

ADVERSE REACTION REPORT

Minnesota Rules 3100.3600 requires that you file this report for any incident that arises from the administration of nitrous oxide inhalation analgesia or of a pharmacological agent for the purpose of general anesthesia, conscious sedation, local anesthesia, analgesia, or anxiolysis that results in a serious or unusual outcome that produces a temporary or permanent physiological injury, harm, or other detrimental effect to one or more of a patient's body system(s). It is NOT necessary to report incidents such as nausea, a single episode of emesis, or mild allergic reaction. **This report and relevant records shall be submitted to the Board of Dentistry within ten days of the incident.**

LICENSEE INFORMATION

Name (please print):

License Number:

Address:

City:

State:

Zip

I. REACTION INFORMATION							
PATIENT ID/INITIALS (In Confidence)	AGE (YRS)	SEX	REACTION ONSET			<p>CHECK ALL APPROPRIATE:</p> <p><input type="checkbox"/> PATIENT DIED</p> <p><input type="checkbox"/> REACTION TREATED WITH RX DRUG</p> <p><input type="checkbox"/> RESULTED IN TREATMENT BY PHYSICIAN AND/OR HOSPITALIZATION</p> <p><input type="checkbox"/> RESULTED IN PERMANENT DISABILITY</p> <p><input type="checkbox"/> NONE OF THE ABOVE</p>	
			MO	DA	YR		
DESCRIBE REACTION(S)							
RELEVANT TESTS/LABORATORY DATA							
II. SUSPECT DRUG(S) INFORMATION							
SUSPECT DRUG(S) (Indicate manufacturer and lot # for vaccines/biologics)					DID REACTION ABATE AFTER STOPPING DRUG?		
DOSE	ROUTE OF ADMINISTRATION				<input type="checkbox"/> YES		<input type="checkbox"/> NO
					<input type="checkbox"/> N/A		

INDICATION(S) FOR USE		DID REACTION REAPPEAR AFTER REINTRODUCTION?		
DATES OF ADMINISTRATION (<i>From/To</i>)	DURATION OF ADMINISTRATION	<input type="checkbox"/> YES	<input type="checkbox"/> NO	<input type="checkbox"/> N/A
III. CONCOMITANT DRUGS AND HISTORY				
CONCOMITANT DRUGS AND DATES OF ADMINISTRATION (<i>Exclude those used to treat reaction</i>)				
OTHER RELEVANT HISTORY (<i>e.g., diagnoses, allergies, pregnancy with LMP, etc.</i>)				
IV. SIGNATURE				
SIGNED:		DATE:		

Please include a copy of the dental and/ or relevant medical records for the patient involved in the adverse reaction event. The record should include:

- All treatment notes relating to the incident (date of incident, follow up care, etc.)
- Anesthesia record (if applicable)
- Medical History
- Dental History
- Current Medications