

**TITLE 9. HEALTH SERVICES**  
**CHAPTER 10. DEPARTMENT OF HEALTH SERVICES**  
**HEALTH CARE INSTITUTIONS: LICENSING**  
**ARTICLE 1. GENERAL**

Section

R9-10-120. Opioid Prescribing and Treatment

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## ARTICLE 1. GENERAL

### R9-10-120. Opioid Prescribing and Treatment

A. This Section does not apply to a health care institution licensed under Article 20 of this Chapter.

B. In addition to the definitions in A.R.S. § §§ 32-3248.01 and 36-401(A) and R9-10-101, the following definitions apply in this Section:

1. “Episode of care” means medical services, nursing services, or health-related services provided by a health care institution to a patient for a specific period of time, ending in discharge, ~~or~~ the completion of the patient’s treatment plan, or 90 days from the start of service provision to the patient, whichever is later.
2. “Order” means to issue written, verbal, or electronic instructions for a specific dose of a specific medication in a specific quantity and route of administration to be obtained and administered to a patient in a health care institution.

C. An administrator of a health care institution where opioids are prescribed or ordered as part of treatment shall:

1. Establish, document, and implement policies and procedures for prescribing or ordering an opioid as part of treatment, to protect the health and safety of a patient, that:
  - a. Cover which personnel members may prescribe or order an opioid in treating a patient and the required knowledge and qualifications of these personnel members;
  - b. As applicable and except when contrary to medical judgment for a patient, are consistent with A.R.S. § 32-3248.01 and the Arizona Opioid Prescribing Guidelines or national opioid-prescribing guidelines, such as guidelines developed by the:
    - i. Centers for Disease Control and Prevention, or
    - ii. U.S. Department of Veterans Affairs and the U.S. Department of Defense;
  - c. ~~Include~~ As applicable, include how, when, and by whom:
    - i. A patient’s profile on the Arizona Board of Pharmacy Controlled Substances Prescription Monitoring Program database is reviewed;
    - ii. An assessment is conducted of a patient’s substance use risk;

- iii. The potential risks, adverse outcomes, and complications, including death, associated with the use of opioids are explained to a patient or the patient's representative;
  - iv. Alternatives to a prescribed or ordered opioid are explained to a patient or the patient's representative;
  - v. Informed consent is obtained from a patient or the patient's representative and, if applicable, in what situations, described in subsection (G), ~~or (H), or (I)~~, informed consent would not be obtained before an opioid is prescribed or ordered for a patient;
  - vi. A patient receiving an opioid is monitored; and
  - vii. The actions taken according to subsections (C)(1)(c)(i) through (vi) are documented;
- d. Address conditions that may impose a higher risk to a patient when prescribing or ordering an opioid as part of treatment, including:
- i. Concurrent use of a benzodiazepine or other sedative-hypnotic medication,
  - ii. History of substance use disorder,
  - iii. Co-occurring behavioral health issue, or
  - iv. Pregnancy;
- e. Cover the criteria for co-prescribing a short-acting opioid antagonist for a patient who is not an inpatient, as defined in R9-10-201;
- f. Include that, if continuing control of a patient's pain after discharge is medically indicated due to the patient's medical condition, a method for continuing pain control will be addressed as part of discharge planning;
- g. Include the frequency of the following for a patient being prescribed or ordered an opioid for longer than a 30-calendar-day period:
- i. Face-to-face interactions with the patient,
  - ii. Conducting an assessment of a patient's substance use risk,
  - iii. Renewal of a prescription or order for an opioid without a face-to-face interaction with the patient, and
  - iv. Monitoring the effectiveness of the treatment;



- f. The name and signature of the individual explaining the use of an opioid to the patient; and
  - g. The signature of the patient or the patient's representative and the date signed.
- D.** Except as provided in subsection (H) or (I), an administrator of a health care institution where opioids are prescribed as part of treatment shall ensure that a medical practitioner authorized by policies and procedures to prescribe an opioid in treating a patient:
- 1. Before prescribing an opioid for a patient of the health care institution:
    - a. Conducts a physical examination of the patient or reviews the documentation from a physical examination conducted during the patient's same episode of care;
    - b. Except as exempted by A.R.S. § 36-2606(G), reviews the patient's profile on the Arizona Board of Pharmacy Controlled Substances Prescription Monitoring Program database;
    - c. Conducts an assessment of the patient's substance use risk or reviews the documentation from an assessment of the patient's substance use risk conducted during the same episode of care by an individual licensed under A.R.S. Title 32 and authorized by policies and procedures to conduct an assessment of the patient's substance use risk;
    - d. Explains to the patient or the patient's representative the risks and benefits associated with the use of opioids or ensures that the patient or the patient's representative understands the risks and benefits associated with the use of opioids, as explained to the patient or the patient's representative by an individual licensed under A.R.S. Title 32 and authorized by policies and procedures to explain to the patient or the patient's representative the risks and benefits associated with the use of opioids;
    - e. Explains If applicable, explains alternatives to a prescribed opioid; and
    - f. Obtains informed consent from the patient or the patient's representative that meets the requirements in subsection (C)(4), including the potential risks, adverse outcomes, and complications associated with the concurrent use of an opioid and a benzodiazepine or another sedative-hypnotic medication, if the patient:
      - i. Is also prescribed or ordered a sedative-hypnotic medication, or
      - ii. Has been prescribed a sedative-hypnotic medication by another medical practitioner;

2. Includes the following information in the patient's medical record, an existing treatment plan, or a new treatment plan developed for the patient: **[This would also be the information required to justify higher doses, as allowed by 32-3248.01.]**
    - a. The patient's diagnosis;
    - b. The patient's medical history, including co-occurring disorders;
    - c. The opioid to be prescribed;
    - d. Other medications or herbal supplements being taken by the patient;
    - e. If applicable:
      - i. The effectiveness of the patient's current treatment,
      - ii. The duration of the current treatment, and
      - iii. Alternative treatments tried by or planned for the patient;
    - f. The expected benefit of the treatment and, if applicable, the benefit of the new treatment compared with continuing the current treatment; and
    - g. Other factors relevant to the patient's being prescribed an opioid; and
  3. If applicable, specifies in the patient's discharge plan how medically indicated pain control will occur after discharge to meet the patient's needs.
- E.** Except as provided in subsection (G) or (H), an administrator of a health care institution where opioids are ordered for administration to a patient in the health care institution as part of treatment shall ensure that a medical practitioner authorized by policies and procedures to order an opioid in treating a patient:
1. Before ordering an opioid for a patient of the health care institution:
    - a. Conducts a physical examination of the patient or reviews the documentation from a physical examination conducted:
      - i. During the patient's same episode of care; or
      - ii. Within the previous 30 calendar days, at a health care institution transferring the patient to the health care institution or by the medical practitioner who referred the patient for admission to the health care institution;
    - b. Except as exempted by A.R.S. § 36-2606(G), reviews the patient's profile on the Arizona Board of Pharmacy Controlled Substances Prescription Monitoring Program database;
    - c. ~~Conducts~~ **Conducts** If medically appropriate based on the physical examination in subsection (E)(1)(a) and the patient's medical history, conducts an assessment of

- the patient's substance use risk or reviews the documentation from an assessment of the patient's substance use risk conducted within the previous 30 calendar days by an individual licensed under A.R.S. Title 32 and authorized by policies and procedures to conduct an assessment of the patient's substance use risk;
- d. ~~Explains to the patient or the patient's representative the risks and benefits associated with the use of opioids or ensures~~ Ensures that the patient or the patient's representative understands the risks and benefits associated with the use of opioids, as explained to the patient or the patient's representative ~~by an individual licensed under A.R.S. Title 32 and authorized by~~ according to policies and procedures ~~to explain to the patient or the patient's representative the risks and benefits associated with the use of opioids;~~ and
  - e. If applicable, explains alternatives to an ordered opioid; and
  - f. ~~Obtains informed consent from the patient or the patient's representative, according to subsection (D)(1)(f); and~~
2. Includes the following information in the patient's medical record, an existing treatment plan, or a new treatment plan developed for the patient:
- a. The patient's diagnosis;
  - b. The patient's medical history, including co-occurring disorders;
  - c. The opioid being ordered and the reason for the order;
  - d. Other medications or herbal supplements being taken by the patient; and
  - e. If applicable:
    - i. The effectiveness of the patient's current treatment,
    - ii. The duration of the current treatment,
    - iii. Alternative treatments tried by or planned for the patient,
    - iv. The expected benefit of a new treatment compared with continuing the current treatment, and
    - v. Other factors relevant to the patient's being ordered an opioid.
- F.** For a health care institution where opioids are administered as part of treatment or where a patient is provided assistance in the self-administration of medication for a prescribed opioid, including a health care institution in which an opioid may be prescribed or ordered as part of treatment, an administrator, a manager as defined in R9-10-801, or a provider, as applicable to the health care institution, shall:

1. Establish, document, and implement policies and procedures for administering an opioid as part of treatment or providing assistance in the self-administration of medication for a prescribed opioid, to protect the health and safety of a patient, that:
  - a. Cover which personnel members may administer an opioid in treating a patient and the required knowledge and qualifications of these personnel members;
  - b. Cover which personnel members may provide assistance in the self-administration of medication for a prescribed opioid and the required knowledge and qualifications of these personnel members;
  - c. Include how, when, and by whom a patient's need for opioid administration is assessed;
  - d. Include how, when, and by whom a patient receiving an opioid is monitored; and
  - e. Cover how, when, and by whom the actions taken according to subsections (F)(1)(c) and (d) are documented;
2. Include in the plan for the health care institution's quality management program a process for:
  - a. Review of incidents of opioid-related adverse reactions or other negative outcomes a patient experiences or opioid-related deaths, and
  - b. Surveillance and monitoring of adherence to the policies and procedures in subsection (F)(1);
3. Except as prohibited by Title 42 Code of Federal Regulations, Chapter I, Subchapter A, Part 2, or as provided in subsection (H)(1), ensure that, if a patient's death may be related to an opioid administered as part of treatment, written notification, in a Department-provided format, is provided to the Department of the patient's death within one working day after the patient's death; and
4. Except as provided in subsection (H), ensure that an individual authorized by policies and procedures to administer an opioid in treating a patient or to provide assistance in the self-administration of medication for a prescribed opioid:
  - a. Before administering an opioid or providing assistance in the self-administration of medication for a prescribed opioid in compliance with an order as part of the treatment for a patient, identifies the patient's need for the opioid;
  - b. Monitors the patient's response to the opioid; and
  - c. Documents in the patient's medical record:





- a. For a patient receiving a surgical procedure or other invasive procedure; or
- b. When changing the type, dosage, or route of administration of an opioid, which had previously been ordered by a medical practitioner of the health care institution for a patient according to the requirements in subsection (E), to meet the patient's needs.

**I.** The requirements in subsections (D)(1)(c) through (f) do not apply to a health care institution's prescribing an opioid as part of treatment for a patient with chronic, intractable pain who has had an established health professional-patient relationship with the prescribing medical practitioner for at least 90 days before the opioid is prescribed.

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