WAC 246-817-760  Moderate sedation with parenteral agents. (1) Training requirements: To administer moderate sedation with parenteral agents, the dentist must have successfully completed a postdoctoral course(s) of sixty clock hours or more which includes training in basic moderate sedation, physical evaluation, venipuncture, technical administration, recognition and management of complications and emergencies, monitoring, and supervised experience in providing moderate sedation to fifteen or more patients. If treating an adult, the dentist must have training in adult sedation. If treating a minor, the dentist must have training in pediatric sedation.

(2) In addition to meeting the criteria in subsection (1) of this section, the dentist must also have a current certification in advanced cardiac life support (ACLS) or pediatric advanced life support (PALS). If treating an adult, the dentist must have ACLS certification. If treating a minor, the dentist must have PALS certification.

(3) The drugs, drug amounts, and techniques used must carry a margin of safety wide enough to render unintended loss of consciousness highly unlikely.

(4) Procedures for administration of moderate sedation with parenteral agents by a dentist and an individual trained in monitoring sedated patients:
   (a) In the treatment setting, a patient receiving moderate sedation with parenteral agents must have that sedation administered by a person qualified under this chapter.
   (b) A patient may not be left alone in a room and must be continually monitored by a dentist with a valid moderate sedation with parenteral agent permit or trained anesthesia monitor.
   (c) An intravenous infusion must be maintained during the administration of a parenteral agent. Two exceptions for intravenous infusion may occur, but reasons why intravenous infusion was not used must be documented for:
      (i) Pediatric sedation cases using agents for brief procedures; and
      (ii) When the pediatric patient is uncooperative or the emotional condition is such that intravenous access is not possible.
   (d) When the operative dentist is also the person administering the moderate sedation with parenteral agents, the operative dentist must be continuously assisted by at least one individual experienced in monitoring sedated patients. If treating an adult, the additional individual must have experience or training in adult sedation. If treating a minor, the additional individual must have experience or training in pediatric sedation.
   (e) In the treatment setting, a patient experiencing moderate sedation with parenteral agents must be visually and tactically monitored by the dentist or an individual trained in monitoring sedated patients. Patient monitoring must include:
      (i) Heart rate;
      (ii) Blood pressure;
      (iii) Respiration;
      (iv) Pulse oximetry; and
      (v) Expired carbon dioxide (CO₂). Two exceptions for expired CO₂ monitoring may occur, but reasons why expired CO₂ monitoring was not used must be documented for:
         (A) Pediatric sedation cases using agents for brief procedures; and
(B) When the pediatric patient is uncooperative or the emotional condition is such that CO₂ monitoring is not possible.

(f) Requirements of immobilization devices for pediatric patients:
   (i) Immobilization devices, such as, papoose boards, must be applied in such a way as to avoid airway obstruction or chest restriction.
   (ii) The pediatric patient head position and respiratory excursions must be checked frequently to ensure airway patency.
   (iii) If an immobilization device is used, a hand or foot must be kept exposed.
   (g) The patient's blood pressure and heart rate must be recorded every five minutes, pulse oximetry recorded every five minutes, and respiration rate must be recorded at least every fifteen minutes.
   (h) The patient's level of consciousness must be recorded prior to the dismissal of the patient.
   (i) Patients receiving moderate sedation with parenteral agents must be accompanied by a responsible adult upon departure from the treatment facility.
   (j) If a patient unintentionally enters a deeper level of sedation, the patient must be returned to a level of moderate sedation as quickly as possible. While returning the patient to the moderate level of sedation, periodic monitoring of pulse, respiration, blood pressure and continuous monitoring of oxygen saturation must be maintained. In such cases, these same parameters must be taken and recorded at appropriate intervals throughout the procedure and vital signs and level of consciousness must be recorded during the sedation and prior to dismissal of the patient.

(5) Dental records must contain appropriate medical history and patient evaluation. Sedation records must be recorded during the procedure in a timely manner and must include:
   (a) Blood pressure;
   (b) Heart rate;
   (c) Respiration;
   (d) Pulse oximetry;
   (e) End-tidal CO₂. Two exceptions for end-tidal CO₂ monitoring may occur, but reasons why end-tidal CO₂ monitoring was not used must be documented for:
      (i) Pediatric sedation cases using agents for brief procedures; and
      (ii) When the pediatric patient is uncooperative or the emotional condition is such that end-tidal CO₂ monitoring is not possible.
   (f) Drugs administered including amounts and time administered;
   (g) Length of procedure; and
   (h) Any complications of sedation.

(6) Equipment and emergency medications: All offices in which moderate sedation with parenteral agents is administered or prescribed must comply with the following equipment standards:
   Office facilities and equipment shall include:
   (a) Suction equipment capable of aspirating gastric contents from the mouth and pharynx;
   (b) Portable oxygen delivery system including full face masks and a bag-valve-mask combination with appropriate connectors capable of delivering positive pressure, oxygen-enriched patient ventilation and oral and nasal pharyngeal airways. If treating an adult, the equipment
must be appropriate for adult sedation. If treating a minor, the equipment must be appropriate for pediatric sedation;
(c) A blood pressure cuff (sphygmomanometer) of appropriate size and stethoscope; or equivalent monitoring devices;
(d) End-tidal CO₂ monitor;
(e) Pulse oximetry; and
(f) An emergency drug kit with minimum contents of:
   (i) Sterile needles, syringes, and tourniquet;
   (ii) Narcotic antagonist;
   (iii) Alpha and beta adrenergic stimulant;
   (iv) Vasopressor;
   (v) Coronary vasodilator;
   (vi) Antihistamine;
   (vii) Parasympatholytic;
   (viii) Intravenous fluids, tubing, and infusion set; and
   (ix) Sedative antagonists for drugs used, if available.
(7) Continuing education: A dentist who administers moderate sedation with parenteral agents must participate in eighteen hours of continuing education or equivalent every three years.
   (a) The education must include instruction in one or more of the following areas:
      (i) Venipuncture;
      (ii) Intravenous sedation;
      (iii) Physiology;
      (iv) Pharmacology;
      (v) Nitrous oxide analgesia;
      (vi) Patient evaluation;
      (vii) Patient monitoring; and
      (viii) Medical emergencies.
   (b) In addition to the education requirements in (a) of this subsection, the dentist must have a current certification in advanced cardiac life support (ACLS) or pediatric advanced life support (PALS) to renew the moderate sedation with parenteral agents permit. Hourly credits earned from certification in BLS, ACLS, or PALS courses may not be used to meet the education requirements in (a) of this subsection to renew a moderate sedation with parenteral agents permit. However, the hourly credits earned in ACLS or PALS certification may be used to meet the requirements of WAC 246-817-440 to renew the dentist license.
(8) A permit of authorization is required. See WAC 246-817-774 for permitting requirements.

[Statutory Authority: RCW 18.32.0365 and 18.32.640. WSR 17-07-037, § 246-817-760, filed 3/8/17, effective 4/8/17; WSR 16-06-106, § 246-817-760, filed 3/1/16, effective 4/1/16; WSR 09-04-042, § 246-817-760, filed 1/30/09, effective 3/2/09. Statutory Authority: RCW 18.32.035. WSR 95-21-041, § 246-817-760, filed 10/10/95, effective 11/10/95.]