



Types of Authorized Recipients – Department of Health or Commissioner of Public Safety

Research current through May 2016.

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Introduction

Each state determines by statute or regulation the persons or entities entitled to access or receive information in the prescription monitoring program database in that particular state. This memorandum sets out those states that allow access to or receipt of database information by the state Department of Health or Commissioner of Public Safety. This does not mean that if a particular state is not listed in this memorandum or the accompanying map that those entities in that state are not allowed access to the information. The following states either specifically include the Department of Health or Commissioner of Public Safety in the list of persons or entities entitled to access or NAMSDDL was informed by the administrator of the state prescription monitoring program that such persons are allowed access.

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New Mexico
ADC 16.19.29

Code of New Mexico Rules (2016)
Title 16. Occupational and Professional Licensing
Chapter 19. Pharmacists
Part 29. Controlled Substance Prescription Monitoring Program

16.19.29. CONTROLLED SUBSTANCE PRESCRIPTION MONITORING PROGRAM

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16.19.29.9 DISCLOSURE OF PRESCRIPTION INFORMATION:

A. Prescription information submitted to the board shall not be subject to Sections 14-2-1 through 14-2-12 of the Inspection of Public Records Act, NMSA 1978, and shall be confidential except as provided in Subsections C through G of 16.19.29.9 NMAC.

B. The board shall maintain procedures to ensure that the privacy and confidentiality of patients and patient information collected, recorded, transmitted, and maintained is not disclosed to persons except as provided in Subsection C through G of 16.19.29.9 NMAC.

C. After receiving a complaint, the board inspectors shall review the relevant prescription information. If there is reasonable cause to believe a violation of law or breach of professional standards may have occurred, the board shall notify the appropriate law enforcement or professional licensing, certification or regulatory agency or entity, and provide prescription information required for an investigation.

D. The board shall be authorized to provide PMP information to the following persons:

(1) persons authorized to prescribe or dispense controlled substances, for the purpose of providing medical or pharmaceutical care for their patients;

(2) a delegate designated by a practitioner; a practitioner, who must also maintain an active account, can designate only one delegate for the purpose of requesting and receiving PMP reports for that practitioner;

(3) state licensing boards, including the medical board, board of nursing, board of veterinary medicine, board of dental health care, board of examiners in optometry, osteopathic examiners board, acupuncture & oriental medicine board, and podiatry board, as the PMP information relates to their licensees;

- (4) professional licensing authorities of other states if their licensees practice in this state or prescriptions provided by their licensees are dispensed in this state;
- (5) local, state and federal law enforcement or prosecutorial officials engaged in an ongoing investigation of an individual in the enforcement of the laws governing licit drugs;
- (6) the state human services department regarding medicaid program recipients;
- (7) a state metropolitan, magistrate and district, or federal court as required by a grand jury subpoena or criminal court order;
- (8) state drug court personnel as authorized by the PMP director;
- (9) personnel of the board for purposes of administration and enforcement of this rule or of 16.19.20 NMAC;
- (10) the prescription monitoring program of another state or group of states with whom the state has established an interoperability agreement;
- (11) a living individual who request's his or her own PMP report in accordance with procedures established under Subsection D of Section 61-11-2 of the Pharmacy Act, NMSA 1978 and Subsection H of 16.19.6.23 NMAC, or an agent authorized by the living individual along with a valid HIPAA release form or court issued subpoena, or;
- (12) a parent to have access to the prescription records about his or her minor child, as his or her minor child's personal representative when such access is not inconsistent with state or other laws;

E. The board shall use de-identified data obtained from the PMP database to identify and report to state and local public health authorities the geographic areas of the state where anomalous prescribing dispensing or use of controlled substances is occurring.

F. The board shall share PMP database data with the department of health for the purpose of tracking inappropriate prescribing and misuse of controlled substances, including drug overdose.

G. The board shall provide data to public or private entities for statistical, research, or educational purposes after removing information that could be used to identify individual patients and persons who have received prescriptions from dispensers.

H. PMP information gained from other states' prescription monitoring programs shall not be subject to civil subpoena, nor shall such information be disclosed, discoverable, or compelled to be produced in any civil proceeding, nor shall such records be deemed admissible as evidence in any civil proceeding for any reason.

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New York
Public Health Law § 3371

McKinney's Consolidated Laws of New York Annotated (2016)
Public Health Law
Chapter 45. Of the Consolidated Laws
Article 33. Controlled Substances
Title VI. Records and Reports

§ 3371. Confidentiality of certain records, reports, and information

1. No person, who has knowledge by virtue of his or her office of the identity of a particular patient or research subject, a manufacturing process, a trade secret or a formula shall disclose such knowledge, or any report or record thereof, except:

- (a) to another person employed by the department, for purposes of executing provisions of this article;
- (b) pursuant to judicial subpoena or court order in a criminal investigation or proceeding;
- (c) to an agency, department of government, or official board authorized to regulate, license or otherwise supervise a person who is authorized by this article to deal in controlled substances, or in the course of any investigation or proceeding by or before such agency, department or board;
- (d) to the prescription monitoring program registry and to authorized users of such registry as set forth in subdivision two of this section;
- (e) to a practitioner to inform him or her that a patient may be under treatment with a controlled substance by another practitioner for the purposes of subdivision two of this section, and to facilitate the department's review of individual challenges to the accuracy of controlled substances histories pursuant to subdivision six of section thirty-three hundred forty-three-a of this article;
- (f) to a pharmacist to provide information regarding prescriptions for controlled substances presented to the pharmacist for the purposes of subdivision two of this section and to facilitate the department's review of individual challenges to the accuracy of controlled substances histories pursuant to subdivision six of section thirty-three hundred forty-three-a of this article;
- (g) to the deputy attorney general for medicaid fraud control, or his or her designee, in furtherance of an investigation of fraud, waste or abuse of the Medicaid program, pursuant to an agreement with the department;

(h) to a local health department for the purpose of conducting public health research or education: (i) pursuant to an agreement with the commissioner; (ii) when the release of such information is deemed appropriate by the commissioner; (iii) for use in accordance with measures required by the commissioner to ensure that the security and confidentiality of the data is protected; and (iv) provided that disclosure is restricted to individuals within the local health department who are engaged in the research or education;

(i) to a medical examiner or coroner who is an officer of or employed by a state or local government, pursuant to his or her official duties; and

(j) to an individual for the purpose of providing such individual with his or her own controlled substance history or, in appropriate circumstances, in the case of a patient who lacks capacity to make health care decisions, a person who has legal authority to make such decisions for the patient and who would have legal access to the patient's health care records, if requested from the department pursuant to subdivision six of section thirty-three hundred forty-three-a of this article or from a treating practitioner pursuant to subparagraph (iv) of paragraph (a) of subdivision two of this section.

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Ohio
§ 4729.80

Baldwin's Ohio Revised Code Annotated (2016)
Title XLVII. Occupations--Professions
Chapter 4729. Pharmacists; Dangerous Drugs
Miscellaneous Provisions

§ 4729.80 Disclosure of database information; disclosure of requests for database information

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(A) If the state board of pharmacy establishes and maintains a drug database pursuant to section 4729.75 of the Revised Code, the board is authorized or required to provide information from the database in accordance with the following:

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(13) On receipt of a request from the director of health, the board shall provide to the director information from the database relating to the duties of the director or the department of health in implementing the Ohio violent death reporting system established under section 3701.93 of the Revised Code.

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Oklahoma
63 § 2-309D

Oklahoma Statutes Annotated (2016)

Title 63. Public Health and Safety

Chapter 2. Uniform Controlled Dangerous Substances Act

Article III. Regulation of Manufacture, Distribution, Dispensing, Prescribing, Administering and Using for Scientific Purposes of Controlled Dangerous Substances

Anti-Drug Diversion Act

§ 2-309D. Central repository information--Confidentiality--Access--Disclosure--Penalties—
Liability

A. The information collected at the central repository pursuant to the Anti-Drug Diversion Act shall be confidential and shall not be open to the public. Access to the information shall be limited to:

1. Peace officers certified pursuant to Section 3311 of Title 70 of the Oklahoma Statutes who are employed as investigative agents of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control;

2. The United States Drug Enforcement Administration Diversion Group Supervisor;

3. The executive director or chief investigator, as designated by each board, of the following state boards:

a. Board of Podiatric Medical Examiners,

b. Board of Dentistry,

c. State Board of Pharmacy,

d. State Board of Medical Licensure and Supervision,

e. State Board of Osteopathic Examiners,

f. State Board of Veterinary Medical Examiners,

g. Oklahoma Health Care Authority,

h. Department of Mental Health and Substance Abuse Services,

i. Board of Examiners in Optometry,

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j. Board of Nursing,

k. Office of the Chief Medical Examiner, and

l. State Board of Health;

4. A multicounty grand jury properly convened pursuant to the Multicounty Grand Jury Act;

5. Medical practitioners employed by the United States Department of Veterans Affairs, the United States Military, or other federal agencies treating patients in this state; and

6. At the discretion of the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, medical practitioners and their staff, including those employed by the federal government in this state.

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Utah

§ 58-37f-301 (eff. until Oct. 30, 2016)

§ 58-37f-301 (eff. Oct. 31, 2016)

West's Utah Code Annotated (2015)

Title 58. Occupations and Professions

Chapter 37F. Controlled Substance Database Act

Part 3. Access

§ 58-37f-301. Access to database

<Text of Section Effective until October 30, 2016>

(1) The division shall make rules, in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, to:

- (a) effectively enforce the limitations on access to the database as described in this part; and
- (b) establish standards and procedures to ensure accurate identification of individuals requesting information or receiving information without request from the database.

(2) The division shall make information in the database and information obtained from other state or federal prescription monitoring programs by means of the database available only to the following individuals, in accordance with the requirements of this chapter and division rules:

- (a) personnel of the division specifically assigned to conduct investigations related to controlled substance laws under the jurisdiction of the division;
- (b) authorized division personnel engaged in analysis of controlled substance prescription information as a part of the assigned duties and responsibilities of their employment;
- (c) a board member if:
 - (i) the board member is assigned to monitor a licensee on probation; and
 - (ii) the board member is limited to obtaining information from the database regarding the specific licensee on probation;
- (d) a member of a diversion committee established in accordance with Subsection 58-1-404(2) if:
 - (i) the diversion committee member is limited to obtaining information from the database regarding the person whose conduct is the subject of the committee's consideration; and

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(ii) the conduct that is the subject of the committee's consideration includes a violation or a potential violation of Chapter 37, Utah Controlled Substances Act, or another relevant violation or potential violation under this title;

(e) in accordance with a written agreement entered into with the department, employees of the Department of Health:

(i) whom the director of the Department of Health assigns to conduct scientific studies regarding the use or abuse of controlled substances, if the identity of the individuals and pharmacies in the database are confidential and are not disclosed in any manner to any individual who is not directly involved in the scientific studies;

(ii) when the information is requested by the Department of Health in relation to a person or provider whom the Department of Health suspects may be improperly obtaining or providing a controlled substance; or

(iii) in the medical examiner's office;

(f) in accordance with a written agreement entered into with the department, a designee of the director of the Department of Health, who is not an employee of the Department of Health, whom the director of the Department of Health assigns to conduct scientific studies regarding the use or abuse of controlled substances pursuant to an application process established in rule by the Department of Health, if:

(i) the designee provides explicit information to the Department of Health regarding the purpose of the scientific studies;

(ii) the scientific studies to be conducted by the designee:

(A) fit within the responsibilities of the Department of Health for health and welfare;

(B) are reviewed and approved by an Institutional Review Board that is approved for human subject research by the United States Department of Health and Human Services; and

(C) are not conducted for profit or commercial gain; and

(D) are conducted in a research facility, as defined by division rule, that is associated with a university or college accredited by one or more regional or national accrediting agencies recognized by the United States Department of Education;

(iii) the designee protects the information as a business associate of the Department of Health; and

(iv) the identity of the prescribers, patients, and pharmacies in the database are de-identified, confidential, not disclosed in any manner to the designee or to any individual who is not directly involved in the scientific studies;

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West's Utah Code Annotated (2015)
Title 58. Occupations and Professions
Chapter 37F. Controlled Substance Database Act
Part 3. Access

§ 58-37f-301. Access to database

<Text of Section Effective October 31, 2016>

(1) The division shall make rules, in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, to:

(a) effectively enforce the limitations on access to the database as described in this part; and

(b) establish standards and procedures to ensure accurate identification of individuals requesting information or receiving information without request from the database.

(2) The division shall make information in the database and information obtained from other state or federal prescription monitoring programs by means of the database available only to the following individuals, in accordance with the requirements of this chapter and division rules:

(a)(i) personnel of the division specifically assigned to conduct investigations related to controlled substance laws under the jurisdiction of the division; and

(ii) the following law enforcement officers, but the division may only provide nonidentifying information, limited to gender, year of birth, and postal ZIP code, regarding individuals for whom a controlled substance has been prescribed or to whom a controlled substance has been dispensed:

(A) a law enforcement agency officer who is engaged in a joint investigation with the division; and

(B) a law enforcement agency officer to whom the division has referred a suspected criminal violation of controlled substance laws;

(b) authorized division personnel engaged in analysis of controlled substance prescription information as a part of the assigned duties and responsibilities of their employment;

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(c) a board member if:

(i) the board member is assigned to monitor a licensee on probation; and

(ii) the board member is limited to obtaining information from the database regarding the specific licensee on probation;

(d) a member of a diversion committee established in accordance with Subsection 58-1-404(2) if:

(i) the diversion committee member is limited to obtaining information from the database regarding the person whose conduct is the subject of the committee's consideration; and

(ii) the conduct that is the subject of the committee's consideration includes a violation or a potential violation of Chapter 37, Utah Controlled Substances Act, or another relevant violation or potential violation under this title;

(e) in accordance with a written agreement entered into with the department, employees of the Department of Health:

(i) whom the director of the Department of Health assigns to conduct scientific studies regarding the use or abuse of controlled substances, if the identity of the individuals and pharmacies in the database are confidential and are not disclosed in any manner to any individual who is not directly involved in the scientific studies;

(ii) when the information is requested by the Department of Health in relation to a person or provider whom the Department of Health suspects may be improperly obtaining or providing a controlled substance; or

(iii) in the medical examiner's office;

(f) in accordance with a written agreement entered into with the department, a designee of the director of the Department of Health, who is not an employee of the Department of Health, whom the director of the Department of Health assigns to conduct scientific studies regarding the use or abuse of controlled substances pursuant to an application process established in rule by the Department of Health, if:

(i) the designee provides explicit information to the Department of Health regarding the purpose of the scientific studies;

(ii) the scientific studies to be conducted by the designee:

(A) fit within the responsibilities of the Department of Health for health and welfare;
(B) are reviewed and approved by an Institutional Review Board that is approved for human subject research by the United States Department of Health and Human Services;
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(C) are not conducted for profit or commercial gain; and

(D) are conducted in a research facility, as defined by division rule, that is associated with a university or college accredited by one or more regional or national accrediting agencies recognized by the United States Department of Education;

(iii) the designee protects the information as a business associate of the Department of Health; and

(iv) the identity of the prescribers, patients, and pharmacies in the database are de-identified, confidential, not disclosed in any manner to the designee or to any individual who is not directly involved in the scientific studies;

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Vermont
18 § 4284

West's Vermont Statutes Annotated (2016)
Title Eighteen. Health
Part 5. Foods and Drugs
Chapter 84A. Vermont Prescription Monitoring System

§ 4284. Protection and disclosure of information

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(b)(1) The Department shall provide only the following persons with access to query the VPMS:

(A) A health care provider, dispenser, or delegate who is registered with the VPMS and certifies that the requested information is for the purpose of providing medical or pharmaceutical treatment to a bona fide current patient.

(B) Personnel or contractors, as necessary for establishing and maintaining the VPMS.

(C) The Medical Director of the Department of Vermont Health Access, for the purposes of Medicaid quality assurance, utilization, and federal monitoring requirements with respect to Medicaid recipients for whom a Medicaid claim for a Schedule II, III, or IV controlled substance has been submitted.

(D) A medical examiner or delegate from the Office of the Chief Medical Examiner, for the purpose of conducting an investigation or inquiry into the cause, manner, and circumstances of an individual's death.

(E) A health care provider or medical examiner licensed to practice in another state, to the extent necessary to provide appropriate medical care to a Vermont resident or to investigate the death of a Vermont resident.

(2) The Department shall provide reports of data available to the Department through the VPMS only to the following persons:

(A) A patient or that person's health care provider, or both, when VPMS reveals that a patient may be receiving more than a therapeutic amount of one or more regulated substances.

(B) A designated representative of a board responsible for the licensure, regulation, or discipline of health care providers or dispensers pursuant to a bona fide specific investigation.

(C) A patient for whom a prescription is written, insofar as the information relates to that patient.

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(D) The relevant occupational licensing or certification authority if the Commissioner reasonably suspects fraudulent or illegal activity by a health care provider. The licensing or certification authority may report the data that are the evidence for the suspected fraudulent or illegal activity to a drug diversion investigator.

(E)(i) The Commissioner of Public Safety, personally, or the Deputy Commissioner of Public Safety, personally, if the Commissioner of Health, personally, or a Deputy Commissioner of Health, personally, makes the disclosure and has consulted with at least one of the patient's health care providers, when the disclosure is necessary to avert a serious and imminent threat to a person or the public.

(ii) The Commissioner of Public Safety, personally, or the Deputy Commissioner of Public Safety, personally, when he or she requests data from the Commissioner of Health, and the Commissioner of Health believes, after consultation with at least one of the patient's health care providers, that disclosure is necessary to avert a serious and imminent threat to a person or the public.

(iii) The Commissioner or Deputy Commissioner of Public Safety may disclose such data received pursuant to this subdivision (E) as is necessary, in his or her discretion, to avert the serious and imminent threat.

(F) A prescription monitoring system or similar entity in another state pursuant to a reciprocal agreement to share prescription monitoring information with the Vermont Department of Health as described in section 4288 of this title.

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