



Sen. Antonio Muñoz

**Filed: 5/3/2019**

10100HB0163sam001

LRB101 04752 SLF 60168 a

1 AMENDMENT TO HOUSE BILL 163

2 AMENDMENT NO. \_\_\_\_\_. Amend House Bill 163 by replacing  
3 everything after the enacting clause with the following:

4 "Section 5. The State Finance Act is amended by adding  
5 Section 5.891 as follows:

6 (30 ILCS 105/5.891 new)

7 Sec. 5.891. The Prescription Monitoring Program Fund.

8 Section 10. The Illinois Controlled Substances Act is  
9 amended by changing Section 316 and by adding Section 322 as  
10 follows:

11 (720 ILCS 570/316)

12 Sec. 316. Prescription Monitoring Program.

13 (a) The Department must provide for a Prescription  
14 Monitoring Program for Schedule II, III, IV, and V controlled

1 substances that includes the following components and  
2 requirements:

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

6 (A) The recipient's name and address.

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

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

10 (D) The date the controlled substance is  
11 dispensed.

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

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

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

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

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

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

26 (K) Any additional information that may be

1 required by the department by administrative rule,  
2 including but not limited to information required for  
3 compliance with the criteria for electronic reporting  
4 of the American Society for Automation and Pharmacy or  
5 its successor.

6 (2) Any information transmitted to the Prescription  
7 Monitoring Program as required by the General Assembly  
8 must:

9 (A) be transmitted when a medication is dispensed to  
10 the patient or to the individual receiving the medication  
11 on behalf of the patient;

12 (B) achieve the "point-of-sale" or "real-time"  
13 reporting within 12 months of enactment of this amendatory  
14 Act of the 101st General Assembly;

15 (C) not impose a financial burden upon a pharmacy which  
16 dispenses to patients;

17 (D) use federal grant funding available to the  
18 Prescription Monitoring Program to avoid a financial  
19 burden to any pharmacy which is required to achieve this  
20 requirement;

21 (E) require that any department or agency of this State  
22 applying for or receiving grant funds which relate to  
23 medication, in any manner, must include funding for the  
24 Prescription Monitoring Program's requirement in this  
25 Section unless the Prescription Monitoring Program  
26 declines the need for additional funding; and

1           (F) be transmitted by any pharmacy at the end of the  
2           business day until the Prescription Monitoring Program  
3           verifies that "point-of-sale" or "real-time" reporting is  
4           functional.

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

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

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

14                   (B) (blank) ~~a computer diskette;~~

15                   (C) (Blank) ~~a magnetic tape;~~ or

16                   (D) (blank). ~~a pharmacy universal claim form or~~  
17 ~~Pharmacy Inventory Control form;~~

18           (4) The Department may impose a civil fine of up to  
19 \$100 per day for willful failure to report controlled  
20 substance dispensing to the Prescription Monitoring  
21 Program. The fine shall be calculated on no more than the  
22 number of days from the time the report was required to be  
23 made until the time the problem was resolved. ~~7~~ and The  
24 fineshall be payable to the Prescription Monitoring  
25 Program.

26           (b) The Department, by rule, may include in the

1 Prescription Monitoring Program certain other select drugs  
2 that are not included in Schedule II, III, IV, or V. The  
3 Prescription Monitoring Program does not apply to controlled  
4 substance prescriptions as exempted under Section 313.

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

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

15 (e) (Blank).

16 (f) Within 2 years ~~one year~~ of January 1, 2018 (the  
17 effective date of Public Act 100-564) ~~this amendatory Act of~~  
18 ~~the 100th General Assembly~~, the Department shall adopt rules  
19 requiring all Electronic Health Records Systems to interface  
20 with the Prescription Monitoring Program application program  
21 on or before January 1, 2021 to ensure that all providers have  
22 access to specific patient records during the treatment of  
23 their patients. These rules shall also address the electronic  
24 integration of pharmacy records with the Prescription  
25 Monitoring Program to allow for faster transmission of the  
26 information required under this Section. The Department shall

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

4 (g) The Department, in consultation with the Advisory  
5 Committee, shall adopt rules allowing licensed prescribers or  
6 pharmacists who have registered to access the Prescription  
7 Monitoring Program to authorize a licensed or non-licensed  
8 designee employed in that licensed prescriber's office or a  
9 licensed designee in a licensed pharmacist's pharmacy, ~~and~~ who  
10 has received training in the federal Health Insurance  
11 Portability and Accountability Act to consult the Prescription  
12 Monitoring Program on their behalf. The rules shall include  
13 reasonable parameters concerning a practitioner's authority to  
14 authorize a designee, and the eligibility of a person to be  
15 selected as a designee. In this subsection (g), "pharmacist"  
16 shall include a clinical pharmacist employed by and designated  
17 by a Medicaid Managed Care Organization providing services  
18 under Article V of the Illinois Public Aid Code under a  
19 contract with the Department of Healthcare ~~Health~~ and Family  
20 Services for the sole purpose of clinical review of services  
21 provided to persons covered by the entity under the contract to  
22 determine compliance with subsections (a) and (b) of Section  
23 314.5 of this Act. A managed care entity pharmacist shall  
24 notify prescribers of review activities.

25 (Source: P.A. 99-480, eff. 9-9-15; 100-564, eff. 1-1-18;  
26 100-861, eff. 8-14-18; 100-1005, eff. 8-21-18; 100-1093, eff.

1 8-26-18; revised 2-20-19.)

2 (720 ILCS 570/322 new)

3 Sec. 322. The Prescription Monitoring Program Fund.

4 (a) There is created in the State treasury a special fund  
5 known as the Prescription Monitoring Program Fund. The  
6 Prescription Monitoring Program Fund shall receive revenue  
7 from:

8 (1) grants;

9 (2) pass-through grants;

10 (3) donations;

11 (4) appropriations;

12 (5) fees charged for services, work, or both performed  
13 as the result of duly authorized requests or contracts such  
14 as subpoenas or research agreements;

15 (6) fees as enacted by the General Assembly; and

16 (7) other legal sources.

17 (b) The Department of Human Services or its successor shall  
18 coordinate to use moneys in the Prescription Monitoring Program  
19 Fund and perform the duties of collecting and reporting  
20 prescription and other medication data as authorized by the  
21 General Assembly.

22 (c) Any surplus funds shall be managed as follows:

23 (1) grant and pass-through fund providers shall be  
24 asked for grant extensions or forgiveness. Otherwise, the  
25 funds shall be refunded;

1           (2) donations shall be used to reduce the Prescription  
2           Monitoring Program appropriation for the next fiscal year  
3           unless otherwise restricted;

4           (3) appropriations in excess of spending shall expire  
5           at the end of the fiscal year unless otherwise authorized;  
6           and

7           (4) fees and other legal sources of revenue may be  
8           retained in the Prescription Monitoring Program Fund for 5  
9           fiscal years. The State Treasurer shall invest excess fees  
10           with the interest deposited into the General Revenue Fund  
11           unless otherwise authorized by the General Assembly.

12           (d) If the Prescription Monitoring Program is able to apply  
13           for and receive a reimbursement grant, it is authorized to  
14           apply for the grant. If a reimbursement grant is awarded, the  
15           Prescription Monitoring Program is authorized to borrow  
16           available funds from the Department of Human Services or the  
17           State Treasurer at no interest."