



Types of Authorized Recipients – Worker’s Compensation Specialists

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 worker's compensation specialists. This does not mean that if a particular state is not listed in this memorandum or the accompanying map that worker's compensation specialists in that state are not allowed access to the information. If such persons fall within the definition of "practitioner" or "health care provider" in the state, he or she may qualify for access to the prescription monitoring program database. The following states either specifically include worker's compensation specialists 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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Arizona
§ 36-2604
§ 23-1026

Arizona Revised Statutes Annotated (2016)
Title 36. Public Health and Safety
Chapter 28. Controlled Substances Prescription Monitoring Program
Article 1. General Provisions

§ 36-2604. Use and release of confidential information

A. Except as otherwise provided in this section, prescription information submitted to the board pursuant to this article is confidential and is not subject to public inspection. The board shall establish procedures to ensure the privacy and confidentiality of patients and that patient information that is collected, recorded and transmitted pursuant to this article is not disclosed except as prescribed in this section.

B. The board or its designee shall review the prescription information collected pursuant to this article. If the board or its designee has reason to believe an act of unprofessional or illegal conduct has occurred, the board or its designee shall notify the appropriate professional licensing board or law enforcement or criminal justice agency and provide the prescription information required for an investigation.

C. The board may release data collected by the program to the following:

1. A person who is authorized to prescribe or dispense a controlled substance, or a delegate who is authorized by the prescriber or dispenser, to assist that person to provide medical or pharmaceutical care to a patient or to evaluate a patient.
2. An individual who requests the individual's own prescription monitoring information pursuant to § 12-2293.
3. A medical practitioner regulatory board established pursuant to title 32, chapter 7, 11, 13, 14, 15, 16, 17, 18, 25 or 29.1 Except as required pursuant to subsection B of this section, the board shall provide this information only if the requesting board states in writing that the information is necessary for an open investigation or complaint.
4. A local, state or federal law enforcement or criminal justice agency. Except as required pursuant to subsection B of this section, the board shall provide this information only if the requesting agency states in writing that the information is necessary for an open investigation or complaint.

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5. The Arizona health care cost containment system administration regarding persons who are receiving services pursuant to chapter 29 of this title.² Except as required pursuant to subsection B of this section, the board shall provide this information only if the administration states in writing that the information is necessary for an open investigation or complaint.

6. A person who is serving a lawful order of a court of competent jurisdiction.

7. A person who is authorized to prescribe or dispense a controlled substance and who performs an evaluation on an individual pursuant to § 23-1026.

8. A county medical examiner or alternate medical examiner who is directing an investigation into the circumstances surrounding a death as described in § 11-593 or a delegate who is authorized by the county medical examiner or alternate medical examiner.

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

E. For the purposes of this section, “delegate” means any of the following:

1. A licensed health care professional who is employed in the office of or in a hospital with the prescriber or dispenser.

2. An unlicensed medical records technician, medical assistant or office manager who is employed in the office of or in a hospital with the prescriber or dispenser and who has received training regarding both the health insurance portability and accountability act privacy standards, 45 Code of Federal Regulations part 164, subpart E, and security standards, 45 Code of Federal Regulations part 164, subpart C.

3. A forensic pathologist, medical death investigator or other qualified person who is assigned duties in connection with a death investigation pursuant to § 11-594.

Arizona Revised Statutes Annotated (2016)

Title 23. Labor

Chapter 6. Workers' Compensation

Article 7. Right to Compensation

§ 23-1026. Periodical medical examination of employee; effect of refusal or obstruction of examination or treatment

A. An employee who may be entitled to compensation under this chapter shall submit himself for medical examination from time to time at a place reasonably convenient for the employee, if and when requested by the commission, his employer or the insurance carrier. A place is reasonably convenient even if it is not where the employee resides if it is the place where the employee was

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injured and the employer or the insurance carrier pays in advance the employee's reasonable travel expenses, including the cost of transportation, food, lodging and loss of pay, if applicable.

B. The request for the medical examination shall fix a time and place having regard to the convenience of the employee, his physical condition and his ability to attend. The employee may have a physician present at the examination if procured and paid for by himself.

C. If the employee refuses to submit to the medical examination or obstructs the examination, his right to compensation shall be suspended until the examination has been made, and no compensation shall be payable during or for such period.

D. A physician who makes or is present at the medical examination provided by this section may be required to testify as to the result thereof.

E. Upon appropriate application and hearing, the commission may reduce or suspend the compensation of an employee who persists in unsanitary or injurious practices tending to imperil or retard his recovery, or who refuses to submit to medical or surgical treatment reasonably necessary to promote his recovery.

F. An employee shall be excused from attending a scheduled medical examination if the employee requests a protective order and the administrative law judge finds that the scheduled examination is unnecessary, would be cumulative or could reasonably be timely scheduled with an appropriate physician where the employee resides. If a protective order is requested the burden is on the employer or insurance carrier to establish that a medical examination should be scheduled at a place other than where the employee resides. If an employee has left this state and the employer or insurance carrier pays in advance the employee's reasonable travel expenses, including the cost of transportation, food, lodging and loss of pay, if applicable, the employer or insurance carrier is entitled to have the employee return to this state one time a year for examination or one time following the filing of a petition to reopen.

G. If a physician performs an examination under this section and is provided data from the Arizona state board of pharmacy pursuant to title 36, chapter 28, the physician may disclose that data to the employee, employer, insurance carrier and the commission.

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Montana
§ 39-71-1110

West's Montana Code Annotated (2015)
Title 39. Labor
Chapter 71. Workers' Compensation
Part 11. Treatment by Designated Providers

§ 39-71-1110. Schedule II and III drugs--prescriber obligation

(1) In order to ensure high-quality health care for an individual with a compensable occupational injury or disease, prescriptions for Schedule II or Schedule III drugs identified in Title 50, chapter 32, part 2, may be carefully monitored for potential abuse, dependence, interaction, and diversion. Ongoing prescriptions for Schedule II and Schedule III drugs may be prescribed only by a treating physician.

(2)(a) A treating physician authorized to prescribe prescription drugs may query the prescription drug registry provided for in Title 37, chapter 7, part 15, prior to the initial prescribing or refilling of a Schedule II or Schedule III drug for treatment of a workers' compensation injury or occupational disease. After consulting the prescription drug registry, a treating physician may decline to prescribe or refill a Schedule II or Schedule III drug if, in the treating physician's judgment, the drug should not be prescribed or refilled.

(b) Prior to the initial prescribing of a Schedule II or Schedule III drug, a treating physician may discuss the risks and benefits of the use of the controlled substance, including risk of tolerance and drug dependence, with the patient or the patient's legal guardian.

(c) A treating physician shall note in the patient's medical file each query conducted.

(3) This section does not apply to a health care provider administering a Schedule II or Schedule III drug under the following circumstances:

- (a) immediately prior to or after surgery;
- (b) at the scene of an emergency;
- (c) in a licensed ambulance; or
- (d) in the emergency department or intensive care unit of a licensed hospital.

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North Dakota
§ 19-03.5-03

West's North Dakota Century Code Annotated (2015)
Title 19. Foods, Drugs, Oils, and Compounds
Chapter 19-03.5. Prescription Drug Monitoring Program

§ 19-03.5-03. Access to prescription information

1. Information submitted to the central repository is confidential and may not be disclosed except as provided in this section.

2. The board shall maintain procedures to ensure that the privacy, confidentiality, and security of patient information collected, recorded, transmitted, and maintained is not disclosed except as provided in this section.

3. Unless disclosure is prohibited by law, the board may provide data in the central repository to:

a. A prescriber for the purpose of providing medical care to a patient, a dispenser for the purpose of filling a prescription or providing pharmaceutical care for a patient, a prescriber or dispenser inquiring about the prescriber's or dispenser's own prescribing activity, or a prescriber or dispenser in order to further the purposes of the program;

b. An individual who requests the prescription information of the individual or the individual's minor child;

c. State boards and regulatory agencies that are responsible for the licensing of individuals authorized to prescribe or dispense controlled substances if the board or regulatory agency is seeking information from the central repository that is relevant to an investigation of an individual who holds a license issued by that board or regulatory agency;

d. Local, state, and federal law enforcement or prosecutorial officials engaged in the enforcement of laws relating to controlled substances who seek information for the purpose of an investigation or prosecution of the drug-related activity or probation compliance of an individual;

e. The department of human services for purposes regarding the utilization of controlled substances by a medicaid recipient or establishment and enforcement of child support and medical support;

f. Workforce safety and insurance for purposes regarding the utilization of controlled substances by a claimant;

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- g. Judicial authorities under grand jury subpoena or court order or equivalent judicial process for investigation of criminal violations of controlled substances laws;
 - h. Public or private entities for statistical, research, or educational purposes after the information is de-identified with respect to any prescriber, dispenser, or patient who received a prescription for a controlled substance;
 - i. A peer review committee which means any committee of a health care organization, composed of health care providers, employees, administrators, consultants, agents, or members of the health care organization's governing body, which conducts professional peer review as defined in chapter 23-34; or
 - j. A licensed addiction counselor for the purpose of providing services for a licensed treatment program in this state.
4. The board shall maintain a record of each person who requests information from the central repository. The board may use the records to document and report statistics and outcomes. The board may provide records of the requests for information to:
- a. A board or regulatory agency responsible for the licensing of individuals authorized to prescribe or dispense controlled substances that is engaged in an investigation of the individual who submitted the request for information from the central repository; and
 - b. Local, state, and federal law enforcement or prosecutorial officials engaged in the enforcement of laws relating to controlled substances for the purpose of an active investigation of an individual who requested information from the central repository.

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Ohio
§ 4729.80
ADC 4123-6-21.4

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

(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:

(1) On receipt of a request from a designated representative of a government entity responsible for the licensure, regulation, or discipline of health care professionals with authority to prescribe, administer, or dispense drugs, the board may provide to the representative information from the database relating to the professional who is the subject of an active investigation being conducted by the government entity.

(2) On receipt of a request from a federal officer, or a state or local officer of this or any other state, whose duties include enforcing laws relating to drugs, the board shall provide to the officer information from the database relating to the person who is the subject of an active investigation of a drug abuse offense, as defined in section 2925.01 of the Revised Code, being conducted by the officer's employing government entity.

(3) Pursuant to a subpoena issued by a grand jury, the board shall provide to the grand jury information from the database relating to the person who is the subject of an investigation being conducted by the grand jury.

(4) Pursuant to a subpoena, search warrant, or court order in connection with the investigation or prosecution of a possible or alleged criminal offense, the board shall provide information from the database as necessary to comply with the subpoena, search warrant, or court order.

(5) On receipt of a request from a prescriber or the prescriber's delegate approved by the board, the board shall provide to the prescriber a report of information from the database relating to a patient who is either a current patient of the prescriber or a potential patient of the prescriber based on a referral of the patient to the prescriber, if all of the following conditions are met:

(a) The prescriber certifies in a form specified by the board that it is for the purpose of providing medical treatment to the patient who is the subject of the request;

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(b) The prescriber has not been denied access to the database by the board.

(6) On receipt of a request from a pharmacist or the pharmacist's delegate approved by the board, the board shall provide to the pharmacist information from the database relating to a current patient of the pharmacist, if the pharmacist certifies in a form specified by the board that it is for the purpose of the pharmacist's practice of pharmacy involving the patient who is the subject of the request and the pharmacist has not been denied access to the database by the board.

(7) On receipt of a request from an individual seeking the individual's own database information in accordance with the procedure established in rules adopted under section 4729.84 of the Revised Code, the board may provide to the individual the individual's own database information.

(8) On receipt of a request from a medical director or a pharmacy director of a managed care organization that has entered into a contract with the department of medicaid under section 5167.10 of the Revised Code and a data security agreement with the board required by section 5167.14 of the Revised Code, the board shall provide to the medical director or the pharmacy director information from the database relating to a medicaid recipient enrolled in the managed care organization, including information in the database related to prescriptions for the recipient that were not covered or reimbursed under a program administered by the department of medicaid.

(9) On receipt of a request from the medicaid director, the board shall provide to the director information from the database relating to a recipient of a program administered by the department of medicaid, including information in the database related to prescriptions for the recipient that were not covered or paid by a program administered by the department.

(10) On receipt of a request from a medical director of a managed care organization that has entered into a contract with the administrator of workers' compensation under division (B)(4) of section 4121.44 of the Revised Code and a data security agreement with the board required by section 4121.447 of the Revised Code,¹ the board shall provide to the medical director information from the database relating to a claimant under Chapter 4121., 4123., 4127., or 4131. of the Revised Code assigned to the managed care organization, including information in the database related to prescriptions for the claimant that were not covered or reimbursed under Chapter 4121., 4123., 4127., or 4131. of the Revised Code, if the administrator of workers' compensation confirms, upon request from the board, that the claimant is assigned to the managed care organization.

(11) On receipt of a request from the administrator of workers' compensation, the board shall provide to the administrator information from the database relating to a claimant under Chapter 4121., 4123., 4127., or 4131. of the Revised Code, including information in the database related to prescriptions for the claimant that were not covered or reimbursed under Chapter 4121., 4123., 4127., or 4131. of the Revised Code.

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(12) On receipt of a request from a prescriber or the prescriber's delegate approved by the board, the board shall provide to the prescriber information from the database relating to a patient's mother, if the prescriber certifies in a form specified by the board that it is for the purpose of providing medical treatment to a newborn or infant patient diagnosed as opioid dependent and the prescriber has not been denied access to the database by the board.

(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.

(14) On receipt of a request from a requestor described in division (A)(1), (2), (5), or (6) of this section who is from or participating with another state's prescription monitoring program, the board may provide to the requestor information from the database, but only if there is a written agreement under which the information is to be used and disseminated according to the laws of this state.

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Baldwin's Ohio Administrative Code Annotated (2016)
4123 Workers' Compensation Bureau
Chapter 4123-6. Health Partnership Program (HPP)

4123-6-21.4 Coordinated services program

The bureau, or a self-insuring employer with a point-of-service adjudication system, may establish a coordinated services program (CSP) that requires an injured worker to obtain prescription medications reimbursed by the bureau or self-insuring employer from a single designated pharmacy and/or prescriber.

(A) Placement in a CSP.

(1) The bureau or self-insuring employer with a point-of-service adjudication system may review an injured worker for possible placement in a CSP if a review of his or her claim indicates the injured worker meets one or more of the following criteria:

(a) Use of three or more different prescribers to obtain prescriptions of the same or comparable medications per three month time frame;

(b) Receipt of prescription drugs from more than two different pharmacies per three month time frame;

(c) Monthly receipt of three or more prescriptions including refills for drugs identified by therapeutic drug class as a narcotic analgesic per three month time frame;

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(d) Monthly receipt of more than two concurrent narcotic analgesics in the same therapeutic drug class per three month time frame;

(e) Monthly receipt of more than two narcotic analgesics in the same therapeutic drug class, more than one benzodiazepine, and more than one sedative-hypnotics per three month time frame.

(2) Upon identification of an injured worker meeting one or more of the criteria identified in paragraphs (A)(1)(a) to (A)(1)(e) of this rule, the bureau or self-insuring employer with a point-of-service adjudication system shall obtain a physician review of the injured worker's most recent twelve months history of prescription medications reimbursed by the bureau or self-insuring employer.

(3) If, based on this physician review, the bureau or self-insuring employer with a point-of-service adjudication system determines that the injured worker's utilization of prescription medications during this period was at a frequency or in an amount that was not medically necessary or appropriate under the criteria set forth in paragraphs (B)(1) to (B)(3) of rule 4123-6-16.2 of the Administrative Code, or was potentially unsafe, the bureau or self-insuring employer may place the injured worker in a CSP.

...

(D) Pharmacies participating in the bureau's CSP.

(1) The bureau shall maintain a list of pharmacies participating in the bureau's CSP that are eligible for selection by an injured worker as a designated pharmacy. To participate in the bureau's CSP, a pharmacy must meet the following criteria:

(a) The pharmacy must be enrolled with the bureau and have a signed agreement with the bureau's pharmacy benefits manager.

(b) The pharmacy must enter into a CSP agreement with the bureau.

(2) Pharmacies participating in the bureau's CSP agree to perform the following monitoring activities:

(a) For each injured worker in the bureau's CSP for whom the pharmacy is the designated pharmacy, the pharmacy shall conduct a bimonthly review of the injured worker's OARRS report from the Ohio board of pharmacy (or a similar automated prescription monitoring report from the injured worker's state of residence).

(b) The pharmacy shall notify the injured worker's prescribing physician of any critical findings discovered in the report. Critical findings are indications of any prescription related activity that could cause harm to the patient, including but not limited to:

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(i) Duplication of therapy,

(ii) Excessive doses of concurrent medications,

(iii) Potential drug interactions or potentiation of side effects.

(c) The pharmacy shall notify BWC in writing whenever reports are made under paragraph (D)(2)(b) of this rule.

(d) BWC may request quarterly documentation of the pharmacy's monitoring activities under paragraphs (D)(2)(a) to (D)(2)(d) of this rule.

(3) Pharmacies participating in the CSP may receive compensation from the bureau under the CSP agreement for services provided as part of the CSP.

(4) Pharmacies participating in the bureau's CSP may terminate their CSP agreement with the bureau and discontinue their participation in the bureau's CSP at any time upon not less than thirty days written notice to the bureau. Pharmacies participating in the bureau's CSP may discontinue providing services to an individual injured worker at any time upon not less than thirty days written notice to the bureau, the injured worker, and the injured worker's authorized representative.

(5) The bureau may terminate the CSP agreement of a pharmacy participating in the bureau's CSP in accordance with the terms of the CSP agreement.

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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:

...

(s) the following licensed physicians for the purpose of reviewing and offering an opinion on an individual's request for workers' compensation benefits under Title 34A, Chapter 2, Workers' Compensation Act, or Title 34A, Chapter 3, Utah Occupational Disease Act:

(i) a member of the medical panel described in Section 34A-2-601;

(ii) a physician employed as medical director for a licensed workers' compensation insurer or an approved self-insured employer; or

(iii) a physician offering a second opinion regarding treatment.

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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:

...

(t) the following licensed physicians for the purpose of reviewing and offering an opinion on an individual's request for workers' compensation benefits under Title 34A, Chapter 2, Workers' Compensation Act, or Title 34A, Chapter 3, Utah Occupational Disease Act:

- (i) a member of the medical panel described in Section 34A-2-601;**
- (ii) a physician employed as medical director for a licensed workers' compensation insurer or an approved self-insured employer; or**
- (iii) a physician offering a second opinion regarding treatment.**

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Washington
§ 70.225.040

West's Revised Code of Washington Annotated (2016)
Title 70. Public Health and Safety
Chapter 70.225. Prescription Monitoring Program

§ 70.225.040. Confidentiality of prescription information--Procedures--Immunity when acting in good faith

(1) Prescription information submitted to the department must be confidential, in compliance with chapter 70.02 RCW and federal health care information privacy requirements and not subject to disclosure, except as provided in subsections (3) and (4) of this section.

(2) The department must 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 in subsections (3) and (4) of this section.

(3) The department may provide data in the prescription monitoring program to the following persons:

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

(b) An individual who requests the individual's own prescription monitoring information;

(c) Health professional licensing, certification, or regulatory agency or entity;

(d) Appropriate law enforcement or prosecutorial officials, including local, state, and federal officials and officials of federally recognized tribes, who are engaged in a bona fide specific investigation involving a designated person;

(e) Authorized practitioners of the department of social and health services and the health care authority regarding medicaid program recipients;

(f) The director or director's designee within the department of labor and industries regarding workers' compensation claimants;

(g) The director or the director's designee within the department of corrections regarding offenders committed to the department of corrections;

(h) Other entities under grand jury subpoena or court order;

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(i) Personnel of the department for purposes of administration and enforcement of this chapter or chapter 69.50 RCW; and

(j) Personnel of a test site that meet the standards under RCW 70.225.070 pursuant to an agreement between the test site and a person identified in (a) of this subsection to provide assistance in determining which medications are being used by an identified patient who is under the care of that person.

(4) The department may provide data to public or private entities for statistical, research, or educational purposes after removing information that could be used to identify individual patients, dispensers, prescribers, and persons who received prescriptions from dispensers.

(5) A dispenser or practitioner acting in good faith is immune from any civil, criminal, or administrative liability that might otherwise be incurred or imposed for requesting, receiving, or using information from the program.

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