioid treatment programs to attempt to obtain written patient consent, document attempts to obtain the written consent, and not transmit information without patient consent. Provides that the documentation obtained shall not be utilized for law enforcement purposes. Provides that treatment of a patient shall not be conditioned upon his or her written consent. Makes other changes.

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A BILL FOR

AN ACT concerning criminal law.

Be it enacted by the People of the State of Illinois, represented in the General Assembly:

Section 5. Findings. The General Assembly finds that:

(1) Prior to August of 2020, the federal Substance Abuse and Mental Health Services Administration (SAMHSA) and the federal Confidentiality of Substance Use Disorder Patient Records set forth at 42 CFR 2, prohibited the sharing of substance use disorder treatment information by opioid treatment programs with prescription monitoring programs.

(2) In August 2020, SAMHSA amended 42 CFR 2 to permit the sharing of substance use disorder treatment information by opioid treatment programs with prescription monitoring programs.

(3) In light of the federal modification to 42 CFR 2 and the protections available under federal and State law and the express requirement of patient consent, the reporting by opioid treatment programs to the prescription monitoring program is permitted and will allow for better coordination of care among treating providers.

Section 10. The Illinois Controlled Substances Act is amended by changing Section 316 as follows:

(720 ILCS 570/316)

Sec. 316. Prescription Monitoring Program.

(a) The Department must provide for a Prescription Monitoring Program for Schedule II, III, IV, and V controlled substances that includes the following components and requirements:

(1) The dispenser must transmit to the central repository, in a form and manner specified by the Department, the following information:

(A) The recipient's name and address.

(B) The recipient's date of birth and gender.

(C) The national drug code number of the controlled substance dispensed.

(D) The date the controlled substance is dispensed.

(E) The quantity of the controlled substance dispensed and days supply.

(F) The dispenser's United States Drug Enforcement Administration registration number.

(G) The prescriber's United States Drug Enforcement Administration registration number.

(H) The dates the controlled substance prescription is filled.

(I) The payment type used to purchase the controlled substance (i.e. Medicaid, cash, third party

insurance).

(J) The patient location code (i.e. home, nursing home, outpatient, etc.) for the controlled substances other than those filled at a retail pharmacy.

(K) Any additional information that may be required by the department by administrative rule, including but not limited to information required for compliance with the criteria for electronic reporting of the American Society for Automation and Pharmacy or its successor.

(2) The information required to be transmitted under this Section must be transmitted not later than the end of the ~~next~~ business day ~~after the date~~ on which a controlled substance is dispensed, or at such other time as may be required by the Department by administrative rule.

(3) A dispenser must transmit the information required under this Section by:

(A) an electronic device compatible with the receiving device of the central repository;

(B) a computer diskette;

(C) a magnetic tape; or

(D) a pharmacy universal claim form or Pharmacy Inventory Control form.

(3.5) The requirements of paragraphs (1), (2), and (3) of this subsection also apply to opioid treatment programs that are licensed or certified by the Department of Human

Services's Division of Substance Use Prevention and Recovery and are authorized by the federal Drug Enforcement Administration to prescribe Schedule II, III, IV, or V controlled substances for the treatment of opioid use disorders. Opioid treatment programs shall attempt to obtain written patient consent, shall document attempts to obtain the written consent, and shall not transmit information without patient consent. Documentation obtained under this paragraph shall not be utilized for law enforcement purposes, as proscribed under 42 CFR 2, as amended by 42 U.S.C. 290dd-2. Treatment of a patient shall not be conditioned upon his or her written consent.

(4) The Department may impose a civil fine of up to $100 per day for willful failure to report controlled substance dispensing to the Prescription Monitoring Program. The fine shall be calculated on no more than the number of days from the time the report was required to be made until the time the problem was resolved, and shall be payable to the Prescription Monitoring Program.

(a-5) Notwithstanding subsection (a), a licensed veterinarian is exempt from the reporting requirements of this Section. If a person who is presenting an animal for treatment is suspected of fraudulently obtaining any controlled substance or prescription for a controlled substance, the licensed veterinarian shall report that information to the local law enforcement agency.

(b) The Department, by rule, may include in the Prescription Monitoring Program certain other select drugs that are not included in Schedule II, III, IV, or V. The Prescription Monitoring Program does not apply to controlled substance prescriptions as exempted under Section 313.

(c) The collection of data on select drugs and scheduled substances by the Prescription Monitoring Program may be used as a tool for addressing oversight requirements of long-term care institutions as set forth by Public Act 96-1372. Long-term care pharmacies shall transmit patient medication profiles to the Prescription Monitoring Program monthly or more frequently as established by administrative rule.

(d) The Department of Human Services shall appoint a full-time Clinical Director of the Prescription Monitoring Program.

(e) (Blank).

(f) Within one year of January 1, 2018 (the effective date of Public Act 100-564), the Department shall adopt rules requiring all Electronic Health Records Systems to interface with the Prescription Monitoring Program application program on or before January 1, 2021 to ensure that all providers have access to specific patient records during the treatment of their patients. These rules shall also address the electronic integration of pharmacy records with the Prescription Monitoring Program to allow for faster transmission of the information required under this Section. The Department shall

establish actions to be taken if a prescriber's Electronic Health Records System does not effectively interface with the Prescription Monitoring Program within the required timeline.

(g) The Department, in consultation with the Prescription Monitoring Program Advisory Committee, shall adopt rules allowing licensed prescribers or pharmacists who have registered to access the Prescription Monitoring Program to authorize a licensed or non-licensed designee employed in that licensed prescriber's office or a licensed designee in a licensed pharmacist's pharmacy who has received training in the federal Health Insurance Portability and Accountability Act and 42 CFR 2 to consult the Prescription Monitoring Program on their behalf. The rules shall include reasonable parameters concerning a practitioner's authority to authorize a designee, and the eligibility of a person to be selected as a designee. In this subsection (g), "pharmacist" shall include a clinical pharmacist employed by and designated by a Medicaid Managed Care Organization providing services under Article V of the Illinois Public Aid Code under a contract with the Department of Healthcare and Family Services for the sole purpose of clinical review of services provided to persons covered by the entity under the contract to determine compliance with subsections (a) and (b) of Section 314.5 of this Act. A managed care entity pharmacist shall notify prescribers of review activities.

(Source: P.A. 100-564, eff. 1-1-18; 100-861, eff. 8-14-18;

100-1005, eff. 8-21-18; 100-1093, eff. 8-26-18; 101-81, eff. 7-12-19; 101-414, eff. 8-16-19.)