

APA-3

**CERTIFICATION OF ADMINISTRATIVE RULES  
FILED WITH THE LEGISLATIVE SERVICES AGENCY  
OTHNI LATHRAM, DIRECTOR**

(Pursuant to Code of Alabama 1975, §41-22-6, as amended).

I certify that the attached is/are correct copy/copies of rule/s as promulgated and adopted on Friday, January 9, 2026, and filed with the agency secretary on Wednesday, January 21, 2026.

**AGENCY NAME:** Board of Dental Examiners of Alabama

**INTENDED ACTION:** Amend

**RULE NO.:** 270-X-2-.17

(If amended rule, give specific paragraph, subparagraphs, etc., being amended)

**RULE TITLE:** Criteria For On-Site Inspection For The Use Of  
General Anesthesia And Parenteral Sedation

**ACTION TAKEN:** State whether the rule was adopted with or without changes from the proposal due to written or oral comments:

**Adopted with changes.** no comments received

NOTICE OF INTENDED ACTION PUBLISHED IN VOLUME XLIV, ISSUE NO. 1, AAM,  
DATED FRIDAY, OCTOBER 31, 2025.

**STATUTORY RULEMAKING AUTHORITY:** 34-9-1-90

(Date Filed)  
(For LRS Use Only)

**REC'D & FILED**

JAN 21, 2026

**LEGISLATIVE SVC AGENCY**

Sharon Thomas

Sharon Thomas

Certifying Officer or his or her  
Deputy

(NOTE: In accordance with §41-22-6(b), as amended, a proposed rule is required to be certified within 90 days after completion of the notice.)

Criteria For On-Site Inspection For The Use Of  
General Anesthesia And Parenteral Sedation/  
Moderate Sedation.

(1) This rule contains the procedures, criteria, and information mandatory for the clinical administration of general anesthesia and parenteral/~~moderate~~ sedation. These shall be used by the inspecting team in determining whether a dentist's facilities, equipment, and personnel have satisfied the requirements imposed by law and this rule for the issuance of a general anesthesia or a parenteral/~~moderate~~ sedation permit.

(a) Clinical Use of General Anesthesia and/or Parenteral/~~Moderate~~ Sedation. Two (2) procedures utilizing general anesthesia and/or parenteral/~~moderate~~ sedation shall be observed. This portion of the evaluation shall not exceed two (2) hours. No evaluation can be considered complete unless this part is included.

(b) Simulated Emergencies.

1. The examiners will simulate the emergency situations/techniques listed below. The permittee and office staff should be competent in managing all of these:

- (i) Laryngospasm
- (ii) Bronchospasm
- (iii) Emesis and aspiration of vomitus
- (iv) Management of foreign bodies in the airway
- (v) Angina pectoris
- (vi) Myocardial infarction
- (vii) Cardiopulmonary resuscitation
- (viii) Hypotension
- (ix) Hypertensive crisis
- (x) Acute allergic reaction
- (xi) Hyperventilation syndrome
- (xii) Convulsion of unknown etiology
- (xiii) Syncope

(IV) A record of the anesthesia and/or sedation shall be made. The anesthetic and other agents and amounts given shall be indicated. Preoperative, intraoperative, and post-operative vital signs shall be recorded and any unusual reactions or complications shall be documented. Starting and ending times for anesthesia shall be recorded. The persons present during the procedure shall be noted.

(V) A record of prescriptions given shall be included.

(e) Office Facilities and Equipment. This section deals with the physical requirements for conducting office anesthesia/sedation.

1. The fundamental physical requirements for the anesthesia and/or sedation facility are:

(i) The Operating Room/Operatory - The operatory shall be large enough to adequately accommodate the patient on a table or in a dental chair and permit the anesthesia and/or sedation team, consisting of the dentist and two or three trained assistants, to move freely about the patient.

(ii) The Operating Table or Dental Chair - The table or chair shall permit the patient to be positioned so the anesthesia team can maintain the airway, allows quick alteration of patient position in an emergency, provides a firm platform for the management of cardiopulmonary resuscitation, and provides easy access to the patient's oral cavity.

(iii) Lighting Systems.

(I) Room lighting shall be adequate to permit evaluation of the patient's skin and mucosal color.

(II) There shall be provisions for auxiliary lighting should the power fail in the operatory. Backup lighting shall be battery powered and of sufficient intensity to permit completion of any procedure underway at the time of general power failure.

(iv) Suction Equipment.

(I) Aspiration shall be provided either by a portable suction unit or by a central suction installation. It is important to provide for auxiliary suction should the pump or electrical power fail. '

(II) Multiple suction tips, including tonsil suction tips, shall be available.

(v) Oxygen and Supplemental Gas-Delivery System.

(I) An oxygen and supplemental gas-delivery system capable of delivering metered oxygen and/or gas under positive pressure shall be required.

(II) Gas outlets for remote delivery systems shall be coded to prevent accidental administration of the wrong gas. Fail-safe mechanisms on anesthetic machines are mandatory.

(vi) Patient Recovery.

(I) Patients shall be retained in the surgery area until all protective reflexes have fully returned unless the dental staff is in immediate attendance at all times in the recovery area to continue vital-sign and airway observations.

(II) A patient recovering from a general anesthetic or sedation procedure shall be monitored in the recovery area. This recovery area shall include sufficient room to treat any emergency situation. The recovery area shall be equipped to provide oxygen under positive pressure and have adequate lighting, access to suction, and electrical outlets for connecting cardiac monitoring and defibrillating equipment.

(vii) Drug and Instrument Preparation and Storage Area-An adequate outpatient facility shall contain an area conducive to the sterile preparation and storage of drugs used in anesthesia and/or sedation. There should be provisions for refrigeration to store certain drugs. The drug preparation area shall include a secure storage site for narcotics and other dangerous drugs.

(viii) Gas Storage Area.

(I) Permissible Categories

I. Gas may be stored in a central location used by one or multiple practitioners within the same building.

II. Gas may be stored in the individual operatory.

III. There shall be reserve tanks of gas not connected for immediate use.

(II) Requirements

I. All gas storage shall be maintained according to local building, fire, and safety codes.

II. Gas stored in a central location shall have a central low-pressure alarm, which shall be easily heard in the treatment area where the procedure is being performed. In lieu of a central alarm, a daily gas log may be maintained and checked by the dentist.

(f) Monitoring.

1. The various methods and physiologic parameters used in monitoring patients shall be designed to immediately detect the changes produced either by dental stimulation or the anesthetics or sedatives employed.

2. All patients shall be monitored when anesthetics and/or sedation are employed. The doctor shall continually observe the patient's status and make moment-to-moment assessments of the patient's condition so necessary adjustments may be made.

3. Mechanical monitoring shall be used with every patient. Blood pressure, cardiac rate, oxygen saturation, end tidal CO<sub>2</sub> (capnography), and electrocardiogram (EKG) all are required to be monitored during the pre-anesthetic and/or pre-sedation and intra- and post-operative intervals.

4. All equipment, both manual and automated, required by this rule for the monitoring of respiration, heart rate, and blood pressure under parts (g), (h), and (i) below must be maintained in each facility location where a dentist possesses a permit to use general anesthesia and/

or parenteral/~~moderate~~ sedation, and shall not be shared or transported between multiple facility locations. Documentation of compliance with manufacturers' recommended maintenance of monitors, anesthesia delivery systems, and other anesthesia-related equipment shall be maintained.

(g) Required Monitoring of Respiration.

1. Oximetry. Oximeter using a peripheral (finger, ear, or toe) transmitted wave-form monitor.

2. Capnography. Capnograph to monitor and measure the concentration or partial pressure of carbon dioxide in the respiratory gases, as well as the competency of the airway for gas exchange. The dentist shall monitor ventilation and/or breathing by monitoring end tidal carbon dioxide unless precluded or invalidated by the nature of the patient, procedure, or equipment.

(h) Required Monitoring of Heart Rate.

1. Pre-cordial stethoscope or pulse oximeter.

2. Electrocardiogram (EKG).

(i) Required Monitoring of Blood Pressure. Systolic pressure, diastolic pressure, and heart rate must be recorded.

1. Sphygmomanometer and stethoscope or an automatic equivalent.

(j) Emergency Drugs

1. At a minimum each facility in which general anesthesia and/or parenteral/~~moderate~~ sedation is used by a dentist, must maintain a secure stock of current emergency drugs from each of the following categories. Any specific drug selected must satisfy current advanced cardiac life support (ACLS) and/or American Association of Oral and Maxillofacial Surgeons (AAOMS) standards:

(i) Anticonvulsant

(ii) Antihypoglycemic

(iii) Anticholinergic

(iv) Antiarrhythmics

- (v) Steroid
- (vi) Antihistamine
- (vii) Cardiac stimulant/antihypotensive agent
- (viii) Analgesic
- (ix) Benzodiazepine antagonist
- (x) Narcotic antagonist
- (xi) Paralytic Agent for Laryngospasms along with the paralytic agent's reversal agent, of any
- (xii) Antihypertensive
- (xiii) Nitrate

2. The aforementioned emergency drugs must be maintained at each facility in which general anesthesia and/or parenteral/~~moderate~~ sedation is used by a dentist and shall not be shared or transported between multiple facility locations.

(k) Personnel.

1. For the administration of parenteral/~~moderate~~ sedation, at least two (2) individuals, each appropriately trained, are required to be present throughout the procedure, consisting of the doctor and an assistant trained to monitor appropriate physiologic variables.

2. For the administration of general anesthesia, at least three (3) individuals, each appropriately trained, are required to be present throughout the procedure, consisting of the doctor who directs the general anesthesia, a person whose responsibilities are observation and monitoring of the patient, and a third person who assists the operating dentist.

(l) The Board shall appoint examiners for the purpose of conducting the on-site inspections of dental facilities, equipment, and personnel as prescribed in Rule 270-X-2-.24.

(m) Upon receipt of an initial application for the administration of general anesthesia or parenteral/~~moderate~~ sedation, a preliminary facility evaluation of the applicant's facility will be conducted by examiners appointed

by the Board. This preliminary facility evaluation will fully assess the safety of the facility, the presence of emergency equipment, the presence of necessary drugs, and the credentials of the individuals who will participate in the procedures. Subsequent to a satisfactory preliminary facility evaluation, a temporary permit for the administration of general anesthesia or parenteral/~~moderate~~ sedation shall be issued. This temporary permit shall be valid for no more than ninety (90) days, until a subsequent evaluation which fully examines the criteria set forth in this rule is conducted by examiners appointed by the Board.

(n) An applicant may receive a single temporary permit for the administration of intravenous anesthesia and normally may not receive a second temporary permit at another location without successfully completing a full anesthesia evaluation.

(o) A second temporary for an additional office that is maintained under the exact same practice identity, such as a satellite office, with the exact same staff may be issued and may stay in effect while the temporary permit is legally maintained pending completion of a full anesthesia evaluation.

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(p) If upon an initial application for the issuance of a permit for the administration of general anesthesia or parenteral/~~moderate~~ sedation, the primary office of a dentist(s) has received a satisfactory on-site inspection and the dentist(s) also applies for the issuance of a permit to administer general anesthesia or parenteral/~~moderate~~ sedation at a secondary office(s) or location(s), any on-site inspection thereof shall be limited only to the dental facility and equipment, provided that the same personnel satisfactorily evaluated at the primary office(s) of the dentist(s) will be engaged or involved in the administration of general anesthesia or parenteral/~~moderate~~ sedation at the said secondary office(s) or location(s). If upon a request

for renewal by a dentist(s) of a permit to administer general anesthesia or parenteral/moderate sedation at both his primary and secondary office(s) or location(s), the Board of Dental Examiners of Alabama determines that an on-site inspection of these office(s) or location(s) is required, the same procedure as outlined above in relation to the initial application for these permits shall be utilized.

(q) The examining team shall submit to the Board the report of their on-site inspection within fourteen (14) days from the date of said inspection. If the results of the initial evaluation are deemed unsatisfactory, the anesthesia certificate is immediately suspended and the applicant must reapply by submitting another application and fee to the Board.

**Author:** Board of Dental Examiners of Alabama

**Statutory Authority:** Code of Ala. 1975, §§34-9-43, 34-9-60, 34-9-63, 34-9-65.

**History:** Filed May 23, 1986. **Amended:** Filed March 8, 1988.

**Amended:** Filed December 20, 1993. **Amended:** Filed February 22, 2012; effective March 28, 2012. **Amended:** Filed January 22, 2018; effective March 8, 2018. **Amended:** Filed July 26, 2018; effective September 9, 2018. **Amended:** Published July 31, 2020; effective September 14, 2020. **Amended:** Published September 29, 2023; effective November 13, 2023. **Amended:** Published February 27, 2026; effective April 13, 2026.