

116TH CONGRESS  
1ST SESSION

# S. 516

To require the use of prescription drug monitoring programs.

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IN THE SENATE OF THE UNITED STATES

FEBRUARY 14, 2019

Ms. KLOBUCHAR (for herself, Mr. PORTMAN, Mr. KING, and Mr. MANCHIN) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

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## A BILL

To require the use of prescription drug monitoring programs.

1       *Be it enacted by the Senate and House of Representa-*  
2       *tives of the United States of America in Congress assembled,*

3       **SECTION 1. SHORT TITLE.**

4       This Act may be cited as the “Prescription Drug  
5       Monitoring Act of 2019”.

6       **SEC. 2. REQUIRING THE USE OF PRESCRIPTION DRUG**  
7               **MONITORING PROGRAMS.**

8       (a) DEFINITIONS.—In this section:

9               (1) CONTROLLED SUBSTANCE.—The term  
10       “controlled substance” has the meaning given the

1 term in section 102 of the Controlled Substances  
2 Act (21 U.S.C. 802).

3 (2) COVERED STATE.—The term “covered  
4 State” means a State that receives funding under  
5 the Harold Rogers Prescription Drug Monitoring  
6 Program established under the Departments of  
7 Commerce, Justice, and State, the Judiciary, and  
8 Related Agencies Appropriations Act, 2002 (Public  
9 Law 107–77; 115 Stat. 748), or under the con-  
10 trolled substance monitoring program under section  
11 3990 of the Public Health Service Act (42 U.S.C.  
12 280g–3).

13 (3) DISPENSER.—The term “dispenser”—

14 (A) means a person licensed or otherwise  
15 authorized by a State to deliver a prescription  
16 drug product to a patient or an agent of the pa-  
17 tient; and

18 (B) does not include a person involved in  
19 oversight or payment for prescription drugs.

20 (4) PDMP.—The term “PDMP” means a pre-  
21 scription drug monitoring program.

22 (5) PRACTITIONER.—The term “practitioner”  
23 means a practitioner registered under section 303(f)  
24 of the Controlled Substances Act (21 U.S.C. 823(f))

1 to prescribe, administer, or dispense controlled sub-  
2 stances.

3 (6) STATE.—The term “State” means each of  
4 the several States and the District of Columbia.

5 (b) REQUIREMENTS.—Beginning 1 year after the  
6 date of enactment of this Act, each covered State shall  
7 require—

8 (1) each prescribing practitioner within the cov-  
9 ered State or their designee, who shall be licensed or  
10 registered healthcare professionals or other employ-  
11 ees who report directly to the practitioner, to consult  
12 the PDMP of the covered State before initiating  
13 treatment with a prescription for a controlled sub-  
14 stance listed in schedule II, III, or IV of section  
15 202(c) of the Controlled Substances Act (21 U.S.C.  
16 812(c)), and every 3 months thereafter as long as  
17 the treatment continues;

18 (2) the PDMP of the covered State to provide  
19 proactive notification to a practitioner when patterns  
20 indicative of controlled substance misuse, including  
21 opioid misuse, are detected;

22 (3) each dispenser within the covered State to  
23 report each prescription for a controlled substance  
24 dispensed by the dispenser to the PDMP not later

1 than 24 hours after the controlled substance is dis-  
2 pensed to the patient;

3 (4) that the PDMP make available a quarterly  
4 de-identified data set and an annual report for pub-  
5 lic and private use, including use by healthcare pro-  
6 viders, health plans and health benefits administra-  
7 tors, State agencies, and researchers, which shall, at  
8 a minimum, meet requirements established by the  
9 Attorney General, in coordination with the Secretary  
10 of Health and Human Services;

11 (5) each State agency that administers the  
12 PDMP to—

13 (A) proactively analyze data available  
14 through the PDMP; and

15 (B) provide reports to law enforcement  
16 agencies and prescriber licensing boards de-  
17 scribing any prescribing practitioner that re-  
18 peatedly fall outside of expected norms or  
19 standard practices for the prescribing practi-  
20 tioner's field; and

21 (6) that the data contained in the PDMP of the  
22 covered State be made available to other States.

23 (c) NONCOMPLIANCE.—If a covered State fails to  
24 comply with subsection (a), the Attorney General or the  
25 Secretary of Health and Human Services may withhold

1 grant funds from being awarded to the covered State  
2 under the Harold Rogers Prescription Drug Monitoring  
3 Program established under the Departments of Com-  
4 merce, Justice, and State, the Judiciary, and Related  
5 Agencies Appropriations Act, 2002 (Public Law 107-77;  
6 115 Stat. 748), or under the controlled substance moni-  
7 toring program under section 3990 of the Public Health  
8 Service Act (42 U.S.C. 280g-3).

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