The table of contents on the first page contains quick links to the referenced page numbers in this Chapter. Refer to the notes at the end of a Section to learn about the history of a rule as it was published in the Arizona Administrative Register.

Sections, Parts, Exhibits, Tables or Appendices codified in this supplement. The list provided contains quick links to the updated rules.

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The release of this Chapter in supplement 18-1 replaces supplement 17-3, 1-17 pages

Please note that the Chapter you are about to replace may have rules still in effect after the publication date of this supplement. Therefore, all superseded material should be retained in a separate binder and archived for future reference.
PREFACE

Under Arizona law, the Department of State, Office of the Secretary of State (Office), accepts state agency rule filings and is the publisher of Arizona rules. The Office of the Secretary of State does not interpret or enforce rules in the Administrative Code. Questions about rules should be directed to the state agency responsible for the promulgation of the rule.

Scott Cancelosi, Director
ADMINISTRATIVE RULES DIVISION

RULES
The definition for a rule is provided for under A.R.S. § 41-1001. “Rule’ means an agency statement of general applicability that implements, interprets, or prescribes law or policy, or describes the procedures or practice requirements of an agency.”

THE ADMINISTRATIVE CODE
The Arizona Administrative Code is where the official rules of the state of Arizona are published. The Code is the official codification of rules that govern state agencies, boards, and commissions.

The Code is separated by subject into titles. Titles are divided into chapters. A chapter includes state agency rules. Rules in chapters are divided into Articles, then Sections. The “R” stands for “rule” with a sequential numbering and lettering outline separated into subsections.

Rules are codified quarterly in the Code. Supplement release dates are printed on the footers of each chapter.

First Quarter: January 1 - March 31
Second Quarter: April 1 - June 30
Third Quarter: July 1 - September 30
Fourth Quarter: October 1 - December 31
For example, the first supplement for the first quarter of 2018 is cited as Supp. 18-1.

Please note: The Office publishes by chapter, not by individual rule section. Therefore there might be only a few sections codified in each chapter released in a supplement. Historical notes at the end of a section provide an effective date and information when a rule was last updated.

AUTHENTICATION OF PDF CODE CHAPTERS
The Office began to authenticate chapters of the Administrative Code in Supp. 18-1 to comply with A.R.S. § 41-1012(B) and A.R.S. § 5302(1), (2)(d) through (e), and (3)(d) through (e).

A certification verifies the authenticity of each Code chapter posted as it is released by the Office of the Secretary of State. The authenticated pdf of the Code includes an integrity mark with a certificate ID. Users should check the validity of the signature, especially if the pdf has been downloaded. If the digital signature is invalid it means the document’s content has been compromised.

HOW TO USE THE CODE
Rules may be in effect before a supplement is released by the Office. Therefore, the user should refer to issues of the Arizona Administrative Register for recent updates to rule Sections.

ARIZONA REVISED STATUTE REFERENCES
The Arizona Revised Statutes (A.R.S.) are available online at the Legislature’s website, www.azleg.gov. An agency’s authority note to make rules is often included at the beginning of a chapter. Other Arizona statutes may be referenced in rule under the A.R.S. acronym.

SESSION LAW REFERENCES
Arizona Session Law references in a chapter can be found at the Secretary of State’s website, under Services-> Legislative Filings.

EXEMPTIONS FROM THE APA
It is not uncommon for an agency to be exempt from the steps outlined in the rulemaking process as specified in the Arizona Administrative Procedures Act, also known as the APA (Arizona Revised Statutes, Title 41, Chapter 6, Articles 1 through 10). Other agencies may be given an exemption to certain provisions of the Act.

An agency’s exemption is written in law by the Arizona State Legislature or under a referendum or initiative passed into law by Arizona voters.

When an agency files an exempt rulemaking package with our Office it specifies the law exemption in what is called the preamble of rulemaking. The preamble is published in the Register online at www.azsos.gov/rules, click on the Administrative Register link.

Editor’s notes at the beginning of a chapter provide information about rulemaking sections made by exempt rulemaking. Exempt rulemaking notes are also included in the historical note at the end of a rulemaking Section.

The Office makes a distinction to certain exemptions because some rules are made without receiving input from stakeholders or the public. Other exemptions may require an agency to propose exempt rules at a public hearing.

EXEMPTIONS AND PAPER COLOR
At one time the office published exempt rules on either blue or green paper. Blue meant the authority of the exemption was given by the Legislature; green meant the authority was determined by a court order. In 2001 the Office discontinued publishing rules using these paper colors.

PERSONAL USE/COMMERCIAL USE
This chapter is posted as a public courtesy online, and is for private use only. Those who wish to use the contents for resale or profit should contact the Office about Commercial Use fees. For information on commercial use fees review A.R.S. § 39-121.03 and 1 A.A.C. 1, R1-1-113.

Rhonda Paschal, managing rules editor, assisted with the editing of this chapter.
ARTICLE 1. GENERAL PROVISIONS

Article 1, consisting of Sections R4-16-101 through R4-16-106, adopted effective June 1, 1984.

Former Article 1, consisting of Sections R4-16-01 through R4-16-10, repealed effective June 1, 1984 (Supp. 84-3).

R4-16-101. Definitions
R4-16-102. Continuing Medical Education
R4-16-103. Rehearing or Review of Board Decision
R4-16-104. Recodified
R4-16-105. Recodified
R4-16-106. Recodified
R4-16-107. Recodified
R4-16-108. Recodified
R4-16-109. Recodified

ARTICLE 2. LICENSURE

Article 2 heading, recodified to Article 3 heading, at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

Article 2, consisting of Sections R4-16-201 through R4-16-205, adopted effective September 22, 1995 (Supp. 95-3).

Section
R4-16-201. Application for License by Examination or Endorsement
R4-16-201. 1. Application for Renewal of License
R4-16-202. Application and Reapplication for Pro Bono Registration
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R4-16-206. Time Frames for Licenses, Permits, and Registrations
R4-16-207. Repealed
Table 1. Time Frames
R4-16-208. Recodified

ARTICLE 3. DISPENSING OF DRUGS

Article 3 heading, recodified from Article 2 heading, at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

Article 3, consisting of Sections R4-16-301 through R4-16-303, adopted effective February 2, 2000 (Supp. 00-1).

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ARTICLE 4. MEDICAL ASSISTANTS

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R4-16-410. Recodified

ARTICLE 5. EXECUTIVE DIRECTOR DUTIES

Article 5, consisting of Sections R4-16-501 through R4-16-505, renumbered by exempt rulemaking at 11 A.A.R. 1056, effective February 18, 2005 (Supp. 05-1).

Article 5, consisting of Sections R4-16-501 through R4-16-505, made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2).

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Article 7, consisting of Sections R4-16-701 through R4-16-707, made by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).

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ARTICLE 1. GENERAL PROVISIONS

R4-16-101. Definitions

Unless the context otherwise requires, definitions prescribed under A.R.S. § 32-1401 and the following apply to this Chapter:

1. “ACLS” means advanced cardiac life support performed according to certification standards of the American Heart Association.
2. “Agent” means an item or element that causes an effect.
3. “Approved medical assistant training program” means a program accredited by any of the following:
   a. The Commission on Accreditation of Allied Health Education Programs;
   b. The Accrediting Bureau of Health Education Schools;
   c. A medical assisting program accredited by any accrediting agency recognized by the United States Department of Education; or
   d. A training program:
      i. Designed and offered by a licensed allopathic physician;
      ii. That meets or exceeds any of the prescribed accrediting programs in subsection (a), (b), or (c); and
      iii. That verifies the entry-level competencies of a medical assistant prescribed under R4-16-402(A).
4. “Auscultation” means the act of listening to sounds within the human body either directly or through use of a stethoscope or other means.
5. “BLS” means basic life support performed according to certification standards of the American Heart Association.
6. “Capnography” means monitoring the concentration of exhaled carbon dioxide of a sedated patient to determine the adequacy of the patient’s ventilatory function.
7. “Deep sedation” means a drug-induced depression of consciousness during which a patient:
   a. Cannot be easily aroused, but
   b. Responds purposefully following repeated or painful stimulation, and
   c. May partially lose the ability to maintain ventilatory function.
8. “Discharge” means a written or electronic documented termination of office-based surgery to a patient.
10. “Emergency” means an immediate threat to the life or health of a patient.
11. “Emergency drug” means a drug that is administered to a patient in an emergency.
12. “General Anesthesia” means a drug-induced loss of consciousness during which a patient:
   a. Is unarousable even with painful stimulus; and
   b. May partially or completely lose the ability to maintain ventilatory, neuromuscular, or cardiovascular function or airway.
13. “Health care professional” means a registered nurse defined in A.R.S. § 32-1601, registered nurse practitioner defined in A.R.S. § 32-1601, physician assistant defined in A.R.S. § 32-2501, and any individual authorized to perform surgery according to A.R.S. Title 32 who participates in office-based surgery using sedation at a physician’s office.
14. “Informed consent” means advising a patient of the:
   a. Purpose for and alternatives to the office-based surgery using sedation,
   b. Associated risks of office-based surgery using sedation, and
   c. Possible benefits and complications from the office-based surgery using sedation.
15. “Inpatient” has the same meaning as in A.A.C. R9-10-201.
16. “Malignant hyperthermia” means a life-threatening condition in an individual who has a genetic sensitivity to inhalant anesthetics and depolarizing neuromuscular blocking drugs that occurs during or after the administration of an inhalant anesthetic or depolarizing neuromuscular blocking drug.
17. “Minimal Sedation” means a drug-induced state during which:
   a. A patient responds to verbal commands,
   b. Cognitive function and coordination may be impaired, and
   c. A patient’s ventilatory and cardiovascular functions are unaffected.
18. “Moderate Sedation” means a drug-induced depression of consciousness during which:
   a. A patient responds to verbal commands or light tactile stimulation, and
   b. No interventions are required to maintain ventilatory or cardiovascular function.
20. “Office-based surgery” means a medical procedure conducted in a physician’s office or other outpatient setting that is not part of a licensed hospital or licensed ambulatory surgical center. (A.R.S. § 32-1401(20)).
21. “PALS” means pediatric life support performed according to certification standards of the American Academy of Pediatrics or the American Heart Association.
23. “Physician” has the same meaning as doctor of medicine as defined in A.R.S. § 32-1401.
24. “Rescue” means to correct adverse physiologic consequences of deeper than intended level of sedation and return the patient to the intended level of sedation.
25. “Sedation” means minimum sedation, moderate sedation, or deep sedation.
26. “Staff member” means an individual who:
   a. Is not a health care professional, and
   b. Assists with office-based surgery using sedation under the supervision of the physician performing the office-based surgery using sedation.
27. “Transfer” means a physical relocation of a patient from a physician’s office to a licensed health care institution.

Historical Note

Former Rule 12. Former Section R4-16-01 repealed, new Section R4-16-101 adopted effective June 1, 1984 (Supp. 84-3). Section repealed, new Section renumbered from R4-16-103 effective September 22, 1995 (Supp. 95-3). Amended by final rulemaking at 8 A.A.R. 830, February 7, 2002 (Supp. 02-1). Amended by final rulemaking at 8 A.A.R. 4270, effective November 18, 2002 (Supp. 02-3). Former Section R4-16-101 recodified to R4-16-102 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). New Section made by final rulemaking at 12 A.A.R. 823, effective February 23, 2006 (Supp. 06-1). Amended by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).

R4-16-102. Continuing Medical Education
A. A physician holding an active license to practice medicine in this state shall complete 40 credit hours of the continuing medical education required by A.R.S. § 32-1434 during the two calendar years preceding biennial registration.

1. The physician shall ensure at least one of the credit hours of continuing medical education is certified as Category 1, as described in subsection (B)(4), and addresses the effective and safe prescribing of opioids;
2. One hour of credit is allowed for each clock hour of participation in continuing medical education activities, unless otherwise designated in subsection (B); and
3. The physician may not carry excess hours of continuing medical education over to another two-year cycle.

B. A physician may claim continuing medical education for the following:

1. Participating in an internship, residency, or fellowship at a teaching institution approved by the American Medical Association, the Association of American Medical Colleges, or the American Osteopathic Association. A physician may claim one credit hour of continuing medical education for each one day of training in a full-time approved program, or for a less than full-time training on a pro rata basis. In this subsection teaching institutions define “full-time.”
2. Participating in an education program for an advanced degree in a medical or medically-related field in a teaching institution approved by the American Medical Association, the Association of American Medical Colleges, or the American Osteopathic Association. A physician may claim one credit hour of continuing medical education for each one day of full-time study or less than a full-time study on a pro rata basis. In this subsection teaching institutions define “full-time.”
3. Participating in full-time research in a teaching institution approved by the American Medical Association, the Association of American Medical Colleges, or the American Osteopathic Association. A physician may claim one credit hour of continuing medical education for each one day of full-time research, or less than full-time research on a pro rata basis. In this subsection teaching institutions define “full-time.”
4. Participating in an education program certified as Category 1 by an organization accredited by the Accreditation Council for Continuing Medical Education, 515 North State Street, Suite 2150, Chicago, Illinois 60610.
5. Participating in a medical education program designed to provide understanding of current developments, skills, procedures, or treatments related to the practice of medicine, that is provided by an organization or institution accredited by the Accreditation Council for Continuing Medical Education.
6. Serving as an instructor of medical students, house staff, other physicians, or allied health professionals from a hospital or other health care institution with a formal training program, if the instructional activities provide the instructor with understanding of current developments, skills, procedures, or treatments related to the practice of allopathic medicine.
7. Publishing or presenting a paper, report, or book that deals with current developments, skills, procedures, or treatments related to the practice of allopathic medicine. The physician may claim one credit hour for each hour preparing, writing, and presenting materials:
   a. Actually published or presented; and
   b. After the date of publication or presentation.

8. A credit hour may be earned for any of the following activities that provide an understanding of current developments, skills, procedures, or treatments related to the practice of allopathic medicine:
   a. Completing a medical education program based on self-instruction that uses videotapes, audiotapes, films, filmstrips, slides, radio broadcasts, or computers;
   b. Reading scientific journals and books;
   c. Preparing for specialty board certification or recertification examinations;
   d. Participating on a staff or quality of care committee, or utilization review committee in a hospital, health care institution, or government agency.

C. If a physician holding an active license to practice medicine in this state fails to meet the continuing medical education requirements under subsection (A) because of illness, military service, medical or religious missionary activity, or residence in a foreign country, upon written application, shall grant an extension of time to complete the continuing medical education.

D. The Board shall mail to each physician a license renewal form that includes a section regarding continuing medical education compliance. The physician shall sign and return the form certified under penalty of perjury that the continuing medical education requirements under subsection (A) are satisfied for the two-calendar-year period preceding biennial renewal. Failure to receive the license renewal form under subsection (A) shall not relieve the physician of the requirements of subsection (A). The Board may randomly audit a physician to verify compliance with the continuing medical education requirements under subsection (A).

Historical Note
Former Rule 16. Former Section R4-16-02 repealed, new Section R4-16-102 adopted effective June 1, 1984 (Supp. 84-3). Section repealed, new Section renumbered from R4-16-106 effective September 22, 1995 (Supp. 95-3). Amended by final rulemaking at 6 A.A.R. 1881, effective May 3, 2000 (Supp. 00-2). Former Section R4-16-102 recodified to R4-16-103; New Section R4-16-103 recodified from R4-16-101 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). Amended by final rulemaking at 24 A.A.R. 182, effective March 10, 2018 (Supp. 18-1).

R4-16-103. Rehearing or Review of Board Decision

A. A motion for rehearing or review shall be filed as follows:
1. Except as provided in subsection (B), any party in a contested case may file a written motion for rehearing or review of the Board’s decision, specifying generally the grounds upon which the motion is based.
2. A motion for rehearing or review shall be filed with the Board and served no later than 30 days after the decision of the Board.
3. For purposes of this Section, “service” has the same meaning as in A.R.S. § 41-1092.09.
4. For purposes of this Section, a document is deemed filed when the Board receives the document.
5. For purposes of the Section, the terms “contested case” and “party” shall have the same meaning as in A.R.S. § 41-1001.

B. If the Board makes a specific finding that it is necessary for a particular decision to take immediate effect to protect the public health and safety, or that a rehearing or review of the Board’s decision is impracticable or contrary to the public interest, the decision shall be issued as a final decision without
E. A rehearing or review may be granted to all or any of the parties. 
F. A rehearing or review, if granted, shall be a rehearing or review only of the question upon which the decision is found erroneous. An order granting a rehearing or review shall specify in the order the grounds for the rehearing or review.
G. Not later than 15 days after a decision is issued, the Board of its own initiative may order a rehearing or review for any reason that it might have granted a rehearing or review on motion of a party. After giving the parties notice and an opportunity to be heard on the matter, the Board may grant a timely-served motion for a rehearing or review, for a reason not stated in the motion. In either case, the Board shall specify in the order the grounds for the rehearing or review.
H. If a motion for rehearing or review is based upon affidavits, they shall be served with the motion. The opposing party may, within 15 days after service, serve opposing affidavits. The Board may extend this period for a maximum of 20 days either by the Board for good cause, or by the parties by written stipulation. The Board may permit reply affidavits.

Historical Note
Former Rule 17; Amended effective August 19, 1977 (Supp. 77-4). Former Section R4-16-03 repealed, new Section R4-16-103 adopted effective June 1, 1984 (Supp. 84-3). Section R4-16-103 renumbered to R4-16-101 effective September 22, 1995 (Supp. 95-3). New Section adopted effective May 20, 1997 (Supp. 97-2). Amended by final rulemaking at 8 A.A.R. 830, February 7, 2002 (Supp. 02-1). Amended by final rulemaking at 8 A.A.R. 4270, effective November 18, 2002 (Supp. 02-3). Former Section R4-16-103 recodified to R4-16-102 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-104. Recodified

Historical Note
Former Rule 18. Former Section R4-16-04 repealed, new Section R4-16-104 adopted effective June 1, 1984 (Supp. 84-3). Section repealed effective September 22, 1995 (Supp. 95-3). New Section adopted effective January 20, 1998 (Supp. 98-1). Section recodified to R4-16-206 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-105. Recodified

Historical Note
Former Rule 19. Former Section R4-16-05 repealed, new Section R4-16-105 adopted effective June 1, 1984 (Supp. 84-3). Section repealed effective September 22, 1995 (Supp. 95-3). New Section adopted effective January 20, 1998 (Supp. 98-1). Section recodified to R4-16-207 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-106. Recodified

Historical Note
Former Rule 21. Former Section R4-16-06 repealed, new Section R4-16-106 adopted effective June 1, 1984 (Supp. 84-3). Section R4-16-106 renumbered to R4-16-102 effective September 22, 1995 (Supp. 95-3). New Section adopted by final rulemaking at 6 A.A.R. 1881, effective May 3, 2000 (Supp. 00-2). Section recodified to R4-16-201 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-107. Recodified

Historical Note
New Section adopted by final rulemaking at 6 A.A.R. 1881, effective May 3, 2000 (Supp. 00-2). Section recodified to R4-16-202 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-108. Recodified

Historical Note
New Section adopted by final rulemaking at 6 A.A.R. 1881, effective May 3, 2000 (Supp. 00-2). Section recodified to R4-16-203 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

Table 1. Recodified

Historical Note
Table 1 adopted effective January 20, 1998 (Supp. 98-1). Table 1 recodified to the end of Article 2 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-109. Recodified

Historical Note
New Section made by final rulemaking at 8 A.A.R. 830, February 7, 2002 (Supp. 02-1). Amended by final rulemaking at 8 A.A.R. 4270, effective November 18, 2002 (Supp. 02-3). Section recodified to R4-16-205 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

ARTICLE 2. LICENSURE

R4-16-201. Application for Licensure by Examination or Endorsement
A. For purposes of this Article, unless otherwise specified:
1. “ABMS” means American Board of Medical Specialties.
5. “LMCC” means Licentiate of the Medical Council of Canada.
6. “NBME” means National Board of Medical Examiners.
7. “Primary source” means the original source or an approved agent of the original source of a specific credential that can verify the accuracy of a qualification reported by an applicant.
8. “SPEX” means Special Purposes Examination.

B. An applicant for licensure to practice medicine by Step 3 of the USMLE or endorsement shall submit the following information on an application form available on request from the Board and on the Board’s website:
1. Applicant’s full name, Social Security number, business and home addresses, primary e-mail address, business and home telephone numbers, and date and place of birth;
2. Name of the school of medicine from which the applicant graduated and date of graduation;
3. A complete list of the applicant’s internship, residency, and fellowship training;
4. List of all licensing examinations taken;
5. Names of the states, U.S. territories, or provinces in which the applicant has applied for or been granted a license or registration to practice medicine, including license number, date issued, and current status of the license;
6. A statement of whether the applicant:
   a. Has had an application for medical licensure denied or rejected by another state or province licensing board, and if so, an explanation;
   b. Has ever had any disciplinary or rehabilitative action taken against the applicant by another licensing board, including other health professions, and if so, an explanation;
   c. Has had any disciplinary actions, restrictions, or limitations taken against the applicant while participating in any type of training program or by any health care provider, and if so, an explanation;
   d. Has been found in violation of a statute, rule, or regulation of any domestic or foreign governmental agency, and if so, an explanation;
   e. Is currently under investigation by any medical board or peer review body, and if so, an explanation;
   f. Has been subject to discipline resulting in a medical license being revoked, suspended, limited, cancelled during investigation, restricted, or voluntarily surrendered, or resulting in probation or entry into a consent agreement or stipulation and if so, an explanation;
   g. Has had hospital privileges revoked, denied, suspended, or restricted, and if so, an explanation;
   h. Has been named as a defendant in a malpractice matter currently pending or that resulted in a settlement or judgment against the applicant, and if so, an explanation;
   i. Has been subjected to any regulatory disciplinary action, including censure, practice restriction, suspension, sanction, or removal from practice, imposed by any agency of the federal or state government, and if so, an explanation;
   j. Has had the authority to prescribe, dispense, or administer medications limited, restricted, modified, denied, surrendered, or revoked by a federal or state agency as a result of disciplinary or other adverse action, and if so, an explanation;
   k. Has been found guilty or entered into a plea of no contest to a felony, a misdemeanor involving moral turpitude in any state, and if so, an explanation;
   l. Whether the applicant is currently certified by any of the American Board of Medical Specialties;
7. Whether the applicant is currently certified by any of the American Board of Medical Specialties;
8. The applicant’s intended specialty;
9. Consistent with the Board’s authority at A.R.S. § 32-1422(B), other information the Board may deem necessary to evaluate the applicant fully;
10. Whether the applicant completed a training unit prescribed by the Board regarding the requirements of A.R.S. Title 32, Chapter 13 and this Chapter;
11. In addition to the answers provided under subsections (B)(1) through (B)(10), the applicant shall answer the following confidential question:
   a. Whether the applicant has received treatment within the last five years for use of alcohol or a controlled substance, prescription-only drug, or dangerous drug or narcotic or a physical, mental, emotional, or nervous disorder or condition that currently affects the applicant’s ability to exercise the judgment and skills of a medical professional; and
   b. If the answer to subsection (B)(11)(a) is yes:
      i. A detailed description of the use, disorder, or condition;
      ii. An explanation of whether the use, disorder, or condition is reduced or ameliorated because the applicant receives ongoing treatment and if so, the name and contact information for all current treatment providers and for all monitoring or support programs in which the applicant is currently participating; and
   c. A copy of any public or confidential agreement or order relating to the use, disorder, or condition, issued by a licensing agency or health care institution within the last five years, if applicable; and
12. A notarized statement, signed by the applicant, verifying the truthfulness of the information provided, and that the applicant has not engaged in any acts prohibited by Arizona law or Board rules, and authorizing release of any required records or documents to complete application review.

C. In addition to the application form required under subsection (B), an applicant for licensure to practice medicine by Step 3 of the USMLE or endorsement shall submit the following:
1. A notarized copy of the applicant’s birth certificate or passport;
2. Evidence of legal name change if the applicant’s legal name is different from that shown on the document submitted under subsection (C)(1);
3. Documentation listed under A.R.S. § 41-1080(A) showing that the applicant’s presence in the U.S. is authorized under federal law;
4. Complete list of all hospital affiliations and medical employment for the five years before the date of application;
5. Verification of any medical malpractice matter currently pending or resulting in a settlement or judgment against the applicant, including a copy of the complaint and either the agreed terms of settlement or the judgment and a narrative statement specifying the nature of the occurrence resulting in the medical malpractice action. An applicant who is unable to obtain a document required under this subsection may apply under subsection (E) a waiver of the requirement;
6. A full set of fingerprints and the processing charge specified in R4-16-205;
7. A paper or digital headshot photograph of the applicant taken no more than 60 days before the date of application; and
8. The fee authorized under A.R.S. § 32-1436 and specified in R4-16-205.

D. In addition to the requirements of subsections (B) and (C), an applicant for licensure to practice medicine by Step 3 of the USMLE or endorsement shall have the following submitted to the Board, electronically or in hard copy, by the primary source, ECFMG, Veridoc, or FCVS:
1. Official transcript or other authentication of graduation from a school of medicine;
2. Verification of completion of postgraduate training;
3. Verification of ECFMG certification if the applicant graduated from an unapproved school of medicine;
4. Examination and Board history report scores for USMLE, FLEX, NBME, and SPEX;
5. Verification of LMCC exam score or state written exam score;
6. Verification of licensure from every state in which the applicant has ever held a medical license;
7. Verification of all hospital affiliations during the five years before the date of application. Under A.R.S. § 32-1422(A)(11)(b), this verification is required to be on the hospital’s official letterhead or the electronic equivalent; and
8. Verification of all medical employment during the five years before the date of application. Under A.R.S. § 32-1422(A)(11)(b), this verification may be submitted by the employer.

E. As provided under A.R.S. § 32-1422(F), the Board may waive a documentation requirement specified under subsections (C)(5) and (D).
1. To obtain a waiver under this subsection, an applicant shall submit a written request that includes the following information:
   a. Applicant’s name;
   b. Date of request;
   c. Document required under subsection (C)(5) or (D) for which waiver is requested;
   d. Detailed description of efforts made by the applicant to provide the document as required under subsection (C)(5) or (D);
   e. Reason the applicant’s inability to provide the document as required under subsection (C)(5) or (D) is due to no fault of the applicant; and
   f. If applicable, documents that support the request for waiver.
2. The Board shall consider the request for waiver at its next regularly scheduled meeting.
3. In determining whether to grant the request for waiver, the Board shall consider whether the applicant:
   a. Made appropriate and sufficient effort to satisfy the requirement under subsection (C)(5) or (D); and
   b. Demonstrated that compliance with the requirement under subsection (C)(5) or (D) is not possible because:
      i. The entity responsible for issuing the required document no longer exists;
      ii. The original of the required document was destroyed by accident or natural disaster;
      iii. The entity responsible for issuing the required document is unable to provide verification because of armed conflict or political strife; or
   iv. Another valid reason beyond the applicant’s control prevents compliance with the requirement under subsection (C)(5) or (D).
4. In determining whether to grant the request for waiver, the Board shall:
   a. Consider whether it is possible for the Board to obtain the required document from other source; and
   b. Request the applicant to obtain and provide additional information the Board believes will facilitate the Board’s decision.
5. If the Board determines the applicant is unable to comply with a requirement under subsection (C)(5) or (D) in spite of the applicant’s best effort and for a reason beyond the applicant’s control, the Board may grant the request for waiver and include the decision in the Board’s official record for the applicant.
6. The Board shall provide the applicant with written notice of its decision regarding the request for waiver. The Board’s decision is not subject to review or appeal.

F. As provided under A.R.S. § 32-1426(B), the Board may require an applicant for licensure by endorsement who passed an examination specified in A.R.S. § 32-1426(A) more than ten years before the date of application to provide evidence the applicant is able to engage safely in the practice of medicine. The Board may consider one or more of the following to determine whether the applicant is able to engage safely in the practice of medicine:
1. If an applicant is board certified by one of the specialties recognized by the ABMS, this criteria is considered met.
2. If an applicant obtains a passing score on a SPEX examination, this criteria is considered met.
3. The Board may also consider any combination of the following:
   a. The applicant’s records,
   b. The applicant’s practice history,
   c. A physical or psychological assessment of the applicant.

Historical Note
Section 4. A.A.C. 16

2. If the answer to subsection (C)(1) is yes:
   a. A detailed description of the use, disorder, or condition; and
   b. An explanation of whether the use, disorder, or condition is reduced or ameliorated because the applicant receives ongoing treatment and if so, the name and contact information for all current treatment providers and for all monitoring or support programs in which the applicant is currently participating; and
   c. A copy of any public or confidential agreement or order relating to the use, disorder, or condition, issued by a licensing agency or health care institution since the last renewal, if applicable.

D. To renew a license, a licensee shall submit the following with the required application form:
   1. If the document submitted under R4-16-201(C)(3) was a limited form of work authorization issued by the federal government, evidence that the licensee’s presence in the U.S. continues to be authorized under federal law;
   2. The renewal fee specified under R4-16-205 and, if applicable, the penalty fee for late renewal; and
   3. An attestation that all information submitted is correct.

Historical Note

New Section made by final exempt rulemaking at 21 A.A.R. 2678, effective October 15, 2015 (Supp. 15-4).
Amended by final rulemaking at 24 A.A.R. 182, effective March 10, 2018 (Supp. 18-1).

R4-16-202. Application and Reapplication for Pro Bono Registration

A. An applicant for a pro bono registration to practice medicine for a maximum of 60 days in a calendar year in Arizona shall submit the following information on an application form available on request from the Board and on the Board’s web site:
   1. Applicant’s full name, Social Security number, business and home addresses, primary e-mail address, and business and home telephone numbers;
   2. List of all states, U.S. territories, and provinces in which the applicant holds an active medical license;
   3. A statement verifying that the applicant:
      a. Agrees to render all medical services without accepting a fee or salary; or
      b. Agrees to perform only initial or follow-up examinations at no cost to the patient or the patient’s family through a charitable organization;

B. In addition to the application form required under subsection (A), an applicant for a pro bono registration to practice medicine shall submit documentation listed under A.R.S. § 41-1080(A) showing that the applicant’s presence in the U.S. is authorized under federal law.

C. An applicant may make application for a pro bono registration annually. A previously registered applicant may apply for a pro bono registration by submitting the following information on an application form available on request from the Board and on the Board’s web site:
   1. Applicant’s full name, home address and telephone number, and primary e-mail address;
   2. Number of previous pro bono registrations;
   3. Name of each state, U.S. territory, and province in which the applicant holds an active medical license;
   4. A statement whether since issuance of the last pro bono registration:
      a. Any disciplinary action has been taken against the applicant, and
      b. Any unresolved complaints are currently pending against the applicant with any state board; and
   5. If the document submitted under R4-16-202(B) was a limited form of work authorization issued by the federal government, evidence that the applicant’s presence in the U.S. continues to be authorized under federal law.

Historical Note

Adopted effective September 22, 1995 (Supp. 95-3). Amended by final rulemaking at 8 A.A.R. 2319, effective May 9, 2002 (Supp. 02-2). Former Section R4-16-202 recodified to R4-16-302; New Section R4-16-202 recodified from R4-16-107 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). Amended by final exempt
B. In addition to the application form required under subsection A, an applicant for a locum tenens registration to practice medicine shall submit the following:

1. Official transcript or other authentication of graduation from a school of medicine;
2. Verification of completion of postgraduate training;
3. A statement completed by the sponsoring Arizona-licensed physician giving the reason for the request for issuance of the registration;
4. Verification of ECFMG certification if the applicant graduated from an unapproved school of medicine; and
5. Verification of licensure from every state in which the applicant has ever held a medical license.

C. In addition to the application form required under subsection (A), an applicant for a locum tenens registration to practice medicine shall submit the following:

1. Documentation listed under A.R.S. § 41-1080(A) showing that the applicant’s presence in the U.S. is authorized under federal law;
2. A full set of fingerprints and the charge specified in R4-16-205;
3. A copy of a government-issued photo identification; and
4. The fee specified under R4-16-205.

Historical Note

R4-16-203. Application for Locum Tenens Registration

A. As specifically authorized under A.R.S. § 32-1436(A), the Board establishes the following charges for the services listed:

1. Application for a license through endorsement, USMLE Step 3, or Endorsement with SPX Examination, $500;
2. Issuance of an initial license, $500, prorated from date of issuance to date of license renewal;
3. Renewal of license for two years, $500;
4. Application to reactivate an inactive license, $500;
5. Annual registration of an approved internship, residency, clinical fellowship program, or short-term residency program, $50;
6. Annual teaching license at an approved school of medicine or at an approved hospital internship, residency, or clinical fellowship program, $250;
7. Five-day teaching permit at an approved school of medicine or at an approved hospital internship, residency, or clinical fellowship program, $100;
8. Initial registration to dispense drugs and devices, $200;
9. Annual renewal to dispense drugs and devices, $150;
10. Penalty fee for late renewal of an active license, $350; and
11. Application for temporary license, $250.

B. As specifically authorized under A.R.S. § 32-1436(B), the Board establishes the following charges for the services listed:

1. Processing fingerprints to conduct a criminal background check, $50;
2. Providing a duplicate license, $50;
3. Verifying a license, $10 per request;
4. Providing a copy of records, documents, letters, minutes, applications, and files, $1 for the first three pages and 25¢ for each additional page;
5. Providing a copy of annual allopathic medical directory, $30; and
6. Providing an electronic medium containing public information about licensed physicians, $100.

R4-16-204. Repealed

Historical Note

R4-16-204. Repealed by final exempt rulemaking at 21 A.A.R. 2678, effective October 15, 2015 (Supp. 05-1).

A. As specifically authorized under A.R.S. § 32-1436(A), the Board establishes and shall collect the following fees, which are nonrefundable unless A.R.S. § 41-1077 applies:

1. Application for a license through endorsement, USMLE Step 3, or Endorsement with SPX Examination, $500;
2. Issuance of an initial license, $500, prorated from date of issuance to date of license renewal;
3. Renewal of license for two years, $500;
4. Application to reactivate an inactive license, $500;
5. Locum tenens registration, $350;
B. For each type of license, permit, or registration issued by the Board, the administrative completeness review time frame under A.R.S. § 41-1072(1) is shown on Table 1 and begins on the date the Board receives an application and all required documentation and information.

1. If the required application is not administratively complete, the Board shall send a written deficiency notice to the applicant.
   a. In the deficiency notice, the Board shall state each deficiency and the information required to complete the application or supporting documentation required to complete the application. In the deficiency notice, the Board shall include a written notice that the application is withdrawn if the applicant does not submit the additional required information or documentation within the time provided for response.
   b. Within the time provided in Table 1 for response to a deficiency notice, the applicant shall submit to the Board the documentation or information specified in the notice. The time frame for the Board to finish the administrative completeness review is suspended from the date of the notice until the date the Board receives the documentation or information from the applicant.

2. Within 30 days after receipt of a deficiency notice, an applicant who disagrees with the deficiency notice may submit to the Board a written request for a hearing regarding the deficiency notice.

3. The Board shall schedule and conduct the applicant’s deficiency hearing according to provisions prescribed under A.R.S. § 32-1427(E).

4. In addition to hearing provisions prescribed under subsection (B)(3), the Board shall send the following to the applicant in writing:
   a. A notice of the scheduled hearing at least 21 days before the hearing date; and
   b. The Board’s decision within 30 days after the hearing and notice of any applicable right of appeal.

C. For each type of license, permit, or registration issued by the Board, the substantive review time frame under A.R.S. § 41-1072(3) is shown on Table 1.

1. The Board may request make a comprehensive written request for additional information from an applicant according to provisions prescribed under A.R.S. § 41-1075 during the substantive review time frame. In any request for additional information, the Board shall include a written notice that the application is withdrawn if the applicant does not submit the additional information within the time provided for response.

2. In response to a single comprehensive written request from the Board under A.R.S. § 41-1075(A), the applicant shall submit the information identified to the Board within the time to respond specified in Table 1. The time frame for the Board to finish the substantive review is suspended from the date the Board sends the comprehensive written request for additional information until the date the Board receives the additional information from the applicant.

3. If the Board determines the applicant does not meet all substantive criteria for a license, permit, or registration as required under A.R.S. Title 32, Chapter 13 or this Chapter, the Board shall send written notice of denial to the applicant. The Board shall include notice of any applicable right of appeal in the denial notice.

4. If the applicant meets all substantive criteria for a license, permit, or registration required under A.R.S. Title 32, Chapter 13 and this Chapter, the Board shall issue the applicable license, permit, or registration to the applicant.

D. An applicant may receive a 30-day extension of the time provided under subsection (B)(1) or (C)(2) by providing written notice to the Board’s Executive Director before the time expires.

E. If a licensee does not apply for license renewal according to the biennial renewal requirement, the licensee’s license expires according to provisions prescribed under A.R.S. § 32-1430(A) unless the licensee is under investigation according to provisions under A.R.S. § 32-3202. If a licensee makes timely application according to the biennial renewal requirement but fails to respond timely to a deficiency notice under subsection (B)(1) or a request for additional information under subsection (C)(2) and fails to request from the Executive Director an extension of time to respond, the licensee’s license expires according to provisions prescribed under A.R.S. § 32-1430(A).

### Historical Note
New Section recodified from R4-16-104 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). Amended by final rulemaking at 11 A.A.R. 2944, effective September 10, 2005 (Supp. 05-3). Amended by final exempt rulemaking at 21 A.A.R. 2678, effective October 15, 2015 (Supp. 15-4).

### R4-16-207. Repealed

### Historical Note
New Section recodified from R4-16-105 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). Amended by final rulemaking at 11 A.A.R. 2944, effective September 10, 2005 (Supp. 05-3). Repealed by final exempt rulemaking at 21 A.A.R. 2678, effective October 15, 2015 (Supp. 15-4).

### Table 1. Time Frames

<table>
<thead>
<tr>
<th>Type of License</th>
<th>Overall Time Frame</th>
<th>Administrative Review Time Frame</th>
<th>Time to Respond to Deficiency Notice</th>
<th>Substantive Review Time Frame</th>
<th>Time to Respond to Request for Additional Information</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial License by Examination or Endorsement</td>
<td>240</td>
<td>120</td>
<td>365</td>
<td>120</td>
<td>90</td>
</tr>
<tr>
<td>Biennial License Renewal</td>
<td>90</td>
<td>45</td>
<td>60</td>
<td>45</td>
<td>60</td>
</tr>
<tr>
<td>Locum Tenens or Pro Bono Registration</td>
<td>120</td>
<td>60</td>
<td>90</td>
<td>60</td>
<td>30</td>
</tr>
<tr>
<td>Teaching License</td>
<td>40</td>
<td>20</td>
<td>30</td>
<td>20</td>
<td>30</td>
</tr>
<tr>
<td>Educational Teaching Permit</td>
<td>20</td>
<td>10</td>
<td>30</td>
<td>10</td>
<td>10</td>
</tr>
</tbody>
</table>
A physician who wishes to dispense a controlled substance as
defined in A.R.S. § 32-1901(12), a prescription-only drug as defined
in A.R.S. § 32-1901(65), or a prescription-only device as defined in A.R.S. § 32-1901(64) shall be currently licensed to practice medicine in Arizona and shall provide to the Board the following:
1. A completed registration form that includes the following information:
   a. The physician’s name, license number, and field of practice;
   b. A list of the types of drugs and devices the physician will dispense; and
c. The location or locations where the physician will dispense a controlled substance, a prescription-only drug, or a prescription-only device.
2. A copy of the physician’s current Drug Enforcement Administration Certificate of Registration for each dispensing location from which the physician will dispense a controlled substance.
3. The fees required in A.R.S. § 32-1436.
B. A physician shall renew a registration to dispense a controlled substance, a prescription-only drug, or a prescription-only device by complying with the requirements in subsection (A) on or before June 30 of each year. If a physician has made timely and complete application for the renewal of a registration, the physician may continue to dispense until the Board approves or denies the renewal application.
C. If the completed annual renewal form, all required documentation, and the fees are not received in the Board’s office on or before June 30, the physician shall not dispense any controlled substances, prescription-only drugs, or prescription-only devices until re-registered. The physician shall re-register by filing for initial registration under subsection (A) and shall not dispense a controlled substance, a prescription-only drug, or a prescription-only device until receipt of the re-registration.

**ARTICLE 3. DISPENSING OF DRUGS**

**R4-16-301. Registration and Renewal**

A. A physician who wishes to dispense a controlled substance as defined in A.R.S. § 32-1901(12), a prescription-only drug as defined in A.R.S. § 32-1901(65), or a prescription-only device as defined in A.R.S. § 32-1901(64) shall be currently licensed to practice medicine in Arizona and shall provide to the Board the following:

1. A completed registration form that includes the following information:
   a. The physician’s name, license number, and field of practice;
   b. A list of the types of drugs and devices the physician will dispense; and
c. The location or locations where the physician will dispense a controlled substance, a prescription-only drug, or a prescription-only device.
2. A copy of the physician’s current Drug Enforcement Administration Certificate of Registration for each dispensing location from which the physician will dispense a controlled substance.
3. The fees required in A.R.S. § 32-1436.

B. All controlled substances and prescription-only drugs dispensed shall be labeled with the following information:
   1. The physician’s name, address, and telephone number;
   2. The date the controlled substance and prescription-only drug is dispensed;
   3. The patient’s name;
   4. The controlled substance and prescription-only drug name, strength, and dosage, form, name of manufacturer, the quantity dispensed, directions for use, and any cautionary statement necessary for the safe and effective use of the controlled substance and prescription-only drug; and
   5. A beyond-use-date not to exceed one year from the date of dispensing or the manufacturer’s expiration date if less than one year.

C. A physician shall secure all controlled substances in a locked cabinet or room and shall control access to the cabinet or room by a written procedure that includes, at a minimum, designation of the persons who have access to the cabinet or room and procedures for recording requests for access to the cabinet or room. This written procedure shall be made available on demand to the Board or its authorized representatives for inspection or copying. Prescription-only drugs shall be stored so as not to be accessible to patients.

D. Controlled substances and prescription-only drugs not requiring refrigeration shall be maintained in an area where the temperature does not exceed 85° F.

E. A physician shall maintain an ongoing dispensing log for all controlled substances and the prescription-only drug nalbuphine hydrochloride (Nubain) dispensed by the physician. The dispensing log shall include the following:
   1. A separate inventory sheet for each controlled substance and prescription-only drug;
   2. The date the drug is dispensed;
   3. The patient’s name;
   4. The dosage, controlled substance and prescription-only drug name, strength, dosage, form, and name of the manufacturer;
   5. The number of dosage units dispensed;
   6. A running total of each controlled substance and prescription-only drug dispensed; and
   7. The signature of the physician written next to each entry.

F. A physician may use a computer to maintain the dispensing log required in subsection (E) if the log is quickly accessible through either on-screen viewing or printing of a copy.

G. This Section does not apply to a prepackaged manufacturer sample of a controlled substance and prescription-only drug, unless otherwise provided by federal law.

**Historical Note**

New Section adopted by final rulemaking at 6 A.A.R. 751, effective February 2, 2000 (Supp. 00-1). Former Section R4-16-301 recodified from R4-16-401; New Section R4-16-301 recodified from R4-16-201 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1). Amended by final rulemaking at 11 A.A.R. 2678, effective October 15, 2015 (Supp. 15-4).
R4-16-302 recodified from R4-16-202 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-303. Prescribing and Dispensing Requirements

A. A physician shall record on the patient’s medical record the name, strength, dosage, and form, of the controlled substance, prescription-only drug, or prescription-only device dispensed, the quantity or volume dispensed, the date the controlled substance, prescription-only drug, or prescription-only device is dispensed, the medical reasons for dispensing the controlled substance, prescription-only drug, or prescription-only device, and the number of refills authorized.

B. Before dispensing a controlled substance, prescription-only drug, or prescription-only device to a patient, a physician shall review the prepared controlled substance, prescription-only drug, or prescription-only device to ensure that:
   1. The container label and contents comply with the prescription, and
   2. The patient is informed of the name of the controlled substance, prescription-only drug, or prescription-only device, directions for use, precautions, and storage requirements.

C. A physician shall purchase all dispensed controlled substances, prescription-only drugs, or prescription-only devices from a manufacturer or distributor approved by the United States Food and Drug Administration, or a pharmacy holding a current permit from the Arizona Board of Pharmacy.

D. The person who prepares a controlled substance, prescription-only drug, or prescription-only device for dispensing shall countersign and date the original prescription form for the controlled substance, prescription-only drug, or prescription-only device.

E. For purposes of this Article, “dispensing” means the delivery of a controlled substance, a prescription-only drug, or a prescription-only device to a patient for use outside the physician’s office.

Historical Note
New Section adopted by final rulemaking at 6 A.A.R. 751, effective February 2, 2000 (Supp. 00-1). Amended by final rulemaking at 6 A.A.R. 4585, effective November 14, 2000 (Supp. 00-4). Former Section R4-16-303 recodified to R4-16-403; New Section R4-16-303 recodified from R4-16-203 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-304. Recordkeeping and Reporting Shortages

A. A physician who dispenses a controlled substance or prescription-only drug shall ensure that an original prescription dispensed from the physician’s office is dated, consecutively numbered in the order in which it is originally dispensed, and filed separately from patient medical records. A physician shall ensure that an original prescription be maintained in three separate files, as follows:
   1. Schedule II controlled substances;
   2. Schedule III, IV, and V controlled substances; and
   3. Prescription-only drugs.

B. A physician shall ensure that purchase orders and invoices are maintained for all controlled substances and prescription-only drugs dispensed for profit and not for profit for three years from the date of the purchase order or invoice. Purchase orders and invoices shall be maintained in three separate files as follows:
   1. Schedule II controlled substances only;
   2. Schedule III, IV, and V controlled substances and nalbuphine; and
   3. All other prescription-only drugs.

C. A physician who discovers a theft or loss of a controlled substance or a dangerous drug, as defined in A.R.S. § 13-3401, from the physician’s office shall:
   1. Immediately notify the local law enforcement agency,
   2. Provide that agency with a written report, and
   3. Send a copy to the Drug Enforcement Administration and the Board within seven days of the discovery.

D. For purposes of this Section, controlled substances are identified, defined, or listed in A.R.S. Title 36, Chapter 27.

Historical Note
New Section recodified from R4-16-204 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-305. Inspections; Denial and Revocation

A. A physician shall cooperate with and allow access to the physician’s office and records for periodic inspection of dispensing practices by the Board or its authorized representative. Failure to cooperate or allow access shall be grounds for revocation of a physician’s registration to dispense a controlled substance, prescription-only drug, or prescription-only device or denial of renewal of the physician’s dispensing registration.

B. Failure to comply with A.R.S. § 32-1491 or this Article constitutes grounds for denial or revocation of dispensing registration.

C. The Board shall revoke a physician’s registration to dispense a controlled substance, prescription-only drug, or prescription-only device upon occurrence of the following:
   1. Suspending, revoking, surrendering, or canceling the physician’s license;
   2. Placing the physician’s license on inactive status;
   3. Failing to timely renew the physician’s license; or
   4. Restricting the physician’s ability to prescribe or administer medication, including loss or expiration of the physician’s Drug Enforcement Administration Certificate of Registration.

D. If the Board denies a physician’s dispensing registration, the physician may appeal the decision by filing a request, in writing, with the Board, no later than 30 days after receipt of the notice denying the registration.

Historical Note
New Section recodified from R4-16-205 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

ARTICLE 4. MEDICAL ASSISTANTS

R4-16-401. Medical Assistant Training Requirements

A. A supervising physician or physician assistant shall ensure that a medical assistant satisfies one of the following training requirements before employing the medical assistant:
   1. Completion of an approved medical assistant training program; or
   2. Completion of an unapproved medical assistant training program and passage of the medical assistant examination administered by either the American Association of Medical Assistants or the American Medical Technologists.

B. This Section does not apply to any person who:
   1. Before February 2, 2000:
      a. Completed an unapproved medical assistant training program and was employed as a medical assistant after program completion; or
      b. Was directly supervised by the same physician, physician group, or physician assistant for a minimum of 2000 hours; or
   2. Completes a United States Armed Forces medical services training program.
R4-16-402. Authorized Procedures for Medical Assistants

A. A medical assistant may perform, under the direct supervision of a physician or a physician assistant, the medical procedures listed in the 2003 revised edition, Commission on Accreditation of Allied Health Education Program's, “Standards and Guidelines for an Accredited Educational Program for the Medical Assistant, Section (III)(C)(3)(a) through (III)(C)(3)(c).” This material is incorporated by reference, does not include any later amendments or editions of the incorporated matter, and may be obtained from the publisher at 35 East Wacker Drive, Suite 1970, Chicago, Illinois 60601, www.caahep.org, or the Arizona Medical Board at 9545 E. Doubletree Ranch Road, Scottsdale, AZ 85258, www.azmd.gov.

B. In addition to the medical procedures in subsection (A), a medical assistant may administer the following under the direct supervision of a physician or physician assistant:
   1. Whirlpool treatments,
   2. Diathermy treatments,
   3. Electronic galvation stimulation treatments,
   4. Ultrasound therapy,
   5. Massage therapy,
   6. Traction treatments,
   7. Transcutaneous Nerve Stimulation unit treatments,
   8. Hot and cold pack treatments, and
   9. Small volume nebulizer treatments.

R4-16-403. Renumbered

R4-16-404. Recodified

R4-16-405. Recodified

R4-16-406. Recodified

R4-16-407. Recodified

R4-16-408. Recodified

R4-16-409. Recodified

R4-16-410. Recodified

ARTICLE 5. EXECUTIVE DIRECTOR DUTIES

R4-16-501. Interim Evaluation and Investigational Interview

A. The executive director may require a physician, who is under investigation by the Board, to submit to a mental, physical, oral, or written medical competency examination after the following:
   1. Reviewing the allegations and investigator’s summary of findings; and
   2. Consulting with and receiving the agreement of the Board’s supervising medical consultant or designee that an examination is necessary.

B. The executive director may request a physician to attend an investigational interview to answer questions regarding a complaint against the physician. Before issuing a request for an investigational interview, the executive director shall review the allegations and facts to determine whether an interview is necessary to provide information the Board needs to adjudicate the case. The executive director shall consult with and receive the agreement of either the investigation supervisor or
C. The executive director shall report to the Board at each regularly scheduled Board meeting, a summary of the number and type of evaluations ordered and completed since the preceding Board meeting.

**Historical Note**
New Section made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2). Former Section R4-16-501 recodified from R4-16-601; New Section R4-16-501 recodified from R4-16-401 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-502. Direct Referral to Formal Interview
The executive director shall refer a case to a formal interview on a future Board meeting agenda, if the medical consultant in cases involving quality of care, the investigative staff, and the lead Board member concur after review of the case that a formal interview is appropriate.

**Historical Note**
New Section made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2). Former Section R4-16-502 recodified from R4-16-602; New Section R4-16-502 recodified from R4-16-402 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

Editor's Note: At the time of publication, A.R.S. § 32-1401(26) (referenced in R4-16-503) was A.R.S. § 32-1401(24). Laws 2003, Ch. 59, § 1, effective 90 days after the close of the First Regular Session of the Forty-sixth Legislature, will change the subparagraph citation to A.R.S. § 32-1401(26) (Supp. 03-2). This Section was subsequently recodified to a different Section in this Chapter. Refer to the historical notes for more information (05-1).

R4-16-503. Request for Inactive Status and License Cancellation
A. If a physician requests inactive status or license cancellation and meets the requirements of A.R.S. §§ 32-1431 and 32-1433, and is not participating in the program defined under A.R.S. § 32-1452, the executive director shall grant the request.
B. The executive director shall provide to the Board at each regularly scheduled Board meeting a list of the individuals granted inactive or cancelled license status since the preceding Board meeting.

**Historical Note**
New Section made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2). Former Section R4-16-503 recodified from R4-16-603; New Section R4-16-503 recodified from R4-16-403 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-504. Interim Consent Agreement
The executive director may enter into an interim consent agreement with a physician if there is evidence that a restriction is needed to mitigate imminent danger to the public health and safety and the investigative staff, the medical consultant, and the lead Board member concur after review of the case that a consent agreement is appropriate.

**Historical Note**
New Section made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2). Former Section R4-16-504 recodified from R4-16-605; New Section R4-16-504 recodified from R4-16-404 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-505. Mediated Case
A. The executive director shall close a case resolved through mediation.
B. The executive director shall provide to the Board at each regularly scheduled Board meeting a list of the physicians whose cases are resolved through mediation since the preceding Board meeting.

**Historical Note**
New Section made by exempt rulemaking at 9 A.A.R. 2274, effective August 12, 2003 (Supp. 03-2). Former Section R4-16-505 recodified from R4-16-606; New Section R4-16-505 recodified from R4-16-405 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-506. Referral to Formal Hearing
A. The executive director may directly refer a case to a formal hearing if the investigative staff, the medical consultant, and the lead Board member concur after review of the physician’s case that a formal hearing is appropriate.
B. The executive director shall provide to the Board at each regularly scheduled Board meeting a list of the physicians whose cases were referred to formal hearing since the preceding Board meeting and whether the referral is for revocation, suspension or is a result of an out-of-state disciplinary action, or is due to complexity of the case.

**Historical Note**
New Section R4-16-506 recodified from R4-16-406 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-507. Dismissal of Complaint
A. The executive director, with the concurrence of the investigative staff, shall dismiss a complaint if the review shows the complaint is without merit and dismissal is appropriate.
B. The executive director shall provide to the Board at each regularly scheduled Board meeting a list of the physicians about whom complaints were dismissed since the preceding Board meeting.

**Historical Note**
New Section R4-16-507 recodified from R4-16-407 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-508. Denial of License
A. The executive director shall deny a license to an applicant who does not meet statutory requirements for licensure if the executive director, in consultation with the investigative staff and the medical consultant concur after reviewing the application, that the applicant does not meet the statutory requirements.
B. The executive director shall provide to the Board at each regularly scheduled Board meeting a list of the physicians whose applications were denied since the preceding Board meeting.

**Historical Note**
New Section R4-16-508 recodified from R4-16-408 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

R4-16-509. Non-disciplinary Consent Agreement
The executive director may enter into a consent agreement under A.R.S. § 32-1451(F) with a physician to limit the physician’s practice or rehabilitate the physician if there is evidence that a licensee is mentally or physically unable to safely engage in the practice of medicine and the investigative staff, the medical consultant, and the lead Board member concur after review of the case that a consent agreement is appropriate.
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**R4-16-510. Appealing Executive Director Actions**

A. Any person aggrieved by an action taken by the executive director may appeal that action to the Board. The aggrieved person shall file a written request to the Board:
1. Thirty days after notification of the action, if personally served; or
2. Thirty-five days after the date on the notification, if mailed.

B. The aggrieved person shall provide, in the written request, evidence showing:
1. An irregularity in the investigative process or the executive director’s review deprived the party of a fair decision; or
2. Misconduct by Board staff, a Board consultant, or the executive director that deprived the party of a fair decision; or
3. Material evidence newly discovered that could have a bearing on the decision and that, with reasonable diligence, could not have been discovered and produced earlier.

C. The fact that the aggrieved party does not agree with the final decision is not grounds for a review by the Board.

D. If an aggrieved person fails to submit a written request within the time specified in subsection (A), the Board is relieved of the requirement to review actions taken by the executive director. The executive director may, however, evaluate newly provided information that is material or substantial in content to determine whether the Board should review the case.

E. If a written request is submitted that meets the requirements of subsection (B):
1. The Board shall consider the written request at its next regularly scheduled meeting.
2. If the written request provides new material or substantial evidence that requires additional investigation, the investigation shall be conducted as expeditiously as possible and the case shall be forwarded to the Board at the first possible regularly scheduled meeting.

**Historical Note**
New Section R4-16-509 recodified from R4-16-409 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

**R4-16-604. Aggravating Factors Considered in Disciplinary Actions**

When determining the degree of discipline, the Board may consider certain factors including, but not limited to, the following:
1. Prior disciplinary offenses;
2. Dishonest or selfish motive;
3. Pattern of misconduct; multiple offenses;
4. Bad faith obstruction of the disciplinary proceeding by intentionally failing to comply with rules or orders of the Board;
5. Submission of false evidence, false statements, or other deceptive practices during the investigative or disciplinary process;
6. Refusal to acknowledge wrongful nature of conduct; and
7. Vulnerability of the victim.

**Historical Note**
New Section R4-16-604 recodified from R4-16-504 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

**R4-16-605. Mitigating Factors Considered in Disciplinary Actions**

When determining the degree of discipline, the Board may consider certain factors including, but not limited to, the following:
1. Absence of prior disciplinary record;
2. Absence of dishonest or selfish motive;
3. Timely good faith effort to rectify consequences of misconduct;
4. Interim rehabilitation;
5. Remoteness of prior offenses; and
6. How much control the physician has of processes in the specific practice setting.

**Historical Note**
New Section R4-16-605 recodified from R4-16-504 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

**ARTICLE 6. DISCIPLINARY ACTIONS**

**R4-16-601. Expired**

**Historical Note**
New Section R4-16-601 recodified from R4-16-501 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

**R4-16-602. Expired**

**Historical Note**
New Section R4-16-602 recodified from R4-16-502 at 11 A.A.R. 1283, effective March 25, 2005 (Supp. 05-1).

**Editor’s Note:** To conform with the renumbering in A.R.S., the Arizona Medical Board requested (under A.R.S. § 41-1011 et seq.) a subsection reference update in R4-16-603 [R05-85]. Please refer to the historical notes for more details (Supp. 05-1).

**R4-16-603. Expired**

**ARTICLE 7. OFFICE-BASED SURGERY USING SEDATION**

**R4-16-701. Health Care Institution License**

A physician who uses general anesthesia in the physician’s office or other outpatient setting that is not part of a licensed hospital or licensed ambulatory surgical center when performing office-based surgery using sedation shall obtain a health care institution license as required by the Arizona Department of Health Services under A.R.S. Title 36, Chapter 4 and 9 A.A.C. 10.

**Historical Note**
New Section made by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).

**R4-16-702. Administrative Provisions**

A. A physician who performs office-based surgery using sedation in the physician’s office or other outpatient setting that is not part of a licensed hospital or licensed ambulatory surgical center shall:
1. Establish, document, and implement written policies and procedures that cover:
A physician shall ensure that each office-based surgery using sedation:

1. Has sufficient education, training, and experience to perform duties assigned;
2. If applicable, has a current license or certification to perform duties assigned; and
3. Performs only those acts that are within the scope of practice established in the staff member’s or health care professional’s governing statutes;

2. Ensure that the office where the office-based surgery using sedation is performed has all equipment necessary:
   a. For the physician to safely perform the office-based surgery using sedation,
   b. For the physician or health care professional to safely administer the sedation,
   c. For the physician or health care professional to monitor the use of sedation, and
   d. For the physician and health care professional administering the sedation to rescue a patient after the sedation is administered to the patient and the patient enters into a deeper state of sedation than what was intended by the physician.

4. Ensure that a copy of the patient’s rights policy is provided to each patient before performing office-based surgery using sedation;

5. Obtain informed consent from the patient before performing an office-based surgery using sedation that:
   a. Authorizes the office-based surgery, and
   b. Authorizes the office-based surgery to be performed in the physician’s office; and

6. Review all policies and procedures every 12 months and update as needed.

B. A physician who performs office-based surgery using sedation shall comply with:

1. The local jurisdiction’s fire code;
2. The local jurisdiction’s building codes for construction and occupancy;
3. The biohazardous waste and hazardous waste standards in 18 A.A.C. 13, Article 14; and
4. The controlled drug administration, supply, and storage standards in 4 A.A.C. 23.

HISTORICAL NOTE
New Section made by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).

R4-16-704. Sedation Monitoring Standards
A physician who performs office-based surgery using sedation shall ensure from the time sedation is administered until post-sedation monitoring begins:

1. A quantitative method of assessing a patient’s oxygenation, such as pulse oximetry, is used when minimal sedation is administered to the patient, and
2. When moderate or deep sedation is administered to a patient:
   a. A quantitative method of assessing the patient’s oxygenation, such as pulse oximetry, is used;
   b. The patient’s ventilatory function is monitored by any of the following:
      i. Direct observation,
      ii. Auscultation, or
      iii. Capnography;
   c. The patient’s circulatory function is monitored during the surgery by:
      i. Having a continuously displayed electrocardiogram,
      ii. Documenting arterial blood pressure and heart rate at least every five minutes, and
      iii. Evaluating the patient’s cardiovascular function by pulse plethysmography;
   d. The patient’s temperature is monitored if the physician expects the patient’s temperature to fluctuate; and
   e. That a licensed and qualified healthcare professional, other than the physician performing the office-based surgery, whose sole responsibility is attending to the patient, is present throughout the office-based surgery.

HISTORICAL NOTE
New Section made by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).

R4-16-705. Perioperative Period; Patient Discharge
A physician performing office-based surgery using sedation shall ensure all of the following:

1. Has a medical condition or other condition that indicates the procedure should not be performed in the physician’s office, or
2. Will require inpatient services at a hospital.

HISTORICAL NOTE
New Section made by final rulemaking at 14 A.A.R. 380, effective January 8, 2008 (Supp. 08-1).
a. The time and date of the patient’s discharge, and
b. A description of the patient’s medical condition at
   the time of discharge; and

6. A patient receives discharge instructions and documents
   in the patient’s medical record that the patient received
   the discharge instructions.

**Historical Note**

New Section made by final rulemaking at 14 A.A.R. 380,
   effective January 8, 2008 (Supp. 08-1).

R4-16-706. Emergency Drugs; Equipment and Space Used
   for Office-Based Surgery Using Sedation

A. In addition to the requirements in R4-16-702(A)(3) and R4-
   16-703(A)(1), a physician who performs office-based surgery
   using sedation shall ensure that the physician’s office has at a
   minimum:

1. The following:
   a. A reliable oxygen source with a SaO₂ monitor;
   b. Suction;
   c. Resuscitation equipment, including a defibrillator;
   d. Emergency drugs; and
   e. A cardiac monitor;

2. The equipment for patient monitoring according to the
   standards in R4-16-704;

3. Space large enough to:
   a. Allow for access to the patient during office-based
      surgery using sedation, recovery, and any emer-
      gency;
   b. Accommodate all equipment necessary to perform
      the office-based surgery using sedation; and
   c. Accommodate all equipment necessary for sedation
      monitoring;

4. A source of auxiliary electrical power available in the
   event of a power failure; and

5. Equipment, emergency drugs, and resuscitative capabili-
   ties required under this Section for patients less than 18
   years of age, if office-based surgery using sedation is per-
   formed on these patients; and

6. Procedures to minimize the spread of infection.

B. A physician who performs office-based surgery using sedation
   shall:

1. Ensure that all equipment used for office-based surgery
   using sedation is maintained, tested, and inspected
   according to manufacturer specifications, and

2. Maintain documentation of manufacturer-recommended
   maintenance of all equipment used in office-based sur-
   gery using sedation.

**Historical Note**

New Section made by final rulemaking at 14 A.A.R. 380,
   effective January 8, 2008 (Supp. 08-1).


A. A physician who performs office-based surgery using sedation
   shall ensure that before a heath care professional participates
   in or staff member assists with office-based surgery using
   sedation, the health care professional and staff member receive
   instruction in the following:

1. Policy and procedure in cases of emergency,
2. Policy and procedure for office evacuation, and
3. Safe and timely patient transfer.

B. When performing office-based surgery using sedation, a phy-
   sician shall not use any drug or agent that trigger malignant
   hyperthermia.

**Historical Note**

New Section made by final rulemaking at 14 A.A.R. 380,
   effective January 8, 2008 (Supp. 08-1).