

117TH CONGRESS  
1ST SESSION

# H. R. 2344

To require the use of prescription drug monitoring programs.

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IN THE HOUSE OF REPRESENTATIVES

APRIL 1, 2021

Mr. RYAN (for himself and Mr. BALDERSON) introduced the following bill;  
which was referred to the Committee on Energy and Commerce

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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 2021”.

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

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

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

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

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

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

20           (5) PRACTITIONER.—The term “practitioner”  
21 means a practitioner registered under section 303(f)  
22 of the Controlled Substances Act (21 U.S.C. 823(f))  
23 to prescribe, administer, or dispense controlled sub-  
24 stances.

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

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

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

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

20          (3) each dispenser within the covered State to  
21          report each prescription for a controlled substance  
22          dispensed by the dispenser to the PDMP not later  
23          than 24 hours after the controlled substance is dis-  
24          pensed to the patient;

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

9           (5) each State agency that administers the  
10 PDMP to—

11           (A) proactively analyze data available  
12 through the PDMP; and

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

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

21           (c) NONCOMPLIANCE.—If a covered State fails to  
22 comply with subsection (a), the Attorney General or the  
23 Secretary of Health and Human Services may withhold  
24 grant funds from being awarded to the covered State  
25 under the Harold Rogers Prescription Drug Monitoring

1 Program established under the Departments of Com-  
2 merce, Justice, and State, the Judiciary, and Related  
3 Agencies Appropriations Act, 2002 (Public Law 107-77;  
4 115 Stat. 748), or under the controlled substance moni-  
5 toring program under section 3990 of the Public Health  
6 Service Act (42 U.S.C. 280g-3).

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