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4lr1353 CF HB 255

By: **The President (By Request – Department of Legislative Services)** Introduced and read first time: January 17, 2014 Assigned to: Finance

A BILL ENTITLED

1 AN ACT concerning

Prescription Drug Monitoring Program – Sunset Extension and Program Evaluation

4 FOR the purpose of continuing the Prescription Drug Monitoring Program in $\mathbf{5}$ accordance with the provisions of the Maryland Program Evaluation Act 6 (Sunset Law) by extending to a certain date the termination provisions relating 7 to the statutory and regulatory authority of the Program; requiring the 8 Department of Legislative Services to conduct a certain evaluation of the 9 Program on or before a certain date and to prepare and submit a certain report in accordance with certain statutory requirements; requiring the Program to 10 submit a certain report to the Governor, the General Assembly, and the 11 12Department of Legislative Services on or before a certain date; repealing the 13 requirement that the technical advisory committee to the Program review requests for certain information before the Program discloses the information to 14a certain person; requiring the Advisory Board on Prescription Drug Monitoring 1516 to include certain information in a certain report; repealing an obsolete 17reporting requirement; and generally relating to the Prescription Drug 18 Monitoring Program.

- 19 BY repealing and reenacting, without amendments,
- 20 Article Health General
- 21 Section 21–2A–05(a), 21–2A–06(b), and 21–2A–07(a)
- 22 Annotated Code of Maryland
- 23 (2009 Replacement Volume and 2013 Supplement)
- 24 BY repealing and reenacting, with amendments,
- 25 Article Health General
- 26 Section 21–2A–05(f)(3), 21–2A–06(c), 21–2A–07(b), and 21–2A–10
- 27 Annotated Code of Maryland
- 28 (2009 Replacement Volume and 2013 Supplement)

EXPLANATION: CAPITALS INDICATE MATTER ADDED TO EXISTING LAW. [Brackets] indicate matter deleted from existing law.



$\frac{1}{2}$	SECTION 1. BE IT ENACTED BY THE GENERAL ASSEMBLY OF MARYLAND, That the Laws of Maryland read as follows:
3	Article – Health – General
4	21–2A–05.
$5 \\ 6$	(a) There is an Advisory Board on Prescription Drug Monitoring in the Department.
7	(f) The Board shall:
	(3) [(i) Provide within 180 days after its first meeting, in accordance with § 2–1246 of the State Government Article, an interim report to the General Assembly setting forth the Board's analysis and recommendations under item (2) of this subsection relating to the design, implementation, and funding of the Program; and
$13 \\ 14 \\ 15$	(ii)] Provide annually to the Governor and, in accordance with § 2–1246 of the State Government Article, the General Assembly [an analysis] A REPORT THAT INCLUDES:
$\begin{array}{c} 16 \\ 17 \end{array}$	(I) THE NUMBER OF PRESCRIBERS REGISTERED WITH AND USING THE PROGRAM;
18 19	(II) THE NUMBER OF DISPENSERS REGISTERED WITH AND USING THE PROGRAM;
$20 \\ 21 \\ 22$	(III) AN ANALYSIS of the impact of the Program on patient access to pharmaceutical care and on curbing prescription drug diversion in the State[, including any]; AND
$\begin{array}{c} 23\\ 24 \end{array}$	(IV) ANY recommendations related to modification or continuation of the Program; and
25	21–2A–06.
$\begin{array}{c} 26\\ 27 \end{array}$	(b) The Program shall disclose prescription monitoring data, in accordance with regulations adopted by the Secretary, to:
28 29	(1) A prescriber, or a licensed health care practitioner authorized by the prescriber, in connection with the medical care of a patient;
30 31	(2) A dispenser, or a licensed health care practitioner authorized by the dispenser, in connection with the dispensing of a monitored prescription drug;

$egin{array}{c} 1 \\ 2 \\ 3 \end{array}$	(3) A federal law enforcement agency or a State or local law enforcement agency, on issuance of a subpoena, for the purpose of furthering an existing bona fide individual investigation;
$\begin{array}{c} 4\\ 5\\ 6\end{array}$	(4) A licensing entity, on issuance of an administrative subpoena voted on by a quorum of the board of the licensing entity, for the purposes of furthering an existing bona fide individual investigation;
$7 \\ 8$	(5) A rehabilitation program under a health occupations board, on issuance of an administrative subpoena;
9 10	(6) A patient with respect to prescription monitoring data about the patient;
$\begin{array}{c} 11 \\ 12 \end{array}$	(7) Subject to subsection (g) of this section, the authorized administrator of another state's prescription drug monitoring program;
$\begin{array}{c} 13\\14\end{array}$	(8) The following units of the Department, on approval of the Secretary, for the purpose of furthering an existing bona fide individual investigation:
15	(i) The Office of the Chief Medical Examiner;
16	(ii) The Maryland Medical Assistance Program;
17	(iii) The Office of the Inspector General;
18	(iv) The Office of Health Care Quality; and
19	(v) The Division of Drug Control; or
$\begin{array}{c} 20\\ 21 \end{array}$	(9) The technical advisory committee established under § $21-2A-07$ of this subtitle for the purposes set forth in subsection (c) of this section.
$\frac{22}{23}$	(c) Before the Program discloses information under subsection (b)(3), (4), (5),[(7),] or (8) of this section, the technical advisory committee to the Program shall:
24	(1) Review the requests for information;
$25 \\ 26 \\ 27$	(2) Provide clinical guidance and interpretation of the information requested to the Secretary to assist in the Secretary's decision on how to respond to a judicial subpoena, administrative subpoena, or other request; and
28 29	(3) Provide clinical guidance and interpretation of the information requested to the authorized recipient of the information.

30 21–2A–07.

1 (a) There is a technical advisory committee to the Program.

2 (b) The purpose of the technical advisory committee is to review requests for 3 information from the Program under § 21–2A–06(b)(3), (4), (5), [(7),] and (8) of this 4 subtitle.

5 21–2A–10.

6 Subject to the evaluation and reestablishment provisions of the Maryland 7 Program Evaluation Act, this subtitle and all regulations adopted under this subtitle 8 shall terminate and be of no effect after July 1, [2016] **2019**.

9 SECTION 2. AND BE IT FURTHER ENACTED, That, on or before January 1, 10 2015, the Prescription Drug Monitoring Program shall submit a report to the Governor 11 and, in accordance with § 2–1246 of the State Government Article, the General 12 Assembly, and the Department of Legislative Services that:

(1) describes efforts to collect and make available, in real-time,
prescription monitoring data;

15 (2) includes recommendations for a long-term funding source to 16 support the Program;

17 (3) provides the status of the Department of Health and Mental18 Hygiene's independent evaluation of the Program; and

(4) discusses the status of any plans to pursue unsolicited reporting or
mandatory utilization of prescription monitoring data by health care providers.

21 SECTION 3. AND BE IT FURTHER ENACTED, That the Department of 22 Legislative Services shall:

(1) conduct a direct full evaluation of the Prescription Drug Monitoring
Program on or before December 1, 2017; and

25 (2) prepare and submit a full evaluation report in accordance with the 26 requirements established under § 8–405(e) and (f) of the State Government Article.

SECTION 4. AND BE IT FURTHER ENACTED, That this Act shall take effect
July 1, 2014.

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