650—29.9 (153) Reporting of adverse occurrences related to sedation, nitrous oxide inhalation analgesia, and antianxiety premedication.

29.9(1) *Reporting.* All licensed dentists in the practice of dentistry in this state must submit a report within a period of seven days to the board office of any mortality or other incident which results in temporary or permanent physical or mental injury requiring hospitalization of the patient during, or as a result of, antianxiety premedication, nitrous oxide inhalation analgesia, or sedation. The report shall include responses to at least the following:

a. Description of dental procedure.

b. Description of preoperative physical condition of patient.

c. List of drugs and dosage administered.

d. Description, in detail, of techniques utilized in administering the drugs utilized.

e. Description of adverse occurrence:

1. Description, in detail, of symptoms of any complications, to include but not be limited to onset, and type of symptoms in patient.
2. Treatment instituted on the patient.
3. Response of the patient to the treatment.

f. Description of the patient's condition on termination of any procedures undertaken.

29.9(2) *Failure to report.* Failure to comply with subrule 29.9(1), when the occurrence is related to the use of sedation, nitrous oxide inhalation analgesia, or antianxiety premedication, may result in the dentist's loss of authorization to administer sedation, nitrous oxide inhalation analgesia, or antianxiety premedication or in any other sanction provided by law.

[**ARC 8614B**, IAB 3/10/10, effective 4/14/10; **ARC 1194C**, IAB 11/27/13, effective 11/4/13]