1	21 NCAC 16Q .0103 is	adopted with change	es as published in 3	7:07 NC	R 543-45 as fol	lows:	
2							
3	21 NCAC 16Q .0103	EQUIPMENT,	PERSONNEL,	AND	CLINICAL	REQUIREMENTS	TO
4			ANESTHESIA OI				
5	(a) Before administerin	g general anesthesia	, moderate conscio	us sedati	on, or moderate	e pediatric conscious se	dation
6	("anesthesia or moderat	e sedation"), or supe	ervising a CRNA to	adminis	ster or <u>an</u> RN ei	nployed to deliver ane	sthesia
7	or moderate sedation, a	dentist shall hold an	unexpired permit is	ssued by	the Board in acc	cordance with this Subo	hapter
8	permitting the dentist to	administer that leve	l of sedation.				
9	(b) Before performing s	sedation procedures i	n a facility other the	an a hosp	oital or credentia	aled surgery center, the	permit
10	holder shall ensure that	the Board has been n	otified that the pern	nit holde	r intends to adm	inister anesthesia or mo	oderate
11	sedation at the facility a	and shall ensure that	the facility has pa	ssed a fa	cility inspection	n by the Board in acco	rdance
12	with this Subchapter. F	or purposes of these	Rules, "credentiale	ed surger	y center" means	s a surgical facility acc	<u>redited</u>
13	by the Joint Commission	n on Accreditation of	Healthcare Organi	zations, t	the Accreditatio	n Association for Ambi	ulatory
14	Health Care, or the Ame	erican Association fo	or Accreditation of	<u>Ambulat</u>	ory Surgery Fac	<u>cilities.</u>	
15	(c) The permit holder sh	nall ensure that the fac	cility where the sed	ation pro	cedure is to be p	performed meets the follower	lowing
16	requirements at the time	of the procedure:					
17	(1) The p	ermit holder shall e	ensure the facility	is equipp	ped as follows	and that the following	listed
18	equip	ment is immediately	available and acces	ssible fro	m the operatory	and recovery rooms:	
19	(A)	an operatory of s	ize and design to p	ermit ac	cess of emerge	ncy equipment and per	sonnel
20		and to permit em	ergency manageme	nt;			
21	(B)	a CPR board or	dental chair with	out enha	ncements s uital	ble for providing eme	rgency
22		treatment;					
23	(C)	lighting as necess	sary for the proced	ure to be	performed, spe	ecific procedures and b	ack-up
24		lighting;					
25	(D)	suction equipmen	nt as necessary for	the proc	edure to be per	rformed, specific proce	edures,
26		including non-ele	ectrical back-up suc	tion;			
27	(E)	positive pressure	oxygen delivery s	ystem, ii	ncluding full fa	ce masks for small, me	edium,
28		and large patient	s, and back-up E-o	cylinder	portable oxyger	n tank apart from the	central
29		system;					
30	(F)	small, medium, a	nd large oral and na	asal airw	ays;		
31	(G)	a blood pressure	monitoring device;				
32	(H)	an EKG monitor;					
33	(I)	a pulse oximeter;					
34	(J)	an automatic exte	ernal defibrillator (A	AED);			
35	(K)	a capnograph;					
36	(L)	a precordial or pr	etracheal stethosco	pe;			
37	(M)	a thermometer:					

1		(iv) vascular access set-up as necessary for the procedure to be performed, specific procedures,
2		including hardware and fluids;
3		(O) a laryngoscope with working batteries;
4		(P) intubation forceps and advanced airway devices;
5		(Q) tonsillar suction with back-up suction;
6		(R) syringes as necessary for the procedure to be performed; specific procedures; and
7		(S) tourniquet and tape.
8	(2)	The permit holder shall ensure all monitoring and other equipment in the facility receives preventive
9		maintenance no less frequently than once per year, including safety and function checks per the
10		manufacturers' recommendations. The permit holder shall maintain documentation of all preventive
11		maintenance performed, and shall ensure equipment is replaced upon its expiration or as clinically
12		required.
13	(3)	The permit holder shall ensure the following unexpired drugs are immediately available and are
14		accessible from the operatory and recovery rooms:
15		(A) epinephrine;
16		(B) atropine;
17		(C) an antiarrhythmic;
18		(D) an antihistamine;
19		(E) an antihypertensive;
20		(F) a bronchodilator;
21		(G) an antihypoglycemic agent;
22		(H) a vasopressor;
23		(I) a corticosteroid;
24		(J) an anticonvulsant;
25		(K) appropriate reversal agents;
26		(L) nitroglycerine; and
27		(M) an antiemetic.
28	(4)	The permit holder shall maintain written emergency and patient discharge protocols accessible from
29		the operatory and recovery rooms. The written emergency manual shall include a protocol for
30		activation of emergency management services for life-threatening complications along with the
31		information set out in Rule .0101(17) of this Section.
32	(5)	The permit holder shall satisfy any additional facility requirements applicable to the level of the
33		permit, as set out in Rule .0202, .0206, .0302, or .0405 of this Subchapter.
34	(d) The permit	holder shall ensure that the following staffing, education, and training requirements are met prior to
35	performing a se	dation procedure:

I	(1)	The permit holder shall provide training to familiarize all auxiliaries in the treatment of clinical
2		emergencies including the following, and shall review and practice responding to clinical
3		emergencies with all auxiliaries as a team and in person every six months;
4		(A) airway obstruction;
5		(B) allergic reactions;
6		(C) angina pectoris;
7		(D) apnea;
8		(E) bradycardia;
9		(F) bronchospasm;
10		(G) cardiac arrest;
11		(H) convulsions;
12		(I) emesis and aspiration;
13		(J) hypertension;
14		(K) hypoglycemia;
15		(L) hypotension;
16		(M) hypoventilation and respiratory arrest;
17		(N) hypoxemia and hypoxia;
18		(O) laryngospasm;
19		(P) myocardial infarction; and
20		(Q) syncope.
21	(2)	All auxiliaries in the facility shall be BLS certified.
22	(3)	Except as set out in Subparagraph (d)(4) of this Rule, the permit holder performing the surgery or
23		other dental procedure shall ensure that an RN or a BLS-certified auxiliary is dedicated to patient
24		monitoring and recording anesthesia or sedation data throughout the sedation procedure.
25	(4)	The requirement set out in Subparagraph (d)(3) of this Rule shall not apply if the permit holder or
26		an additional sedation provider is dedicated to patient care and monitoring regarding anesthesia or
27		moderate sedation throughout the sedation procedure and is not performing the surgery or other
28		dental procedure. The additional sedation provider shall be:
29		(A) a dentist holding a permit or mobile permit in satisfaction of this Subchapter to administer
30		the anesthesia or sedation level at the facility where the sedation procedure is performed;
31		(B) an anesthesiologist licensed and practicing in accordance with the rules of the North
32		Carolina Medical Board; or
33		(C) a CRNA licensed and practicing in accordance with the rules of the North Carolina Board
34		of Nursing, under the supervision and direction of the permit holder who shall ensure the
35		level of sedation administered does not exceed the level of the sedation allowed by the
36		permit holder's permit.

1	(5)	The permit holder shall satisfy any additional staffing, education, and training requirements
2		applicable to the level of the permit, as set out in Rule .0202, .0302, or .0405 of this Subchapter.
3	(e) Before star	ting any sedation procedure, the permit holder shall conduct a pre-operative patient evaluation which
4	shall include, b	ut is not limited to, include the following:
5	(1)	evaluate evaluating the patient for health risks relevant to the potential sedation procedure;
6	(2)	evaluate evaluating the patient's food and fluid intake following the ASA guidelines for pre-
7		operative fasting applicable to elective procedures involving the administration of anesthesia or
8		moderate sedation. The ASA guidelines are incorporated by reference, including subsequent
9		amendments and editions, and may be accessed at https://www.asahq.org at no cost; and
10	(3)	satisfy satisfying any additional requirements for pre-operative patient evaluation and procedures
11		applicable to the level of the permit, as set out in Rule .0202, .0302, or .0405 of this Subchapter.
12	(f) During the	sedation procedure:
13	(1)	Prescriptions intended to accomplish procedural sedation, including enteral dosages, shall be
14		administered only under the direct supervision of the permit holder.
15	(2)	If IV sedation is used, IV infusion shall be administered before the start of the procedure and
16		maintained until the patient is ready for discharge.
17	(3)	Capnography shall be used to monitor patients unless an individual patient's behavior or condition
18		prevents use of capnography. In that event, the permit holder shall document in the sedation record
19		the clinical reason capnography could not be used.
20	(4)	The permit holder shall ensure the patient's base line baseline vital signs are taken and recorded,
21		including temperature, SPO2, blood pressure, and pulse.
22	(5)	The permit holder shall ensure the patient's blood pressure, oxygen saturation, ET CO2 (unless
23		capnography cannot be used), pulse, and respiration rates ("vital sign information") are monitored
24		continuously in a manner that enables the permit holder to view vital sign trends throughout the
25		procedure.
26	(6)	The permit holder shall ensure the intraoperative vital sign information is recorded on the anesthesia
27		or sedation record contemporaneously throughout the procedure in intervals of five minutes or less
28		for patients over twelve years old, and in intervals of ten minutes or less for pediatric patients twelve
29		years old or younger.
30	(7)	The permit holder shall satisfy any additional requirements for operative procedures applicable to
31		the level of the permit, as set out in Rule .0202, .0302, or .0405 of this Subchapter.
32	(g) Post-operat	tive monitoring and discharge shall include the following:
33	(1)	The permit holder or an auxiliary under his or her direct supervision shall monitor the patient's post-
34		operative vital signs sign information until the patient is recovered and is ready for discharge from
35		the office. Recovery from anesthesia or moderate sedation shall include documentation of the
36		following:
37		(A) stable cardiovascular function;

1		(B)	uncompromised airway patency;
2		(C)	patient arousable and protective reflexes intact;
3		(D)	state of hydration within normal limits;
4		(E)	patient can talk, if applicable;
5		(F)	patient can sit unaided, if applicable;
6		(G)	patient can ambulate with minimal assistance, if applicable; and
7		(H)	for a special needs patient, the pre-sedation level of responsiveness or the level as close as
8			possible for that patient shall be achieved.
9	(2)	Before	allowing the patient to leave the office, the permit holder shall determine that the patient has
10		met th	e recovery criteria set out in Subparagraph (g)(1) of this Rule and the following discharge
11		criteria	1:
12		(A)	oxygenation, circulation, activity, skin color, and level of consciousness are stable and have
13		(D)	been documented;
14		(B)	explanation and documentation of written post-operative instructions have been provided
15		(6)	to the patient or a person responsible for the patient at time of discharge; and
16		(C)	a person authorized by or responsible for the patient is available to transport the patient
17			after discharge.
18	•		nall maintain the following in the patient treatment records for 10 years:
19	(1)	-	tient's current written medical history history, including known allergies and previous
20		surger	
21	(2)	-	operative assessment as set out in Paragraph (e) of this Rule;
22	(3)		at to the procedure and to the anesthesia or sedation, signed by the patient or guardian,
23			ying the procedure and its risks and benefits, the level of anesthesia or sedation and its risks
24			nefits, and the date signed;
25	(4)	the and	esthesia or sedation record that shall include:
26		(A)	the patient's base line baseline vital signs and intraoperative vital sign information as set
27			out in Subparagraphs (f)(4)-(6) of this Rule;
28		(B)	the printed or downloaded vital sign information from the capnograph. A permit holder's
29			failure to maintain capnograph documentation, except as set out in Subparagraph (f)(3) of
30			this Rule, shall be deemed a failure to monitor the patient as required pursuant to this
31			Subchapter;
32		(C)	procedure start and end times;
33		(D)	gauge of needle and location of IV on the patient, if used;
34		(E)	the total amount of any local anesthetic administered during the procedure;
35		(F)	any analgesic, sedative, pharmacological, or reversal agent, or other drugs administered
36			during the procedure, including route of administration, dosage, strength, time, and

1		sequence of administration, with separate entries for each increment of medication that is
2		titrated to effect;
3		(G) documentation of complications or morbidity, and clinical responses; and
4		(H) status of patient upon discharge, including documentation of satisfying the requirements
5		set out in Paragraph (g) of this Rule; and
6	(5)	any additional documentation applicable to the level of the permit, as set out in Rule .0202, .0302,
7		or .0405 of this Subchapter.
8		
9	History Note:	Authority G.S. 90-28; 90-30.1; 90-31.1; 90-48;
10		Eff. February 1, 2023.

21 NCAC 16Q .0104 is adopted with changes as published in 37:07 NCR 545 as follows: REQUIREMENTS FOR INSPECTIONS AND EVALUATIONS 21 NCAC 16Q .0104 (a) During a facility inspection pursuant to the rules of this Subchapter, for a dentist applying for or holding a permit to administer general anesthesia, moderate conscious sedation, or moderate pediatric conscious sedation, the applicant or permit holder shall demonstrate satisfaction of the requirements set forth in Rule .0103(c) and (d) of this Section. (b) During an evaluation, for a dentist applying for or holding a permit to administer general anesthesia, moderate conscious sedation, or moderate pediatric conscious sedation, the applicant or permit holder shall demonstrate the administration of anesthesia or sedation in accordance with the level of the permit, and shall demonstrate competency including but not limited to in the following areas: (1) pre-operative patient evaluation and procedures, including the requirements set forth in Rule .0103(e) of this Section; (2) operative procedures, including the deployment of an intravenous delivery system and the requirements set forth in Rule .0103(f) of this Section; (3) post-operative patient monitoring and discharge, including the requirements set forth in Rule .0103(g) of this Section; and **(4)** treatment of the clinical emergencies set out in Rule .0103(d)(1) of this Section. (c) During the evaluation, the applicant shall take a written examination on the topics set forth in Paragraph (b) Subparagraphs (b)(1)-(4) of this Rule. The applicant shall obtain a passing score on the written examination by answering 80 percent of the examination questions correctly. If the applicant fails to obtain a passing score on the written examination, he or she may be re-examined in accordance with Rule .0204(h), .0306(h), or .0408(h) of this Subchapter.

(d) The permit holder shall be subject to re-evaluation every five years. Each facility where the permit holder

1 of 1

administers anesthesia or sedation shall be subject to a facility inspection upon annual renewal of the permit.

History Note: Authority G.S. 90-28; 90-30.1; 90-48;

Eff. February 1, 2023.

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1	21 NCAC 16Q .0202 is amended as published in 37:07 NCR 546-48 as follows:
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3	21 NCAC 16Q .0202 GENERAL ANESTHESIA EQUIPMENT AND CLINICAL REQUIREMENTS
4	(a) A dentist administering holding or applying for a permit to administer general anesthesia shall ensure that the
5	facility where the general anesthesia is administered meets the following requirements: be subject to the requirements
6	set out in Section .0100 of this Subchapter.
7	(b) In addition to the drugs listed in Rule .0103(c)(3) of this Subchapter, an unexpired neuromuscular blocking agen
8	shall be immediately available and be accessible from the operatory and recovery rooms.
9	(1) The facility shall be equipped with the following:
10	(A) an operatory of size and design to permit access of emergency equipment and personne
11	and to permit emergency management;
12	(B) a CPR board or dental chair without enhancements, suitable for providing emergency
13	treatment;
14	(C) lighting as necessary for specific procedures and back up lighting;
15	(D) suction equipment as necessary for specific procedures, including non electrical back up
16	suction;
17	(E) positive pressure oxygen delivery system, including full face masks for small, medium
18	and large patients, and back up E cylinder portable oxygen tank apart from the centra
19	system;
20	(F) small, medium, and large oral and nasal airways;
21	(G) blood pressure monitoring device;
22	(H) EKG monitor;
23	(I) pulse oximeter;
24	(J) automatic external defibrillator (AED);
25	(K) precordial stethoscope or capnograph;
26	(L) thermometer;
27	(M) vascular access set up as necessary for specific procedures, including hardware and fluids
28	(N) laryngoscope with working batteries;
29	(O) intubation forceps and advanced airway devices;
30	(P) tonsillar suction with back up suction;
31	(Q) syringes as necessary for specific procedures; and
32	(R) tourniquet and tape.
33	(2) The following unexpired drugs shall be maintained in the facility and with access from the operatory
34	and recovery rooms:
35	(A) Epinephrine;
36	(B) Atropine;
37	(C) antiarrhythmic;

1		(D) antihistamine;
2		(E) antihypertensive;
3		(F) bronchodilator;
4		(G) antihypoglycemic agent;
5		(H) vasopressor;
6		(I) corticosteroid;
7		(J) anticonvulsant;
8		(K) muscle relaxant;
9		(L) appropriate reversal agents;
10		(M) nitroglycerine;
11		(N) antiemetic; and
12		(O) Dextrose.
13	(3)	The permit holder shall maintain written emergency and patient discharge protocols. The permit
14		holder shall also provide training to familiarize auxiliaries in the treatment of clinical emergencies.
15	(4)	The permit holder shall maintain the following records for 10 years:
16		(A) Patient's current written medical history, including a record of known allergies and
17		previous surgeries;
18		(B) Consent to general anesthesia, signed by the patient or guardian, identifying the risks and
19		benefits, level of anesthesia, and date signed;
20		(C) Consent to the procedure, signed by the patient or guardian identifying the risks, benefits,
21		and date signed; and
22		(D) Patient base line vital signs, including temperature, SPO2, blood pressure, and pulse.
23	(5)	The anesthesia record shall include:
24		(A) base line vital signs, blood pressure (unless patient behavior prevents recording), oxygen
25		saturation, ET CO2 if capnography is utilized, pulse and respiration rates of the patient
26		recorded in real time at 15 minute intervals;
27		(B) procedure start and end times;
28		(C) gauge of needle and location of IV on the patient, if used;
29		(D) status of patient upon discharge; and
30		(E) documentation of complications or morbidity.
31	(6)	The facility shall be staffed with at least two BLS certified auxiliaries, one of whom shall be
32		dedicated to patient monitoring and recording general anesthesia or sedation data throughout the
33		sedation procedure. This Subparagraph shall not apply if the dentist permit holder is dedicated to
34		patient care and monitoring regarding general anesthesia or sedation throughout the sedation
35		procedure and is not performing the surgery or other dental procedure.
36	(b) During an	inspection or evaluation, the applicant or permit holder shall demonstrate the administration of
37	anesthesia while	the evaluator observes, and shall demonstrate competency in the following areas:

1	(1) monitoring of blood pressure, pulse, ET CO2 if capnography is utilized, and respiration;
2	(2) drug dosage and administration;
3	(3) treatment of untoward reactions including respiratory or cardiac depression;
4	(4) sterile technique;
5	(5) use of BLS certified auxiliaries;
6	(6) monitoring of patient during recovery; and
7	(7) sufficiency of patient recovery time.
8	(c) During an inspection or evaluation, the applicant or permit holder shall demonstrate competency in the treatment
9	of the following clinical emergencies:
10	(1) laryngospasm;
11	(2) bronchospasm;
12	(3) emesis and aspiration;
13	(4) respiratory depression and arrest;
14	(5) angina pectoris;
15	(6) myocardial infarction;
16	(7) hypertension and hypotension;
17	(8) syncope;
18	(9) allergic reactions;
19	(10) convulsions;
20	(11) bradycardia;
21	(12) hypoglycemia;
22	(13) cardiac arrest; and
23	(14) airway obstruction.
24	(d) During the evaluation, the permit applicant shall take a written examination on the topics set forth in Paragraphs
25	(b) and (c) of this Rule. The permit applicant must obtain a passing score on the written examination by answering 80
26	percent of the examination questions correctly. If the permit applicant fails to obtain a passing score on the written
27	examination that is administered during the evaluation, he or she may be re examined in accordance with Rule
28	.0204(h) of this Section.
29	(e) A general anesthesia permit holder shall evaluate a patient for health risks before starting any anesthesia procedure.
30	(f) Post operative monitoring and discharge shall include the following:
31	(1) the permit holder or a BLS certified auxiliary under his or her direct supervision shall monitor the
32	patient's vital signs throughout the sedation procedure until the patient is recovered as defined by
33	Subparagraph (f)(2) of this Rule and is ready for discharge from the office; and
34	(2) recovery from general anesthesia shall include documentation of the following:
35	(A) cardiovascular function stable;
36	(B) airway patency uncompromised;
37	(C) patient arousable and protective reflexes intact;

1		(D) state of hydration within normal limits;
2		(E) patient can talk, if applicable;
3		(F) patient can sit unaided, if applicable;
4		(G) patient can ambulate, if applicable, with minimal assistance; and
5		(H) for the special needs patient or a patient incapable of the usually expected responses, the
6		pre sedation level of responsiveness or the level as close as possible for that patient shall
7		be achieved; and
8	(3)	before allowing the patient to leave the office, the dentist shall determine that the patient has met
9		the recovery criteria set out in Subparagraph (f)(2) of this Rule and the following discharge criteria:
10		(A) oxygenation, circulation, activity, skin color, and level of consciousness are stable and have
11		been documented;
12		(B) explanation and documentation of written postoperative instructions have been provided
13		to the patient or a person responsible for the patient at time of discharge; and
14		(C) a person authorized by the patient is available to transport the patient after discharge.
15		
16	History Note:	Authority G.S. 90-28; 90-30.1; 90-48;
17		Eff. February 1, 1990;
18		Amended Eff. June 1, 2017; November 1, 2013; August 1, 2002; August 1, 2000;
19		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. January 9,
20		2018;
21		Amended Eff February 1 2023: February 1 2019: August 1 2018

I	21 NCAC 16Q .0302 is amended with changes as published in 37:07 NCR 548-50 as follows:
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3	21 NCAC 16Q .0302 MODERATE PARENTERAL AND ENTERAL CONSCIOUS SEDATION
4	CLINICAL REQUIREMENTS AND EQUIPMENT
5	(a) A dentist administering holding or applying for a permit to administer moderate conscious sedation or supervising
6	any CRNA employed to administer or RN employed to deliver moderate conscious sedation shall ensure that the
7	facility where the sedation is administered meets the following requirements: be subject to the requirements set out is
8	Section .0100 of this Subchapter.
9	(b) In addition to the drugs listed in Rule .0103(c)(3) of this Subchapter, an unexpired muscle relaxant shall b
10	immediately available and be accessible from the operatory and recovery rooms.
11	(c) As part of the pre-operative assessment required by Rule .0103(e) of this Subchapter, the permit holder shall
12	evaluate the patient for health risks as follows:
13	(1) a patient who is medically stable and who is ASA I or II shall be evaluated by reviewing the patient
14	current medical history and medication use; or
15	(2) a patient who is not medically stable or who is ASA III or higher shall be evaluated by the permit
16	holder's consultation with the patient's primary care physician or consulting medical specialis
17	regarding the potential risks posed by the planned dental procedure.
18	(d) During the sedation procedure, a moderate conscious sedation permit holder shall not administer anesthetic of
19	sedative agents:
20	(1) designed by the manufacturer for use in administering general anesthesia or deep sedation;
21	(2) <u>determined by the manufacturer to be</u> contraindicated for use in moderate conscious sedation; or
22	(3) in amounts exceeding the manufacturers' maximum recommended dosages, unless the permit holde
23	documents in the sedation record the clinical reason for exceeding the maximum recommended
24	dosage for the patient.
25	(1) The facility shall be equipped with the following:
26	(A) an operatory of size and design to permit access of emergency equipment and personne
27	and to permit emergency management;
28	(B) a CPR board or a dental chair without enhancements, suitable for providing emergency
29	treatment;
30	(C) lighting as necessary for specific procedures and back up lighting;
31	(D) suction equipment as necessary for specific procedures, including non electrical back up
32	suction;
33	(E) positive pressure oxygen delivery system, including full face masks for small, medium
34	and large patients and back up E cylinder portable oxygen tank apart from the centra
35	system;
36	(F) small, medium, and large oral and nasal airways;
37	(G) blood pressure monitoring device;

1		(H) EKG monitor;
2		(I) pulse oximeter;
3		(J) automatic external defibrillator (AED);
4		(K) precordial stethoscope or capnograph;
5		(L) thermometer;
6		(M) vascular access set up as necessary for specific procedures, including hardware and fluids;
7		(N) laryngoscope with working batteries;
8		(O) intubation forceps and advanced airway devices;
9		(P) tonsillar suction with back up suction;
10		(Q) syringes as necessary for specific procedures; and
11		(R) tourniquet and tape.
12	(2)	The following unexpired drugs shall be maintained in the facility and with access from the operatory
13		and recovery rooms:
14		(A) Epinephrine;
15		(B) Atropine;
16		(C) antiarrhythmie;
17		(D) antihistamine;
18		(E) antihypertensive;
19		(F) bronchodilator;
20		(G) antihypoglycemic agent;
21		(H) vasopressor;
22		(I) corticosteroid;
23		(J) anticonvulsant;
24		(K) muscle relaxant;
25		(L) appropriate reversal agents;
26		(M) nitroglycerine;
27		(N) antiemetic; and
28		(O) Dextrose.
29	(3)	The permit holder shall maintain written emergency and patient discharge protocols. The permit
30		holder shall also provide training to familiarize auxiliaries in the treatment of clinical emergencies;
31	(4)	The dentist shall maintain the following records for at least 10 years:
32		(A) patient's current written medical history and pre operative assessment;
33		(B) drugs administered during the procedure, including route of administration, dosage,
34		strength, time, and sequence of administration; and
35		(C) a sedation record;
26	(5)	The codetion record shall include:

1	(A) base line vital signs, blood pressure (unless patient behavior prevents recording), oxygen
2	saturation, ET CO2 if capnography is utilized, pulse and respiration rates of the patient
3	recorded in real time at 15 minute intervals;
4	(B) procedure start and end times;
5	(C) gauge of needle and location of IV on the patient, if used;
6	(D) status of patient upon discharge;
7	(E) documentation of complications or morbidity; and
8	(F) consent form, signed by the patient or guardian, identifying the procedure, risks and
9	benefits, level of sedation, and date signed; and
10	(6) The following conditions shall be satisfied during a sedation procedure:
11	(A) The facility shall be staffed with at least two BLS certified auxiliaries, one of whom shall
12	be dedicated to patient monitoring and recording sedation data throughout the sedation
13	procedure. This Subparagraph shall not apply if the dentist permit holder is dedicated to
14	patient care and monitoring regarding sedation throughout the sedation procedure and is
15	not performing the surgery or other dental procedure; and
16	(B) If IV sedation is used, IV infusion shall be administered before the start of the procedure
17	and maintained until the patient is ready for discharge.
18	(b) During an inspection or evaluation, the applicant or permit holder shall demonstrate the administration of moderate
19	conscious sedation on a patient, including the deployment of an intravenous delivery system, while the evaluator
20	observes. During the demonstration, the applicant or permit holder shall demonstrate competency in the following
21	areas:
22	(1) monitoring blood pressure, pulse, ET CO2 if capnography is utilized, and respiration;
23	(2) drug dosage and administration;
24	(3) treatment of untoward reactions including respiratory or cardiac depression if applicable;
25	(4) sterile technique;
26	(5) use of BLS certified auxiliaries;
27	(6) monitoring of patient during recovery; and
28	(7) sufficiency of patient recovery time.
29	(c) During an inspection or evaluation, the applicant or permit holder shall demonstrate competency to the evaluator
30	in the treatment of the following clinical emergencies:
31	(1) laryngospasm;
32	(2) bronchospasm;
33	(3) emesis and aspiration;
34	(4) respiratory depression and arrest;
35	(5) angina pectoris;
36	(6) myocardial infarction;
37	(7) hypertension and hypotension:

1	(8) allergic reactions;
2	(9) convulsions;
3	(10) syncope;
4	(11) bradycardia;
5	(12) hypoglycemia;
6	(13) cardiac arrest; and
7	(14) airway obstruction.
8	(d) During the evaluation, the permit applicant shall take a written examination on the topics set forth in Paragraphs
9	(b) and (c) of this Rule. The permit applicant must obtain a passing score on the written examination by answering 80
10	percent of the examination questions correctly. If the permit applicant fails to obtain a passing score on the written
11	examination that is administered during the evaluation, he or she may be re examined in accordance with Rule
12	.0306(h) of this Section.
13	(e) A moderate conscious sedation permit holder shall evaluate a patient for health risks before starting any sedation
14	procedure as follows:
15	(1) a patient who is medically stable and who is ASA I or II shall be evaluated by reviewing the patient's
16	current medical history and medication use or;
17	(2) a patient who is not medically stable or who is ASA III or higher shall be evaluated by a consultation
18	with the patient's primary care physician or consulting medical specialist regarding the potential
19	risks posed by the procedure.
20	(f) Post operative monitoring and discharge:
21	(1) the permit holder or a BLS certified auxiliary under his or her direct supervision shall monitor the
22	patient's vital signs throughout the sedation procedure until the patient is recovered as defined in
23	Subparagraph (f)(2) of this Rule and is ready for discharge from the office.
24	(2) recovery from moderate conscious sedation shall include documentation of the following:
25	(A) cardiovascular function stable;
26	(B) airway patency uncompromised;
27	(C) patient arousable and protective reflexes intact;
28	(D) state of hydration within normal limits;
29	(E) patient can talk, if applicable;
30	(F) patient can sit unaided, if applicable;
31	(G) patient can ambulate, if applicable, with minimal assistance; and
32	(H) for the special needs patient or patient incapable of the usually expected responses, the pre-
33	sedation level of responsiveness or the level as close as possible for that patient shall be
34	achieved.
35	(3) before allowing the patient to leave the office, the dentist shall determine that the patient has met
36	the recovery criteria set out in Subparagraph (f)(2) of this Rule and the following discharge criteria:

I		(A) oxygenation, circulation, activity, skin color, and level of consciousness are stable, and
2		have been documented;
3		(B) explanation and documentation of written postoperative instructions have been provided
4		to the patient or a person responsible for the patient at the time of discharge; and
5		(C) a person authorized by the patient is available to transport the patient after discharge.
6		
7	History Note:	Authority G.S. 90-28; 90-30.1; 90-48;
8		Eff. February 1, 1990;
9		Amended Eff. August 1, 2002; August 1, 2000;
10		Temporary Amendment Eff. December 11, 2002;
11		Amended Eff. June 1, 2017; November 1, 2013; July 1, 2010; July 3, 2008; August 1, 2004;
12		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. January 9,
13		2018;
14		Amended Eff. <u>February 1, 2023;</u> February 1, 2019; August 1, 2018.

16 5 of 5

1	21 NCAC 16Q .0405 is amended with changes as published in 37:07 NCR 550-52 as follows:					
2						
3	21 NCAC 16Q .0405	MODERATE	PEDIATRIC	CONSCIOUS	SEDATION	CLINICAL
4			NTS AND EQUIPN			
5	(a) A dentist administer		-		-	
6	ensure that the facility			meets the following	requirements: be	subject to the
7	requirements set out in		*			
8	(b) In addition to the	-	. , . ,	*	expired muscle re	elaxant shall be
9	immediately available a	and be accessible from	m the operatory and	d recovery rooms.		
10	(c) In addition to the	-		· -	-	
11	manual shall include as	signments to be perf	formed in the event	t of emergency by a	BLS-certified aux	iliary dedicated
12	to patient monitoring.					
13	(d) In addition to the re	equirements set out i	in Rule .0103(e) of	this Subchapter con	cerning pre-operat	ive procedures,
14	the permit holder shall	ensure that patients	who have been ad	ministered moderate	pediatric conscio	ous sedation are
15	monitored for alertness	s, responsiveness, b	reathing, and skin	coloration during	waiting periods b	efore operative
16	procedures by the perm	<u>it holder or an auxili</u>	ary dedicated to pa	tient monitoring.		
17	(e) As part of the pre-	-operative assessmer	nt required by Rule	e .0103(e) of this Su	bchapter, the peri	nit holder shall
18	evaluate the patient for	health risks as follow	<u>ws:</u>			
19	<u>(1) a pati</u>	ent who is medically	stable and who is A	ASA I or II shall be ev	valuated by review	ing the patient's
20	curre	nt medical history an	nd medication use;	<u>or</u>		
21	<u>(2)</u> a pati	ent who is not medi-	cally stable or who	is ASA III or highe	r shall be evaluate	d by the permit
22	holde	er's consultation with	h the patient's prin	nary care physician	or consulting me	edical specialist
23	regare	ding the potential ris	ks posed by the pla	nned dental procedu	re.	
24	(f) If a patient immobil	lization device is use	d, the permit holde	r shall ensure that:		
25	(1) the de	evice is applied to av	oid airway obstruct	tion or chest restricti	on;	
26	(2) the p	atient's head positio	on and respiratory	excursions are chec	ked frequently to	ensure airway
27	<u>paten</u>	cy:				
28	(3) a han	d or foot is kept expo	osed; and			
29	(4) the pa	atient is [<mark>attended</mark>] <mark>u</mark>	nder observation b	<mark>y the permit holder (</mark>	or a BLS-certified	auxiliary at all
30	times	<u>.</u>				
31	(g) During the sedation procedure, a moderate pediatric conscious sedation permit holder shall not administe			not administer		
32	anesthetic or sedative a	gents:				
33	(1) desig	ned by the manufact	urer for use in adm	inistering general and	esthesia or deep se	edation;
34	(2) deter	mined by the manu	facturer to be cont	traindicated for use	in moderate pedi	atric conscious
35	sedati	ion; or				

1	(3)	in amounts exceeding the manufacturers' maximum recommended dosages, unless the permit holder
2		documents in the sedation record the clinical reason for exceeding the maximum recommended
3		dosage for the patient.
4	(h) In addition t	to the requirements set out in Rule .0103(h) of this Subchapter concerning the patient treatment record,
5	the permit holde	er shall maintain documentation of pre-sedation instructions and information provided to the patient
6	or person respon	nsible for the patient, which shall include:
7	<u>(1)</u>	objectives of the sedation;
8	(2)	anticipated changes in patient behavior during and after sedation;
9	(3)	instructions to the person responsible for a patient transported in a [ear] child safety seat regarding
10		patient head position to avoid airway obstruction;
11	<u>(4)</u>	a 24-hour telephone number for the permit holder or his or her BLS-certified auxiliaries; and
12	<u>(5)</u>	instructions on limitations of activities and dietary precautions.
13	(i) For purposes	s of Rule .0104(b)(2) of this Subchapter, during an evaluation, a moderate pediatric conscious sedation
14	permit holder o	r applicant shall demonstrate competency in the deployment of an intravenous delivery system as
15	<u>follows:</u>	
16	<u>(1)</u>	a permit holder or applicant who uses intravenous sedation shall demonstrate the administration of
17		moderate pediatric conscious sedation on a live patient, including the deployment of an intravenous
18		delivery system; and
19	(2)	a permit holder or applicant who does not use intravenous sedation shall describe the proper
20		deployment of an intravenous delivery system and shall demonstrate the administration of moderate
21		pediatric conscious sedation on a live patient.
22	(1)	The facility shall be equipped with the following:
23		(A) an operatory of size and design to permit access of emergency equipment and personnel
24		and to permit emergency management;
25		(B) a CPR board or a dental chair without enhancements, suitable for providing emergency
26		treatment;
27		(C) lighting as necessary for specific procedures and back up lighting;
28		(D) suction equipment as necessary for specific procedures, including non electrical back up
29		suction;
30		(E) positive pressure oxygen delivery system, including full face masks for small, medium,
31		and large patients and back up E cylinder portable oxygen tank apart from the central
32		system;
33		(F) small, medium, and large oral and nasal airways;
34		(G) blood pressure monitoring device;
35		(H) EKG monitor;
36		(I) pulse oximeter;
37		(J) automatic external defibrillator (AED);

18 2 of 6

1		(K) precordial stethoscope or capnograph;
2		(L) thermometer;
3		(M) vascular access set up as necessary for specific procedures, including hardware and fluids;
4		(N) laryngoscope with working batteries;
5		(O) intubation forceps and advanced airway devices;
6		(P) tonsillar suction with back up suction;
7		(Q) syringes as necessary for specific procedures; and
8		(R) tourniquet and tape.
9	(2)	The following unexpired drugs shall be maintained in the facility and with access from the operatory
10		and recovery rooms:
11		(A) Epinephrine;
12		(B) Atropine;
13		(C) antiarrhythmie;
14		(D) antihistamine;
15		(E) antihypertensive;
16		(F) bronchodilator;
17		(G) antihypoglycemic agent;
18		(H) vasopressor;
19		(I) corticosteroid;
20		(J) anticonvulsant;
21		(K) muscle relaxant;
22		(L) appropriate reversal agents;
23		(M) nitroglycerine;
24		(N) antiemetic; and
25		(O) Dextrose.
26	(3)	The permit holder shall maintain written emergency and patient discharge protocols. The permit
27		holder shall also provide training to familiarize auxiliaries in the treatment of clinical emergencies;
28	(4)	The following records are maintained for at least 10 years:
29		(A) patient's current written medical history and pre-operative assessment;
30		(B) drugs administered during the procedure, including route of administration, dosage,
31		strength, time, and sequence of administration;
32		(C) a sedation record; and
33		(D) a consent form, signed by the patient or a guardian, identifying the procedure, risks and
34		benefits, level of sedation, and date signed;
35	(5)	The sedation record shall include:

1	(A) base line vital signs, blood pressure (unless patient behavior prevents recording), oxygen
2	saturation, ET CO2 if capnography is utilized, pulse and respiration rates of the patient
3	recorded in real time at 15 minute intervals;
4	(B) procedure start and end times;
5	(C) gauge of needle and location of IV on the patient, if used;
6	(D) status of patient upon discharge; and
7	(E) documentation of complications or morbidity; and
8	(6) The following conditions shall be satisfied during a sedation procedure:
9	(A) the facility shall be staffed with at least two BLS certified auxiliaries, one of whom shall
10	be dedicated to patient monitoring and recording sedation data throughout the sedation
11	procedure. This Subparagraph shall not apply if the dentist permit holder is dedicated to
12	patient care and monitoring regarding sedation throughout the sedation procedure and is
13	not performing the surgery or other dental procedure; and
14	(B) when IV sedation is used, IV infusion shall be administered before the commencement of
15	the procedure and maintained until the patient is ready for discharge.
16	(b) During an inspection or evaluation, applicants and permit holders who use intravenous sedation shall demonstrate
17	the administration of moderate pediatric conscious sedation on a live patient, including the deployment of an
18	intravenous delivery system, while the evaluator observes. Applicants and permit holders who do not use IV sedation
19	shall describe the proper deployment of an intravascular delivery system to the evaluator and shall demonstrate the
20	administration of moderate pediatric conscious sedation on a live patient while the evaluator observes.
21	(c) During the demonstration, all applicants and permit holders shall demonstrate competency in the following areas:
22	(1) monitoring blood pressure, pulse, and respiration;
23	(2) drug dosage and administration;
24	(3) treatment of untoward reactions including respiratory or cardiac depression if applicable;
25	(4) sterile technique;
26	(5) use of BLS certified auxiliaries;
27	(6) monitoring of patient during recovery; and
28	(7) sufficiency of patient recovery time.
29	(d) During an inspection or evaluation, the applicant or permit holder shall demonstrate competency in the treatment
30	of the following clinical emergencies:
31	(1) laryngospasm;
32	(2) bronchospasm;
33	(3) emesis and aspiration;
34	(4) respiratory depression and arrest;
35	(5) angina pectoris;
36	(6) myocardial infarction;
37	(7) hypertension and hypotension;

1	(8)	allergic reactions;
2	(9)	convulsions;
3	(10)	syncope;
4	(11)	bradycardia;
5	(12)	hypoglycemia;
6	(13)	cardiac arrest; and
7	(14)	airway obstruction.
8	(e) During the ev	raluation, the permit applicant shall take a written examination on the topics set forth in Paragraphs
9	(c) and (d) of this	Rule. The permit applicant must obtain a passing score on the written examination by answering 80
10	percent of the exa	amination questions correctly. If the permit applicant fails to obtain a passing score on the written
11	examination that	is administered during the evaluation, he or she may be re examined in accordance with Rule
12	.0408(h) of this S	ection.
13	(f) A moderate p	ediatric conscious sedation permit holder shall evaluate patients for health risks before starting any
14	sedation procedur	e as follows:
15	(1)	a patient who is medically stable and who is ASA I or II shall be evaluated by reviewing the patient's
16		current medical history and medication use; or
17	(2)	a patient who is not medically stable or who is ASA III or higher shall be evaluated by a consultation
18		with the patient's primary care physician or consulting medical specialist regarding the potential
19		risks posed by the procedure.
20	(g) Patient monit	oring:
21	(1)	Patients who have been administered moderate pediatric conscious sedation shall be monitored for
22		alertness, responsiveness, breathing, and skin coloration during waiting periods before operative
23		procedures.
24	(2)	The permit holder or a BLS certified auxiliary under his or her direct supervision shall monitor the
25		patient's vital signs throughout the sedation procedure until the patient is recovered as defined in
26		Subparagraph (g)(3) of this Rule and is ready for discharge from the office.
27	(3)	Recovery from moderate pediatric conscious sedation shall include documentation of the following:
28		(A) cardiovascular function stable;
29		(B) airway patency uncompromised;
30		(C) patient arousable and protective reflexes intact;
31		(D) state of hydration within normal limits;
32		(E) patient can talk, if applicable;
33		(F) patient can sit unaided, if applicable;
34		(G) patient can ambulate, if applicable, with minimal assistance; and
35		(H) for the special needs patient or a patient incapable of the usually expected responses, the
36		pre sedation level of responsiveness or the level as close as possible for that patient shall
37		be achieved.

1	(4)	Before allowing the patient to leave the office, the dentist shall determine that the patient has met
2		the recovery criteria set out in Subparagraph (g)(3) of this Rule and the following discharge criteria:
3		(A) oxygenation, circulation, activity, skin color, and level of consciousness are stable, and
4		have been documented;
5		(B) explanation and documentation of written postoperative instructions have been provided
6		to a person responsible for the patient at time of discharge; and
7		(C) a person responsible for the patient is available to transport the patient after discharge, and
8		for the patient for whom a motor vehicle restraint system is required, an additional
9		responsible individual is available to attend to the patient.
10		
11	History Note:	Authority G.S. 90-28; 90-30.1; 90-48;
12		Eff. June 1, 2017;
13		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. January 9,
14		2018;
15		Amended Eff. <u>February 1, 2023;</u> February 1, 2019; August 1, 2018.

1 21 NCAC 16Q .0703 is amended with changes as published in 37:07 NCR 552-53 as follows: 2 3 21 NCAC 16Q .0703 REPORTS OF ADVERSE OCCURRENCES 4 (a) A dentist who holds a permit to administer general anesthesia or sedation shall submit an adverse occurrence 5 report to the Board within 72 hours after each adverse occurrence related to the administration of general anesthesia 6 or sedation that results in the death of a patient if the patient dies or has permanent organic brain dysfunction within 7 24 hours of after the procedure, administration of general anesthesia or sedation. Sedation permit holders shall cease 8 administration of sedation until the Board has investigated the death or permanent organic brain dysfunction and 9 approved resumption of permit privileges. General anesthesia permit holders shall cease administration of general 10 anesthesia and sedation until the Board has reviewed the incident adverse occurrence report and approved resumption 11 of permit privileges. 12 (b) A dentist who holds a permit to administer general anesthesia or sedation shall submit an adverse occurrence 13 report to the Board within 30 days after each adverse occurrence related to if the patient is admitted to a hospital on 14 inpatient status for a medical emergency or physical injury within 24 hours after the administration of general 15 anesthesia or sedation sedation that results in permanent organic brain dysfunction of a patient occurring within 24 hours of the procedure or that results in physical injury or severe medical emergencies, causing hospitalization of a 16 17 patient occurring within 24 hours of the procedure. 18 (c) The adverse occurrence report shall be in writing and shall include the following: 19 dentist's name, license number and permit number; (1) 20 (2) date and time of the occurrence; 21 (3) facility where the occurrence took place; 22 (4) name and address of the patient; 23 (5) surgical procedure involved; type and dosage of sedation or anesthesia utilized in the procedure; 24 (6)25 **(7)** circumstances involved in the occurrence; and 26 (8) the entire patient treatment record including anesthesia records. (d) Upon receipt of any such report, report submitted pursuant to this Rule, the Board shall investigate and shall take 27 28 disciplinary action if the evidence demonstrates that a licensee has violated the Dental Practice Act set forth in Article 29 2 of G.S. Chapter 90 of the General Statutes or the Board's rules of this Chapter. 30 31 History Note: Authority G.S. 90-28; 90-30.1; 90-41; 90-48;

Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. January 9,

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Eff. April 1, 2016;

Amended Eff. February 1, 2023.

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1 of 1 23