THE GENERAL ASSEMBLY OF PENNSYLVANIA

HOUSE BILL No. 1005 Session of 2019

INTRODUCED BY COX, BERNSTINE, BROWN, IRVIN, KIRKLAND, MASSER, MILLARD, MURT, NEILSON, PICKETT, PYLE, READSHAW, SAYLOR, STRUZZI, ZIMMERMAN AND HEFFLEY, APRIL 9, 2019

AS AMENDED ON THIRD CONSIDERATION, HOUSE OF REPRESENTATIVES, JUNE 24, 2019

AN ACT

1 2 3 4 5 6	Amending the act of October 27, 2014 (P.L.2911, No.191), entitled "An act providing for prescription drug monitoring; creating the ABC-MAP Board; establishing the Achieving Better Care by Monitoring All Prescriptions Program; and providing for unlawful acts and penalties," further providing for purpose, for definitions, for powers and duties of board and	
7	for establishment of program; and providing for requirements	<
8	for first responder agencies and hospital emergency	
9 10	departments and for requirements for coroners and medical examiners ; AND FURTHER PROVIDING FOR ACCESS TO PRESCRIPTION	< <
11	INFORMATION.	_
12	The General Assembly of the Commonwealth of Pennsylvania	
13	hereby enacts as follows:	
14	Section 1. Section 2 of the act of October 27, 2014	
15	(P.L.2911, No.191), known as the Achieving Better Care by	
16	Monitoring All Prescriptions Program (ABC-MAP) Act, is amended	
17	to read:	
18	Section 2. Purpose.	
19	This act is intended to increase the quality of patient care	
20	by giving prescribers and dispensers access to a patient's	
21	prescription medication history, including, but not limited to,	

any history of a drug-related overdose event, through an 1 2 electronic system that will alert medical professionals to 3 potential dangers for purposes of making treatment determinations. The act further intends that patients will have 4 a thorough and easily obtainable record of their prescriptions 5 for purposes of making educated and thoughtful health care 6 7 decisions. Additionally, the act seeks to aid regulatory and law 8 enforcement agencies in the detection and prevention of fraud, drug abuse and the criminal diversion of controlled substances. 9 10 Section 2. Section 3 of the act is amended by adding definitions to read: 11 12 Section 3. Definitions. 13 The following words and phrases when used in this act shall have the meanings given to them in this section unless the 14 15 context clearly indicates otherwise: 16 * * "Drug related overdose death." An incident where an over-17 <---18 the counter drug, prescription or controlled substance or 19 illegal substance is the primary or secondary cause of death of 20 an individual or may have been a contributing factor to the 21 death of an individual. 22 "Drug-related overdose event." As follows: 23 (1) An incidence of a physical state resulting from 24 intentionally or unintentionally consuming or administering a 25 toxic or otherwise harmful level of an over-the-counter drug, 26 prescription or controlled substance or illegal substance that may be suspected by any of the following: 27 28 (i) An observation of symptoms requiring medic 29 response. 30 (ii) A clinical suspicion of a drug overdose.

20190HB1005PN2238

- 2 -

1	(iii) A positive urine toxicology screen for a
2	controlled substance or a negative urine toxicology
3	screen if there are no other conditions to explain the
4	clinical symptoms.
5	(2) The term may include, but is not limited to, any of
6	the following events that resulted from consuming drugs:
7	(i) Central nervous system depression resulting in a
8	decreased heart rate and breathing, loss of consciousness
9	<u>or death.</u>
10	(ii) Stimulant effects resulting in an increased or
11	irregular heart rate, agitation or hypertension.
12	(iii) Hallucinations, seizures or unresponsiveness.
13	"DRUG-RELATED OVERDOSE EVENT." AN INCIDENCE OF A PHYSICAL <
14	STATE RESULTING FROM INTENTIONALLY OR UNINTENTIONALLY CONSUMING
15	OR ADMINISTERING A TOXIC OR OTHERWISE HARMFUL LEVEL OF
16	CONTROLLED PRESCRIPTION MEDICATION OR ILLEGAL SUBSTANCE THAT MAY
17	BE SUSPECTED BY ANY OF THE FOLLOWING:
18	(1) AN OBSERVATION OF SYMPTOMS REQUIRING AN EMERGENT
19	MEDICAL RESPONSE.
20	(2) A CLINICAL SUSPICION OF A DRUG OVERDOSE.
21	"First responder." A firefighter, law enforcement officer or
22	emergency medical services provider.
23	"First responder agency." A Federal, State, local
24	governmental or nongovernmental agency that employs first
25	responders. The term includes an emergency medical services
26	agency as defined in 35 Pa.C.S. § 8103 (relating to
27	<u>definitions).</u>
28	* * *
29	"SINGLE COUNTY AUTHORITY." THE AGENCY DESIGNATED TO PLAN AND <
30	COORDINATE DRUG AND ALCOHOL PREVENTION, INTERVENTION AND

20190HB1005PN2238

- 3 -

TREATMENT SERVICES FOR A GEOGRAPHIC AREA, WHICH MAY CONSIST OF 1 ONE OR MORE COUNTIES, AND TO ADMINISTER THE PROVISIONS OF SUCH 2 SERVICES FUNDED THROUGH THE AGENCY. 3 * * * 4 5 Section 3. Section 5 of the act is amended to read: <---SECTION 3. SECTION 5(5)(V) OF THE ACT IS AMENDED AND THE 6 <---7 PARAGRAPH IS AMENDED BY ADDING SUBPARAGRAPHS TO READ: Section 5. Powers and duties of board. 8 9 The board shall have the following powers and duties: (1) Evaluate and secure a vendor of an electronic 10 <---11 prescription monitoring system for the purpose of carrying out the provisions of this act. 12 13 (2) Appoint an advisory group comprised of dispensers, 14 prescribers, law enforcement officials, addiction 15 specialists, patient and privacy advocates and individuals 16 with expertise considered important to the operation of the program. All members shall have varying perspectives and will-17 18 provide input and recommendations to the board regarding the 19 establishment and maintenance of the program. The advisory 20 group shall not exceed 12 members. 21 (3) Create a written notice to be used by prescribers 2.2 and used or displayed by dispensers to provide notice to-23 patients that information regarding prescriptions for-24 controlled substances and drug-related overdose events is 25 being collected by the program and that the patient has a 26 right to review and correct the information with the program. The notice must include all of the following: 27 28 (i) The manner in which the patient may access the patient's personal information. The notice shall state-29 30 that one time quarterly patient access shall be at no-20190HB1005PN2238

- 4 -

1 cost.

2	(ii) An explanation of the program and the program's
3	authorized users.
4	(iii) The program's record retention policies.
5	(iv) An explanation that prescription information is
6	confidential and is not subject to the act of February
7	14, 2008 (P.L.6, No.3), known as the Right to Know Law.
8	(v) Any cost associated with accessing the
9	information more than once during each calendar quarter.
10	(4) Phase in an enforcement process so that dispensers
11	and prescribers may transition and have adequate time to make-
12	the necessary changes to their operating systems.
13	* * * * <
14	(5) Develop policies and procedures to:
15	(i) Require more frequent reporting of prescription <
16	medication information under section 7 should technology-
17	permit and so long as there is little or no fiscal impact-
18	to the Commonwealth or those required to report. Any-
19	change in the frequency of reporting shall be made in-
20	collaboration with the Board of Pharmacy and the Board of
21	Pharmacy's members to ensure that a pharmacy is able to
22	accommodate the change.
23	(ii) Evaluate the information in the system.
24	(iii) Allow for authorized department personnel to
25	conduct internal reviews, analyses and interpret the data
26	contained in the system.
27	(iv) Safeguard the release of information to-
28	authorized users and department personnel and ensure the-
29	privacy and confidentiality of patients and patient
30	information.

20190HB1005PN2238

- 5 -

1	* * *	<
2	(v) Aid prescribers <u>AND FIRST RESPONDERS</u> in	<
3	identifying at-risk individuals and referring them to	
4	SINGLE COUNTY AUTHORITIES, drug addiction treatment	<
5	professionals and programs.	
6	(v.1) Aid prescribers AND FIRST RESPONDERS in	<
7	identifying individuals with a history of drug overdoses	-
8	in order to provide alternative treatment options.	
9	(vi) Establish professionally developed criteria,	<
10	with the advice of the advisory group, that generates	
11	referrals of prescription monitoring information to the	

appropriate licensing board in the Department of State. A 13 referral may only be generated when the system produces 14 an alert that there is a pattern of irregular data for a-15 dispenser or prescriber which appears to deviate from the 16 clinical standard.

(vii) Provide training to prescribers and dispensers 17 18 on the use of the system.

19 (viii) Assist professional organizations whose 20 members prescribe, monitor or treat patients or dispense-21 controlled substances to patients to develop educational 22 programs for those members relating to prescribing-23 practices, pharmacology, controlled substance abuse and 24 clinical standards, including:

25 (A) identification of those at risk for 26 controlled substance abuse; and 27 (B) referral and treatment options for patients. 28 (ix) Permit individuals employed by prescribers, 29 pharmacies and dispensers to query the system as

designees so long as each individual designee has a 30

12

- 6 -

1	unique identifier when accessing the system and set	
2	explicit standards to qualify individuals authorized to-	
3	query the system and to ensure the security of the system-	
4	when used by a designee.	
5	(x) Keep pace with technological advances that	
6	facilitate the interoperability of the system with other-	
7	states' prescription drug monitoring systems and	
8	electronic health information systems.	
9	(xi) Evaluate the costs and benefits of the program.	
10	(xii) Convene the advisory group at least annually.	
11	(xiii) Direct the department to operate and maintain	
12	the program on a daily basis.	
13	(xiv) Review the program for the purpose of	
14	compiling statistics, research and educational materials	
15	and outreach.	
16	(xv) Identify any controlled substance that has been	
17	shown to have limited or no potential for abuse and	
18	therefore should not be reported to the program.	
19	(xvi) Require and ensure registration of all	
20	prescribers and dispensers with the program.	
21	* * *	
22	(xvii) Identify additional medications that could	
23	assist prescribers in making treatment options for	
24	patients who are at risk for a substance use disorder.	
25	Section 4. Section 6(b)(1) of the act is amended and the	
26	<pre>section SUBSECTION is amended by adding paragraphs to read:</pre>	
27	Section 6. Establishment of program.	
28	* * *	
29	(b) Program componentsThe program shall:	
30	(1) Provide an electronic system of controlled	
201	90HB1005PN2238 - 7 -	

1	substances prescribed and dispensed in this Commonwealth <u>and</u>
2	of drug-related overdose events that occurred in this
3	Commonwealth.
4	* * *
5	(6) Establish a protocol for hospital emergency
6	departments and first responder agencies to ensure data
7	submitted to the system with respect to drug-related overdose_
8	<u>events is not duplicative.</u>
9	(7) Provide drug-related overdose death event
10	information, including any drugs that contributed to the
11	overdose, on the patient's program record.
12	* * *
13	Section 5. The act is amended by adding $\frac{1}{2}$ A SECTION <
14	to read:
15	Section 7.1. Requirements for first responder agencies and
16	hospital emergency departments.
17	(a) SubmissionA first responder agency or hospital
18	emergency department shall, in the format determined by the
19	department, electronically submit drug-related overdose event
20	information to the department.
21	(b) Data elementsAll of the following information THAT IS <
22	AVAILABLE AND REASONABLY ABLE TO BE IDENTIFIED DURING A REVIEW
23	OF THE INDIVIDUAL'S MEDICAL RECORDS shall be provided by a first
24	responder agency or hospital emergency department:
25	(1) The full name, date of birth, gender and address of
26	an THE individual who experienced a drug-related overdose <
27	event.
28	(2) The date and time of the drug-related overdose
29	event.
30	(3) The address where the individual was picked up or
201	90HB1005PN2238 - 8 -

1	where the drug-related overdose event took place.
2	(4) Whether an emergency opioid antagonist was
3	administered to the individual.
4	(5) The location where the emergency opioid antagonist
5	was administered, if available.
6	(6) The amount of emergency opioid antagonist
7	administered, if available.
8	(7) Whether the drug-related overdose event resulted in
9	death.
10	(8) The suspected or confirmed drug involved in the
11	drug-related overdose event.
12	(c) FrequencyA first responder agency or hospital
13	emergency department shall submit all information required under
14	subsection (b) to the program no later than 72 hours after a <
15	drug-related overdose event was reported 14 DAYS AFTER THE <
16	COMPLETION OF THE ACUTE EPISODE OF CARE.
17	(d) DefinitionAs used in this section, the term
18	"emergency opioid antagonist" means a medication approved by the
19	United States Food and Drug Administration to reverse the
20	effects of an opioid drug.
21	Section 7.2. Requirements for coroners and medical examiners. <
22	(a) Submission. A county coroner or medical examiner in
23	this Commonwealth shall electronically submit data, in the
24	format published under subsection (c), on a drug related
25	overdose death to the department within five business days of
26	finalizing the cause and manner of the drug related overdose
27	death.
28	(b) Contents of data. In complying with subsection (a), a
29	county coroner or medical examiner shall provide all of the
30	following information to the department:

20190HB1005PN2238

- 9 -

1	(1) Demographic information of the decedent, including
2	but not limited to, the full name, address and date of birth
3	of the decedent.
4	(2) The toxicology report.
5	(3) The autopsy report.
6	(4) The circumstances of the drug related overdose
7	death.
8	(c) Publication. The department shall transmit a notice of
9	the format for data submission under subsection (a) to the
10	Legislative Reference Bureau for publication in the Pennsylvania
11	Bulletin within 30 days of the effective date of this
12	subsection.
13	(d) Public reports. The department shall use the data
14	submitted under subsection (a) to compile publicly available
15	reports containing statistics and patterns relating to drug-
16	related overdose deaths on a quarterly basis to help identify
17	threats to public health and safety.
18	(e) Liability Any individual who, in good faith, provides
19	data to the department under this section shall not be subject
20	to any civil or criminal liability as a result of providing the
21	data.
22	Section 6. Section 9 heading of the act is amended to read:
23	SECTION 6. SECTION 9 HEADING AND (B)(3) OF THE ACT ARE <
24	AMENDED AND SUBSECTION (B) IS AMENDED BY ADDING A PARAGRAPH TO
25	READ:
26	Section 9. Access to prescription information and drug-related
27	overdose event information.
28	* * *
29	(B) AUTHORIZED USERSTHE FOLLOWING INDIVIDUALS MAY QUERY <
30	THE SYSTEM ACCORDING TO PROCEDURES DETERMINED BY THE BOARD AND

20190HB1005PN2238

- 10 -

2 * * *

3 (3) (1) THE OFFICE OF ATTORNEY GENERAL SHALL QUERY THE
4 SYSTEM ON BEHALF OF ALL LAW ENFORCEMENT AGENCIES,
5 INCLUDING, BUT NOT LIMITED TO, THE OFFICE OF THE ATTORNEY
6 GENERAL AND FEDERAL, STATE AND LOCAL LAW ENFORCEMENT
7 AGENCIES FOR:

8 (A) SCHEDULE II CONTROLLED SUBSTANCES AS 9 INDICATED IN THE ACT OF APRIL 14, 1972 (P.L.233, 10 NO.64), KNOWN AS THE CONTROLLED SUBSTANCE, DRUG, 11 DEVICE AND COSMETIC ACT, AND IN THE MANNER DETERMINED 12 BY THE PENNSYLVANIA ATTORNEY GENERAL PURSUANT TO 28 13 PA. CODE § 25.131 (RELATING TO EVERY DISPENSING 14 PRACTITIONER);

(B) ALL OTHER SCHEDULES UPON RECEIPT OF A COURT
ORDER OBTAINED BY THE REQUESTING LAW ENFORCEMENT
AGENCY. UPON RECEIPT OF A MOTION UNDER THIS CLAUSE,
THE COURT MAY ENTER AN EX PARTE ORDER GRANTING THE
MOTION IF THE LAW ENFORCEMENT AGENCY HAS DEMONSTRATED
BY A PREPONDERANCE OF THE EVIDENCE THAT:

(I) THE MOTION PERTAINS TO A PERSON WHO IS
THE SUBJECT OF AN ACTIVE CRIMINAL INVESTIGATION
WITH A REASONABLE LIKELIHOOD OF SECURING AN
ARREST OR PROSECUTION IN THE FORESEEABLE FUTURE;
AND

26 (II) THERE IS REASONABLE SUSPICION THAT A27 CRIMINAL ACT HAS OCCURRED.

(II) DATA OBTAINED BY A LAW ENFORCEMENT AGENCY UNDER
THIS PARAGRAPH SHALL ONLY BE USED TO ESTABLISH PROBABLE
CAUSE TO OBTAIN A SEARCH WARRANT OR ARREST WARRANT.

20190HB1005PN2238

- 11 -

1 (III) REQUESTS MADE TO THE OFFICE OF ATTORNEY 2 GENERAL TO OUERY THE SYSTEM UNDER THIS PARAGRAPH SHALL BE 3 MADE IN A FORM OR MANNER PRESCRIBED BY THE OFFICE OF ATTORNEY GENERAL AND SHALL INCLUDE THE COURT ORDER, WHEN 4 5 APPLICABLE. EACH INDIVIDUAL DESIGNEE OF THE OFFICE OF 6 ATTORNEY GENERAL SHALL HAVE A UNIQUE IDENTIFIER WHEN 7 ACCESSING THE SYSTEM. 8 (IV) THE OFFICE OF ATTORNEY GENERAL SHALL NOT OUERY 9 THE SYSTEM FOR INFORMATION REGARDING A DRUG-RELATED 10 OVERDOSE EVENT. * * * 11 12 (12) AN AUTHORIZED EMPLOYEE OF A COUNTY OR MUNICIPAL 13 HEALTH DEPARTMENT MAY OUERY THE SYSTEM IF THE EMPLOYEE HAS A 14 UNIQUE IDENTIFIER WHEN ACCESSING THE SYSTEM AND THE EMPLOYEE ACCESSES THE SYSTEM FOR ANY OF THE FOLLOWING PURPOSES: 15 16 (I) DEVELOPING EDUCATIONAL PROGRAMS RELATING TO PRESCRIBING PRACTICES AND CONTROLLED SUBSTANCE ABUSE. 17 18 (II) IDENTIFYING AT-RISK INDIVIDUALS FOR THE PURPOSE 19 OF CONNECTING THEM WITH ADDICTION TREATMENT PROFESSIONALS 20 AND PROGRAMS, INCLUDING SINGLE COUNTY AUTHORITIES. 21 (III) COMPILING EPIDEMIOLOGICAL DATA TO ENSURE THE 22 SECURITY OF THE SYSTEM WHEN AN AUTHORIZED EMPLOYEE OF A 23 COUNTY OR MUNICIPAL HEALTH DEPARTMENT ACCESSES THE 24 SYSTEM. 25 * * * 26 Section 7. This act shall take effect in 180 days.

20190HB1005PN2238

- 12 -