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3	10A NCAC 15.	0611 COMPUTED TOMOGRAPHY (CT) X-RAY SYSTEMS	
4	(a) This Rule pro	ovides special requirements for human diagnostic use of computed tomography (CT) x-ray equipment	
5	The uses of Cone	e Beam CT, Veterinary CT, CT Simulation, and CT attenuation correction shall be exempt from this	
6	Rule. The provis	sions of this Rule are in addition to, and not in substitution for, the Rules in Sections .0100, .0200	
7	.0600, .0900, .10	00, and .1600 of this Chapter.	
8	(b) The following	ng definitions shall apply to this Rule:	
9	(1)	"CT qualified expert (CT QE)" means an individual who is registered or is providing service for a	
10		registered facility where they are employed, as required by Section .0200 of this Chapter. The	
11		individual shall have the following education and experience:	
12		(A) a master's or doctoral degree in physics, medical physics, biophysics, radiological physics	
13		medical health physics, or equivalent disciplines from an accredited a college or university	
14		university accredited by an agency recognized by the U.S. Department of Education, and	
15		(B) three years work experience in a clinical CT environment. The work experience shall be	
16		supervised and documented by a board certified medical physicist; physicist certified in	
17		the specialty area of diagnostic medical physics by the American Board of Radiology, the	
18		Canadian College of Physicists in Medicine, or the American Board of Medical Physics	
19		or	
20		(C) (B) certification in the specific subfield(s) of specialty area of diagnostic medical physics with	
21		its associated medical health physics aspect by an appropriate national certifying body the	
22		American Board of Radiology, the Canadian College of Physicists in Medicine, or the	
23		American Board of Medical Physics and shall abide by the certifying body's requirements	
24		for continuing education.	
25	(2)	"general supervision" means the activity is performed under the qualified supervisor's overal	
26		direction and control but the qualified supervisor's physical presence is not shall not be required	
27		during the activity.	
28	(3)	"personal supervision" means overall direction, control control, and training of an individual by a	
29		qualified supervisor who must shall be physically present during the activities performed by the	
30		supervised individual.	
31	(c) Equipment a	nd Installation Requirements	
32	(1)	CT x-ray systems shall meet the requirements of 21 CFR 1020.33 as incorporated by reference in	
33		Rule .0117(a)(3) of this Chapter.	
34	(2)	The operator of a CT scanner shall be able to maintain aural communication with the patient from	
35		a shielded position at the control panel.	
36	(d) Personnel R	equirements. Individuals who operate CT x-ray systems shall: shall be specifically trained on the	
37	operational features of the unit and:		

10A NCAC 15 .0611 is adopted with changes as published in 31:19 NCAC 1862-1864 as follows:

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1	(1)	hold (CT) registration with the American Registry of Radiologic Technologists (ARRT); or
2	(2)	be a Registered Technologist (R.T.) by the ARRT with registration in radiography (R) or a Certifie
3		Nuclear Medicine Technologist by the Nuclear Medicine Technology Certification Board; thes
4		individuals shall document training and experience that is equivalent to that required to attain (CT
5		registration with the ARRT; or
6	(3)	be in training under the personal supervision of an individual that meets the requirements of
7		Paragraph (d) (d)(1) or (d)(2) of this Rule; and Rule.
8	(4)	be specifically trained on the operational features of the unit.
9	(e) System Per	formance Evaluations
10	(1)	Performance evaluations of the CT x-ray system shall be performed by, or under the general
11		supervision of, a CT QE who assumes the responsibility for the evaluation.
12	(2)	The performance evaluation of a CT x-ray system shall be performed within 30 days of installatio
13		and at least every 14 months.
14	(3)	Performance evaluation standards and tolerances shall meet manufacturer's specifications of
15		standards and tolerances for the CT x-ray system from the American College of Radiology (ACR
16		and the American Association of Physicists in Medicine (AAPM). (AAPM) incorporated herein b
17		reference including subsequent amendments and editions. These standards and tolerances may b
18		found at no charge on the ACR website at https://www.arc.org and the AAPM websites. website a
19		www.aapm.org.
20	(4)	The performance evaluation shall include the following as applicable to the design of the scanner:
21		(A) geometric factors and alignment including alignment light accuracy, and table increment
22		accuracy;
23		(B) image localization from a scanned projection radiograph (localization image);
24		(C) radiation beam width;
25		(D) image quality including high-contrast (spatial) resolution, low-contrast resolution, image
26		uniformity, noise, and artifact evaluation;
27		(E) CT number accuracy;
28		(F) image quality for acquisition workstation display devices; and
29		(G) a review of the results of the routine QC, as set forth in Paragraph (f) of this Rule; and
30	(5)	The performance evaluation shall also include the evaluation of radiation output and patient dos
31		indices for the following clinical protocols if performed:
32		(A) pediatric head;
33		(B) pediatric abdomen;
34		(C) adult head;
35		(D) adult abdomen; and
36		(E) brain perfusion.

1	(6)	Evaluation of radiation output shall be performed with a dosimetry system that is calibrated. The
2		dosimetry system shall have been calibrated within the preceding two years by persons registered
3		to provide such services pursuant to Rule .0205 of this Chapter.
4	(7)	The performance evaluation shall be documented and maintained for inspection by the Agency. The
5		documentation shall include the name of the CT QE performing or supervising the evaluation, as
6		well as any other individual(s) individuals participating in the evaluation under the general
7		supervision of the CT QE. The documentation shall be retained for 14 months.
8	(f) Routine Qua	ality Control (QC)
9	(1)	A routine QC program for the CT system shall be developed by or have written approval by a CT
10		QE and include:
11		(A) instructions for the routine QC;
12		(B) intervals for QC testing;
13		(C) acceptable tolerances for the QC tests;
14		(D) use of a water equivalent phantom to evaluate each day of clinical use: noise, CT number
15		accuracy, and artifacts; and
16		(E) routine QC tests that may be performed in place of system performance evaluations after
17		equipment repairs or maintenance. This shall include the process for obtaining approval
18		from the CT QE prior to conducting testing.
19	(2)	The duties in the routine QC program, as described in Part (f)(1) of this Rule, shall be conducted by
20		individuals that meet the requirements of Part (d) of this Rule or individuals approved by the CT
21		QE.
22	(3)	The routine QC shall be documented and maintained for inspection by the Agency. The records
23		shall be retained for 14 months.
24	(g) Operating	Requirements. The following information shall be accessible to the CT operator during use of the
25	machine and wh	nile performing routine QC:
26	(1)	instructions on performing routine QC;
27	(2)	a schedule of routine QC;
28	(3)	any allowable variations set by the CT QE for the indicated parameters;
29	(4)	the results of the most recent routine QC completed on the system; and
30	(5)	established scanning protocols.
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32	History Note:	Authority G.S. 104E-7; 104E-11; 104E-12;
33		Eff. October 1, 2017.