Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Kissinger, Regina < regina.kissinger@dhhs.nc.gov>

Sent: Friday, October 11, 2024 10:18 AM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Wiggs, Travis C <travis.wiggs@oah.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Thank you! Hope you have a great weekend.

Regina Kissinger, R.T.(R) (M) ARRT Health Physicist Healing Arts Supervisor <u>Division of Health Service Regulation, Radiation Protection</u> NC Department of Health and Human Services

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From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Friday, October 11, 2024 9:57 AM

To: Wiggs, Travis C < travis.wiggs@oah.nc.gov; Kissinger, Regina < regina.kissinger@dhhs.nc.gov

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Thanks for all of your help! Have a great weekend!!

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov>

Sent: Friday, October 11, 2024 9:51 AM

To: Black, Shanah <<u>shanah.black@dhhs.nc.gov</u>>; Kissinger, Regina <<u>regina.kissinger@dhhs.nc.gov</u>>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good morning,

I intend to recommend to the RRC that all the final revised rules be approved at the October 30th meeting. Please submit all revised rules via email to oah.rules@oah.nc.gov no later than 5pm on October 18, 2024. The electronic copy must be saved as the official rule name (XX NCAC XXXX). Please include me on the email.

Thank you both for your work.

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10A NCAC 15 .0802.docx; 10A NCAC 15 .0803.docx; 10A NCAC 15 .0804.docx; 10A

NCAC 15 .0805.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Thursday, October 10, 2024 6:57 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Hey Travis,

Please see attached.

Thanks for all of your help.

Have a great night.

10A NCAC 15 .0802 is amended with changes as published in 38:19 NCR 1219-1229 as follows:

10A NCAC 15.0802 DEFINITIONS

- In addition to terms found in Rule .0104 of this Chapter Chapter, the following definitions shall apply to this Section:
- 5 (1) "Accredited bomb squad" means a law enforcement agency utilizing certified bomb technicians.
 - (2) <u>"Accessible surface" means the external or outside surface of the enclosure or housing provided by the manufacturer or designer of the RGD. This includes the high-voltage generator, doors, access panels, latches, control knobs, and other permanently mounted hardware, and including the plane across the exterior edge of any opening.</u>
 - (2)(3) "Analytical RGD equipment" means equipment that uses electronic means to generate ionizing radiation for the purpose of examining the microstructure of materials, materials i.e. using direct x-ray transmission, x-ray diffraction, x-ray fluorescence, and x-ray spectroscopy.
 - (3)(4) "Analytical RGD system" means a group of local and remote components utilizing x-rays to determine the elemental composition or to examine the microstructure of materials.
 - (4) "Bomb detection RGDs" means RGDs used for the sole purpose of remotely detecting explosive devices.
 - (5) "Certified bomb technician" means a member of an accredited bomb squad who has <u>successfully</u> completed the FBI Hazardous Devices School. Information pertaining to this program can be found <u>on the school website</u> at http://www.fbi.gov/about-us/cirg/hazardous-devices.
 - (6) "Certifiable cabinet x-ray system" means an existing uncertified RGD that has been modified to meet the certification requirements specified in 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (7) "Certified cabinet x-ray system" means an RGD utilized in an enclosed, interlocked cabinet, such that the radiation machine will not operate unless all openings are securely closed. These systems shall be certified in accordance with 21 CFR 1010.2 1010.2, as incorporated by reference in Rule .0117 of this Chapter, as being manufactured and assembled pursuant to the provisions of 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (8) "Collimator" means a device or mechanism by which the x-ray beam is restricted in size.
 - (9) "Control panel" means that the part of the x-ray control where the switches, knobs, pushbuttons, and other hardware are, that are necessary located for manually setting the technique factors.
 - (10) "Electron Beam Device" means any device using electrons below 1MeV to heat, join, or otherwise irradiate materials.
 - (11) "Enclosed beam RGD" means an RGD with all possible x-ray beam paths contained in a chamber, coupled chambers, or other beam-path-confinement devices devices, to prevent any part of the body from intercepting the beam during normal operations. Normal access to the primary beam path, such as a sample chamber door, shall be interlocked with the high voltage of the x-ray tube or the shutter

1		for the beam to be considered "enclosed." An open-beam device placed in an interlocked enclosure
2		is considered an "enclosed beam" unless there are provisions for routine bypassing of the interlocks.
3	<u>(12)</u>	"Emergency procedure" means the written pre-planned steps to be taken in the event of actual or
4		suspected radiation exposure of an individual exceeding administrative or regulatory limits. limits
5		found in Rule 10A NCAC 15 .1601(a)(8) and .1601(a)(15). This procedure shall include the names
6		and telephone numbers of individuals to be contacted contacted, as well as directives for processing
7		individual monitoring devices.
8	(12) (13	(1) "Fail-safe characteristics" means a design feature that causes the radiation beam to terminate, port
9		shutters to close, or otherwise prevents emergence of the primary beam, beam upon the failure of a
10		safety or warning device. For example, if an "X-ray On" light indicator or light indicator, shutter
11		indicator shutter indicator, or interlock fails, the radiation beam shall terminate.
12	<u>(14)</u>	"Gauging device" means a mechanism containing a source of ionizing radiation that is designed and
13		manufactured for the purpose of determining or controlling thickness, density, level, interface
14		location, or qualitative or quantitative composition of materials. It may include components such as
15		radiation shields, useful-beam controls, and other safety features in order to meet the requirements
16		or specifications. specifications of the device.
17	<u>(15)</u>	"General-use system" means a security screening system that delivers an effective dose of 25
18		microrem (0.25 microSv) or less per screening.
19	(13) (16) "Hand-held x-ray system" means any device or equipment that is portable and used for similar
20		purposes as analytical RGD equipment.
21	(14)	"Hybrid gauge" means an x-ray gauge device utilizing both x-ray and radioactive sources.
22	(15)	"Industrial radiography" means RGDs used to make radiographic images to examine the structure
23		of materials by nondestructive methods. These RGDs shall not be contained in a cabinet and are not
24		permanent installations.
25	<u>(17)</u>	"Individual responsible for radiation protection" means a person who has the knowledge and
26		responsibility to apply appropriate radiation rules, for persons registered with the agency in
27		accordance with Section .0200 of this Chapter, commensurate with the scope of the activities
28		authorized by the registrant.
29	(17) (18	"Inspection Zone" means the area established for the purpose of controlling access where screening
30		is performed. Areas controlled due to the presence of radiation shall include areas of ingress, egress,
31		gates, portals, and traffic paths. The area outside of the inspection zone shall not exceed the limits
32		of Rule .1601(a)(13) of this Section.
33	(18) (19	"Interlock" means a feature designed to prevent access to an area of radiation hazard by preventing
34		entry or by automatically removing the hazard.
35	(16)<mark>(20</mark>	"Ion implantation equipment, low-energy" means any enclosed device operating below 1MeV used
36		to accelerate elemental ions and implant them in other materials.
37	(17)<mark>(21</mark>	Theakage radiation" means radiation emanating from the source assembly housing except for:

l	(A) the primary beam;
2	(B) scatter radiation emanating from other components (e.g., shutter or collimator); and
3	(C) radiation produced when the beam on "beam on" switch or timer is not activated.
4	(21)(22) "Limited-use system" means a screening system that is capable of delivering an effective dose
5	greater than 25 microrem (0.25 microSv) per screening screening, but shall not exceed an effective
6	dose of 1 mrem (10 microSv) per screening, screening,
7	(18)(23) "Local components" means part of an RGD x-ray system and include areas that are struck by x-rays
8	x rays, such as radiation source housings, port and shutter assemblies, collimators, sample holders,
9	cameras, goniometers, detectors, and shielding, but do not include power supplies, transformers,
10	amplifiers, readout devices, and control panels.
11	(19)(24) "Mobile RGD" means RGD equipment mounted on a permanent base with wheels or casters for
12	moving while completely assembled.
13	(20)(25) "Normal operating procedures" means step-by-step instructions necessary to accomplish a task.
14	These procedures shall include sample insertion and manipulation, equipment alignment, routine
15	maintenance by the registrant, and data recording procedures that are related to radiation safety.
16	(21)(26) "Open-beam RGD" means a device or system designed in such a way that the primary beam is not
17	completely enclosed during normal operation operation, and when used for analysis, gauging, or
18	imaging imaging, in which an individual could accidentally place some part of their body in the
19	primary beam or stray radiation path during normal operation.
20	(22) "Permanent radiographic installation" means an RGD utilized in an enclosed shielded room, cell, or
21	vault that allows entry when the RGD is not energized.
22	(23)(27) "Portable RGD" means RGD equipment designed to be carried by hand.
23	(24)(28) "Primary beam" means radiation that passes through an aperture of the source assembly housing by
24	a direct path from the radiation source.
25	(25)(29) "Radiation generating device (RGD)" means any system, device, subsystem, or machine component
26	that may generate generate, by electronic means means, x-rays or particle radiation above 5 keV,
27	but below 1 MeV, and not used for healing arts parts on humans or animals. Examples of RGDs are
28	the following may be used as a:
29	(A) analytical RGD equipment mobile RGD;
30	(B) certified and certifiable cabinet x-ray systems portable RGD; or
31	(C) gauging devices using x-ray sources; stationary RGD.
32	(D) hybrid gauging devices;
33	(E) e beam welders;
34	(F) baggage scanners;
35	(G) industrial radiography RGDs; and
36	(H) permanent radiographic installations.

1	(26)<u>(30)</u>	"Remote components" means parts of an RGD x-ray system that are not struck by x-rays x-rays,
2		such as power supplies, transformers, amplifiers, readout devices, and control panels.
3	(30) (31)	"Safety Device" means a device, interlock or system that prevents the entry of any portion of an
4		individual's body into the primary x-ray beam or that will cause the beam to shut off upon entry into
5		its path.
6	(27)<mark>(32)</mark>	"Scattered radiation" means radiation, other than leakage radiation, that during passage through
7		matter, has been deviated in direction or has been modified by a decrease in energy.
8	(32) (33)	"Screening" means the sum of scans necessary for a security screening system to image concealed
9		objects as intended by the system design under normal operating conditions.
10	(33) (34)	"Security screening device" means a non-human use open-beam device designed for the detection
11		of contraband or weapons concealed in baggage, mail, packages, or other structures. These devices
12		include bomb detection devices used for the sole purpose of detecting explosive devices.
13	(34) (35)	"Security screening system" means a system specifically designed to detect contraband and weapons
14		concealed on a person and is used for the sole purpose of public safety and security evaluation by
15		law enforcement.
16	(28)<mark>(36)</mark>	"Shutter" means an adjustable device, generally made of lead or other high atomic number material,
17		fixed to a source assembly housing to intercept, block, or collimate the primary beam.
18	(29)<mark>(37)</mark>	"Source" means the point of origin of the radiation, such as the focal spot of an x-ray tube.
19	(30)<mark>(38)</mark>	"Stationary RGD" means RGD equipment that is installed or placed in a fixed location.
20	(31)<mark>(39)</mark>	"Stray radiation" means the sum of leakage and scatter radiation emanating from the source
21		assembly or other components components, except for the primary beam, and radiation produced
22		when the beam on switch or timer is not activated.
23	(39) (40)	"Warning device" means an audible or visible signal that warns individuals of a potential radiation
24		<u>hazard.</u>
25	(32)<mark>(41)</mark>	"X-ray generator" means the part of an x-ray system that provides the accelerating (high) voltage
26		and current for the x-ray tube.
27	(33)	"X ray gauge" means an x ray producing device designed and manufactured for the purpose of
28		detecting, measuring, gauging, or controlling thickness, density, level, or interface location of
29		manufactured products.
30	(41) (42)	"X-ray source housing" means the portion of an RGD system which contains the x-ray tube and
31		emitting target. The housing often contains radiation shielding material or inherently provides
32		shielding.
33		
34	History Note:	Authority G.S. 104E-7;
35		Eff. February 1, 1980;
36		Transferred and Recodified from 15A NCAC 11 .0802 Eff. February 1, 2015;
37		Amended Eff. October 1, 2015;

1	Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22
2	2019. <u>2019;</u>
3	Amended Eff. November 1, 2024

1	10A NCAC 15 .0803 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0803 EQUIPMENT REQUIREMENTS PERSONNEL REQUIREMENTS
4	(a) Certified cabinet x ray systems shall meet the requirements of 21 CFR 1020.40 as incorporated by reference in
5	Rule .0117(a)(3) of this Chapter.
6	(b) All certified and certifiable cabinet x-ray systems shall:
7	(1) be constructed so that, the radiation emitted from the system shall not exceed an exposure of 0.5
8	milliroentgen (mR) in one hour at any point five centimeters outside the external surface; and
9	(2) have a fail safe interlock that prevents irradiation when the cabinet, chamber, or coupled chambers
10	are open.
11	(c) Open beam analytical RGD systems shall be equipped with a safety device that prevents the entry of any portion
12	of an individual's body into the primary x-ray beam path that causes the beam to be shut off upon entry into its path.
13	(d) Open beam analytical RGDs shall be provided with a visible and legible indication of:
14	(1) x ray tube status (ON OFF) located near the radiation source housing, if the primary beam is
15	controlled in this manner; or
16	(2) shutter status (OPEN CLOSED) or beam status (ON OFF) located near each port on the radiation
17	source housing, if the primary beam is controlled in this manner.
18	(e) Warning devices on open beam analytical RGDs shall be labeled so that their purpose is identified. On open beam
19	analytical RGDs installed after February 1, 1980, warning devices and lights shall have fail safe characteristics.
20	(f) Unused ports on radiation source housings for open-beam RGDs shall be secured in the closed position in a manner
21	that will prevent unintended opening.
22	(g) Each port on the radiation source housing on open beam analytical RGDs installed after February 1, 1980 and
23	designed to accommodate interchangeable components shall be equipped with a shutter that cannot be opened unless
24	a collimator or a component coupling is connected to the port.
25	(h) Portable open beam analytical RGDs that shall be manufactured to be used hand held without safety devices are
26	exempt from the requirements of Paragraph (c) of this Rule and shall be constructed according to International
27	Standard IEC 62495 that is incorporated by reference and includes subsequent amendments. This standard can be
28	downloaded for one hundred twenty one dollars (\$121.00) at the following website:
29	http://webstore.ansi.org/FindStandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx.aspx.aspx.aspx.aspx.aspx.aspx.asp
30	geNum=0&SearchTermsArray=null%7cIEC+62495+Ed.+1.0+en%3a2011%7cnull.
31	(i) A registrant may apply to the agency, as defined in Rule .0104 of this Chapter, for an exemption from the
32	requirement of a safety device. This request shall include:
33	(1) a description of the safety devices;
34	(2) the reason safety devices cannot be used; and
35	(3) a description of the alternative methods that will be employed to minimize the possibility of an
36	accidental exposure, including procedures to assure that operators and others in the area will be
37	informed of the absence of safety devices.

1	(j) Analytical RGDs shall be provided with a visible and legible label(s) bearing the radiation symbol and the words:
2	(1) "CAUTION HIGH INTENSITY X RAY BEAM," or words having a similar meaning, near the
3	exit port to identify the location of the beam; and
4	(2) "CAUTION RADIATION THIS EQUIPMENT PRODUCES RADIATION WHEN
5	ENERGIZED", or words having a similar meaning, near any switch that energizes an x-ray tube, if
6	the radiation source is an x-ray tube.
7	(k) Warning lights labeled with the words "X RAYS ON," or other words having similar meaning, shall be located:
8	(1) near any switch that activates the high voltage to energize an x-ray tube; or
9	(2) in a conspicuous location near the radiation source housing and radiation beam(s) and visible from
10	all instrument access areas.
11	(1) Warning lights shall activate when the x-ray tube is energized.
12	(m) Each x ray tube housing shall be:
13	(1) constructed that when all shutters are closed the leakage radiation measured at a distance of five
14	centimeters from its surface is not capable of producing an exposure in excess of 2.5 millirem
15	(mrem)/ (25 microsieverts µSv) in one hour; and
16	(2) if the tube housing is the primary shielding for the x ray tube, does not produce x rays when the
17	housing is opened or disassembled.
18	(n) Each x ray generator shall be supplied with a protection cabinet which limits leakage radiation measured at a
19	distance of five centimeters from its surface such that it is not capable of producing an exposure in excess of 0.25
20	mrem/2.5μSv in one hour.
21	$ \hbox{(o) Permanent radiographic installations and industrial radiography RGDs shall comply with the requirements of Rule } \\$
22	.0807 of this Section.
23	(a) The registrant registrant, as defined in 10A NCAC 15 .0104(130), shall document the scope of training and
24	instruction required for the RGD in use.
25	(b) No individual shall be permitted to operate or maintain RGDs unless the individual has received instruction in the
26	basic principles of radiation protection, training specific to the manufacturer's recommendations for safe operation
27	and unique features of the RGD in use, and instruction in the operating and emergency procedures. Instruction and
28	training shall include:
29	(1) Basic principles of radiation protection:
30	(A) radiation fundamentals;
31	(B) source and magnitude of common sources of radiation exposure;
32	(C) units of radiation dose and measurements:
33	(D) potential hazards, biological effects of ionizing radiation, and recognition of symptoms of
34	an acute localized exposure;
35	(E) ALARA (As Low As Reasonably Achievable) principles for radiation protection concepts
36	of time, distance, and shielding to minimize radiation exposure;
37	(F) declared pregnancy policy;

I		(G) occupational, embryo/tetus, and public dose limits; and
2		(H) proper use of individual monitoring devices and survey instruments.
3	(2)	Device specific training for each RGD:
4		(A) hands-on training for proper use;
5		(B) radiation hazards associated with use;
6		(C) precautions to take or measures required to minimize radiation exposure;
7		(D) procedures to prevent unauthorized use; and
8		(E) agency rules regarding use.
9	(3)	Operating and emergency procedure requirements of Rule .0804 in this Section.
10	(c) Records of i	nstruction and training for each individual operating RGDs, documenting that the requirements of this
11	Rule have been	met, shall be maintained and available for agency review during inspection.
12	<u>(d) <mark>Individuals</mark> j</u>	Persons who will be operating the RGD shall be able to demonstrate an understanding in safe operating
13	procedures and	use of the RGD according to the manufacturer's specifications and to an authorized representative of
14	the Radiation Pr	rotection Section.
15	(e) Each registr	ant shall provide ring or wrist individual monitoring devices to individuals:
16	(1)	operating open-beam RGDs; and
17	<u>(2)</u>	performing maintenance on an RDG, if the maintenance procedures require the presence of a
18		primary x-ray beam when any local component in the RGD is disassembled or removed.
19		
20	History Note:	Authority G.S. 104E-7;
21		Eff. February 1, 1980;
22		Transferred and Recodified from 15A NCAC 11 .0803 Eff. February 1, 2015;
23		Amended Eff. October 1, 2015;
24		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
25		2019. <u>2019:</u>
26		<u>Amended Eff.</u> <mark>October </mark>

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer individual responsible
29	for radiation protection as defined in 10A NCAC 15 .0802(17). protection. as defined in Rule.0104 of this Chapter.
30	
31	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
32	all operators and support staff for review during the use of an RGD.
33	(d) Normal operating and emergency procedures shall include the following:
34	(1) safe use of the RGD;
35	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
36	<u>and</u>

any unusual occurrence or malfunction that may involve exposure to radiation. (c) Open beam and portable handheld RGDs (l) Registrants shall have operating procedures developed to ensure radiation protective measure (A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation where the primary x-ray beam is not intercepted by a detector device during operation (C) available to all individuals operating the RGD. (2) Operators shall not do the following while operating an RGD: (A) point the primary beam at any individual including him or herself; (B) allow their hand to approach the primary beam: or (C) hold a sample. If a sample is small and it is necessary to hold the sample while operating and emergency procedures shall be paleed in a shielded sample enclosure. (f) Operating and emergency procedures shall be available for review by the individual responsible for radiation protection agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the radiation safety officer india responsible for radiation protection and manufacturer of the RGD. (i) Safety Devices (i) Testing (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD testoms are established and approved in writing by the radiation safety officer india responsible for radiation protection. (B) Testing records shall include the date test was performed, the list of safety devices to the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. (C) Records of the testing shall be retained by the registrant for a	1	(3)	instructions on reporting to the radiaton safety officer individual responsible for radiation protection
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(1) Registrants shall have operating procedures developed to ensure radiation protective measure (A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation (C) available to all individuals operating the RGD. (C) operators shall not do the following while operating an RGD: (A) point the primary beam at any individual including him or herself; (B) allow their hand to approach the primary beam; or (C) hold a sample, If a sample is small and it is necessary to hold the sample while operating and emergency procedures shall be paralled in a shielded sample enclosure. (f) Operating and emergency procedures shall be available for review by the individual responsible for radiation agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the radiation safety officer individual responsible for radiation protection and manufacturer of the RGD. (i) Safety Devices (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD be taken out of service until corrective action is performed or temporary administ controls are established and approved in writing by the radiation-safety officer individual responsible for radiation protection. (B) Testing records shall include the date test was performed, the list of safety devices in the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. (C) Records of the testing shall be retained by the registrant for agency review of inspection. (A) No individual shall bypass a safety device unless the person has obtained approval the radiation protection pro	3		any unusual occurrence or malfunction that may involve exposure to radiation.
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22 (A) Safety devices including interlocks, shutters, and warning lights shall be tested 23 annually for proper operation on all RGDs in use. If any safety device fails, the RGD 24 be taken out of service until corrective action is performed or temporary administ 25 controls are established and approved in writing by the radiation safety officer indiv 26 responsible for radiation protection. 27 (B) Testing records shall include the date test was performed, the list of safety devices to 28 the survey instrument used, the calibration date, the results of the test, the name of 29 individual that performed the test, and any corrective actions for a failed test. 30 (C) Records of the testing shall be retained by the registrant for agency review of 31 inspection. 32 (2) Bypassing 33 (A) No individual shall bypass a safety device unless the person has obtained approval 34 the radiation safety officer individual responsible for radiation protection. Procedur 35 bypassing a safety device shall be incorporated into the radiation protection progra	20	(i) Safety Device	<u>es</u>
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31 <u>inspection.</u> 32 (2) <u>Bypassing</u> 33 (A) <u>No individual shall bypass a safety device unless the person has obtained approval 34 <u>the radiation safety officer individual responsible for radiation protection. Procedur</u> 35 <u>bypassing a safety device shall be incorporated into the radiation protection progra</u></u>	29		individual that performed the test, and any corrective actions for a failed test.
32 (2) Bypassing (A) No individual shall bypass a safety device unless the person has obtained approval the radiation safety officer individual responsible for radiation protection. Procedur bypassing a safety device shall be incorporated into the radiation protection progra	30		(C) Records of the testing shall be retained by the registrant for agency review during
No individual shall bypass a safety device unless the person has obtained approval the radiation safety officer individual responsible for radiation protection. Procedur bypassing a safety device shall be incorporated into the radiation protection progra	31		inspection.
the radiation safety officer individual responsible for radiation protection. Procedur bypassing a safety device shall be incorporated into the radiation protection progra	32	(2)	Bypassing
bypassing a safety device shall be incorporated into the radiation protection progra	33		(A) No individual shall bypass a safety device unless the person has obtained approval from
	34		the radiation safety officer individual responsible for radiation protection. Procedures for
36 <u>the radiation safety officer individual responsible for radiation protection, as set fo</u>	35		bypassing a safety device shall be incorporated into the radiation protection program by

1		Rule .1601 of this Chapter, and the operating procedures as set forth in Paragraph (c) of
2		this Rule.
3		(B) The written approval, as granted by the individual responsible for radiation protection, shall
4		include the start and end date of approval.
5		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
6		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
7		ray source housing and the control panel during the bypassing period.
8	<u>(j) <mark>Prior to an ir</mark></u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
9	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
10	<u>(1)</u>	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
11	(2)	shutters;
12	(3)	collimators; or
13	<u>(4)</u>	beam stops.
14		
15	History Note:	Authority G.S. 104E-7(a)(2);
16		Eff. February 1, 1980;
17		Amended Eff. January 1, 1994;
18		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
19		Amended Eff. October 1, 2015;
20		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
21		2019. <u>2019:</u>
22		Amended Eff. November 1, 2024.

1	10A NCAC 15.	.0805 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2		
3	10A NCAC 15	.0805 OPERATING REQUIREMENTS AREA REQUIREMENTS
4	(a) RGDs shall	be operated by individuals that have completed the training requirements of Rule .0806 of this Section.
5	(b) Normal ope	rating procedures shall be written and available to all RGD operators and support staff.
6	(c) No individ	ual shall be permitted to operate RGDs in any manner other than that specified in the operating
7	procedures unle	ss the person has obtained written approval of the individual responsible for radiation or the Radiation
8	Safety Officer (RSO) as defined in Rule .0104 of this Chapter.
9	(d) No individu	nal shall bypass a safety device unless the person has obtained the approval of the person responsible
10	for radiation saf	ety or the RSO. This process shall be incorporated into the radiation protection program by the RSO,
11	as set forth in R	ule .1603(a), and the operating procedures as set forth in Rule .0603(a)(1)(B). The written approval,
12	as granted by the	e RSO, shall include an expiration date. When a safety device has been bypassed, a legible sign bearing
13	the words "SAF	ETY DEVICE NOT WORKING" or words having a similar meaning shall be placed on the radiation
14	source housing	and the control panel during the bypassing period.
15	(e) Prior to an i	ndividual modifying the:
16	(1)	x ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
17	(2)	— shutters;
18	(3)	— collimators; or
19	(4)	— beam stops
20	the individual sl	nall determine the tube is off and will remain off until safe conditions have been restored.
21	(f) Safety devic	es including interlocks, shutters, and warning lights shall be tested once annually for proper operation
22	on all RGDs in	operation. Records of the testing shall be retained by the registrant for three years.
23	(g) No individu	al shall hold a sample or object while it is being irradiated.
24	(a) Each radiati	on area, as defined in Rule .1601(a)(3) of this Chapter, containing RGDs shall be:
25	<u>(1)</u>	conspicuously posted with caution signs, in accordance with the requirements of Rule .1601(a)(34)
26		of this Chapter, bearing the words "CAUTION - RADIATION AREA", or words having a similar
27		meaning; and
28	<u>(2)</u>	supervised continuously during operation of the RDG or shall utilize one or more of the following:
29		(A) door interlocks;
30		(B) entry monitors; or
31		(C) engineering controls.
32	(b) Access to ea	ach restricted area where an individual may receive a dose equivalent exceeding 100 mrem in any year,
33	but does not exc	eed levels of a radiation area, shall be designated as a controlled area. The area shall be controlled by:
34	(1)	visibly separating adjacent uncontrolled areas so doses do not exceed the limits of Rule .1601(a)(15)
35		of this Chapter; and
36	(2)	posting a sign bearing the words "Warning: X-rays in Use", or words having a similar meaning.

I	(c) The local co	omponents of RGDs shall be located and arranged to include sufficient shielding or access control to
2	ensure no radia	tion levels exist in any area surrounding the local components that result in a dose to an individual
3	present in exces	s of the dose limits in Rule .1601(a)(15) of this Chapter.
4	(d) Surveys sha	all be performed for each RGD, as set forth in Rule .1601(a)(23) of this Chapter, to show compliance
5	with Paragraph	(c) of this Rule.
6	<u>(1)</u>	Radiation survey instruments shall be:
7		(A) capable of measuring the radiation energies of the RGD surveyed; and
8		(B) calibrated annually when a frequency is not recommended by the manufacturer.
9	(2)	Equipment surveys shall confirm radiation levels do not exceed the requirements of Rule
10		.0806(c)(7); (d)(3); .0806(d)(3); and (h)(2) .0806(h)(2) of this Section. Surveys shall be performed:
11		(A) prior to initial use and include testing of warning and safety devices;
12		(B) prior to use following any change in the initial arrangement, including the number or type
13		of local components in the system or x-ray tube source housing;
14		(C) prior to use following any maintenance requiring the disassembly or removal of a local
15		component in the system or x-ray tube source housing that could affect the radiation
16		exposure to personnel; and
17		(D) during the performance of calibration, maintenance, or any alignment procedure if the
18		presence of a primary x-ray beam is required while any local component in the system is
19		disassembled or removed.
20	(3)	A registrant may apply to the agency for approval of procedures differing from those in
21		Subparagraph (d) of this Rule, provided that the registrant demonstrates satisfactory compliance
22		with Paragraph (c) of this Rule.
23	<u>(4)</u>	Records shall be available for agency review during inspection.
24	(e) RGDs in Ru	ale .0806(i) and .0807(2) of this Section, installed after the effective date of this Rule, shall ensure the
25	following provis	sions are met:
26	<u>(1)</u>	A floor plan with equipment arrangement shall be submitted to the agency for review and
27		acknowledgement prior to installation of the system. The floor plan shall include:
28		(A) the proposed location of the system;
29		(B) direction of the useful beam;
30		(C) adjacent areas; and
31		(D) location of the operator.
32	(2)	An area radiation survey shall be performed prior to initial use to show compliance with dose limits
33		of the rules in Section .1600 of this Chapter. The survey shall include:
34		(A) a drawing of the room indicating the location of the x-ray tube and orientation of the useful
35		beam;
36		(B) radiation levels at the operator location and adjacent areas;
37		(C) survey instrument used; and

	(D) name of the service provider that is registered registered, in accordance with Rule .0205 of
	this Chapter Chapter, and date the survey was performed.
<u>(3)</u>	Modifications to the room, RGD, or adjacent areas that may increase the radiation dose to any
	individual shall require a new survey.
<u>(4)</u>	Records of the floor plan with equipment arrangement and survey shall be available for review by
	an authorized representative of the Radiation Protection Section agency review during inspection.
History Note:	Authority G.S. 104E-7; 104E-12;
	Eff. February 1, 1980;
	Transferred and Recodified from 15A NCAC 11 .0805 Eff. February 1, 2015;
	Amended Eff. January 1, 2016; October 1, 2015;
	Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
	2019. <u>2019;</u>
	Amended Eff. November 1, 2024.
	(4)

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Kissinger, Regina < regina.kissinger@dhhs.nc.gov>

Sent: Thursday, October 10, 2024 3:30 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Thank you so much, Mr. Wiggs. We will make the changes and send them back to you.

Sincerely,

Regina Kissinger, R.T.(R) (M) ARRT Health Physicist Healing Arts Supervisor <u>Division of Health Service Regulation, Radiation Protection</u> NC Department of Health and Human Services

Office <u>919-814-2335</u> regina.kissinger@dhhs.nc.gov

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To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

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Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov>

Sent: Thursday, October 10, 2024 3:27 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Below are a few minor changes to finalize this set of rules. Please reply with the revisions.

.0802, on pg. 2, (12) line 5, please add "Rule" before "10A...."

.0803, on pg. 2, line 23, add a space after "15" that shows "15 .0104(130)."

.0804, at the end of line 29, please add "15" to the cited Rule to make it read "10A NCAC 15 .0802(17)". This is my fault as I failed to include the "15" in my last request.

.0805, on pg. 3, line 6, add "an" before "authorized".

Thanks,

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings

Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10A NCAC 15 .0803.docx; 10A NCAC 15 .0804.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Thursday, October 10, 2024 12:41 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Thanks for your quick response. Please see attached.

Thanks

1	10A NCAC 15 .0803 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0803 EQUIPMENT REQUIREMENTS PERSONNEL REQUIREMENTS
4	(a) Certified cabinet x ray systems shall meet the requirements of 21 CFR 1020.40 as incorporated by reference in
5	Rule .0117(a)(3) of this Chapter.
6	(b) All certified and certifiable cabinet x-ray systems shall:
7	(1) be constructed so that, the radiation emitted from the system shall not exceed an exposure of 0.5
8	milliroentgen (mR) in one hour at any point five centimeters outside the external surface; and
9	(2) have a fail safe interlock that prevents irradiation when the cabinet, chamber, or coupled chambers
10	are open.
11	(c) Open beam analytical RGD systems shall be equipped with a safety device that prevents the entry of any portion
12	of an individual's body into the primary x-ray beam path that causes the beam to be shut off upon entry into its path.
13	(d) Open beam analytical RGDs shall be provided with a visible and legible indication of:
14	(1) x ray tube status (ON OFF) located near the radiation source housing, if the primary beam is
15	controlled in this manner; or
16	(2) shutter status (OPEN CLOSED) or beam status (ON OFF) located near each port on the radiation
17	source housing, if the primary beam is controlled in this manner.
18	(e) Warning devices on open beam analytical RGDs shall be labeled so that their purpose is identified. On open beam
19	analytical RGDs installed after February 1, 1980, warning devices and lights shall have fail safe characteristics.
20	(f) Unused ports on radiation source housings for open-beam RGDs shall be secured in the closed position in a manner
21	that will prevent unintended opening.
22	(g) Each port on the radiation source housing on open beam analytical RGDs installed after February 1, 1980 and
23	designed to accommodate interchangeable components shall be equipped with a shutter that cannot be opened unless
24	a collimator or a component coupling is connected to the port.
25	(h) Portable open beam analytical RGDs that shall be manufactured to be used hand held without safety devices are
26	exempt from the requirements of Paragraph (c) of this Rule and shall be constructed according to International
27	Standard IEC 62495 that is incorporated by reference and includes subsequent amendments. This standard can be
28	downloaded for one hundred twenty one dollars (\$121.00) at the following website:
29	http://webstore.ansi.org/FindStandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx.aspx.aspx.aspx.aspx.aspx.aspx.asp
30	geNum=0&SearchTermsArray=null%7cIEC+62495+Ed.+1.0+en%3a2011%7cnull.
31	(i) A registrant may apply to the agency, as defined in Rule .0104 of this Chapter, for an exemption from the
32	requirement of a safety device. This request shall include:
33	(1) a description of the safety devices;
34	(2) the reason safety devices cannot be used; and
35	(3) a description of the alternative methods that will be employed to minimize the possibility of an
36	accidental exposure, including procedures to assure that operators and others in the area will be
37	informed of the absence of safety devices.

I	(j) Analytical RGDs shall be provided with a visible and legible label(s) bearing the radiation symbol and the words:
2	(1) "CAUTION HIGH INTENSITY X RAY BEAM," or words having a similar meaning, near the
3	exit port to identify the location of the beam; and
4	(2) "CAUTION RADIATION THIS EQUIPMENT PRODUCES RADIATION WHEN
5	ENERGIZED", or words having a similar meaning, near any switch that energizes an x-ray tube, if
6	the radiation source is an x-ray tube.
7	(k) Warning lights labeled with the words "X RAYS ON," or other words having similar meaning, shall be located:
8	(1) near any switch that activates the high voltage to energize an x-ray tube; or
9	(2) in a conspicuous location near the radiation source housing and radiation beam(s) and visible from
10	all instrument access areas.
11	(l) Warning lights shall activate when the x-ray tube is energized.
12	(m) Each x ray tube housing shall be:
13	(1) constructed that when all shutters are closed the leakage radiation measured at a distance of five
14	centimeters from its surface is not capable of producing an exposure in excess of 2.5 millirem
15	(mrem)/ (25 microsieverts μSv) in one hour; and
16	(2) if the tube housing is the primary shielding for the x ray tube, does not produce x rays when the
17	housing is opened or disassembled.
18	(n) Each x ray generator shall be supplied with a protection cabinet which limits leakage radiation measured at a
19	distance of five centimeters from its surface such that it is not capable of producing an exposure in excess of 0.25
20	mrem/2.5μSv in one hour.
21	(o) Permanent radiographic installations and industrial radiography RGDs shall comply with the requirements of Rule
22	.0807 of this Section.
23	(a) The registrant registrant, as defined in 10A NCAC 15.0104(130), shall document the scope of training and
24	instruction required for the RGD in use.
25	(b) No individual shall be permitted to operate or maintain RGDs unless the individual has received instruction in the
26	basic principles of radiation protection, training specific to the manufacturer's recommendations for safe operation
27	and unique features of the RGD in use, and instruction in the operating and emergency procedures. Instruction and
28	training shall include:
29	(1) Basic principles of radiation protection:
30	(A) radiation fundamentals;
31	(B) source and magnitude of common sources of radiation exposure;
32	(C) units of radiation dose and measurements;
33	(D) potential hazards, biological effects of ionizing radiation, and recognition of symptoms of
34	an acute localized exposure;
35	(E) ALARA (As Low As Reasonably Achievable) principles for radiation protection concepts
36	of time, distance, and shielding to minimize radiation exposure;
37	(F) declared pregnancy policy;

I		(G) occupational, embryo/tetus, and public dose limits; and
2		(H) proper use of individual monitoring devices and survey instruments.
3	(2)	Device specific training for each RGD:
4		(A) hands-on training for proper use;
5		(B) radiation hazards associated with use;
6		(C) precautions to take or measures required to minimize radiation exposure;
7		(D) procedures to prevent unauthorized use; and
8		(E) agency rules regarding use.
9	(3)	Operating and emergency procedure requirements of Rule .0804 in this Section.
10	(c) Records of i	nstruction and training for each individual operating RGDs, documenting that the requirements of this
11	Rule have been	met, shall be maintained and available for agency review during inspection.
12	<u>(d) <mark>Individuals</mark> j</u>	Persons who will be operating the RGD shall be able to demonstrate an understanding in safe operating
13	procedures and	use of the RGD according to the manufacturer's specifications and to an authorized representative of
14	the Radiation Pr	rotection Section.
15	(e) Each registr	ant shall provide ring or wrist individual monitoring devices to individuals:
16	<u>(1)</u>	operating open-beam RGDs; and
17	<u>(2)</u>	performing maintenance on an RDG, if the maintenance procedures require the presence of a
18		primary x-ray beam when any local component in the RGD is disassembled or removed.
19		
20	History Note:	Authority G.S. 104E-7;
21		Eff. February 1, 1980;
22		Transferred and Recodified from 15A NCAC 11 .0803 Eff. February 1, 2015;
23		Amended Eff. October 1, 2015;
24		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
25		2019. <u>2019:</u>
26		<u>Amended Eff.</u> <mark>October </mark>

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer individual responsible
29	for radiation protection as defined in 10A NCAC .0802(17). protection. as defined in Rule.0104 of this Chapter.
30	
31	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
32	all operators and support staff for review during the use of an RGD.
33	(d) Normal operating and emergency procedures shall include the following:
34	(1) safe use of the RGD;
35	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
36	<u>and</u>

any unusual occurrence or malfunction that may involve exposure to radiation. (c) Open beam and portable handheld RGDs (l) Registrants shall have operating procedures developed to ensure radiation protective measure (A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation where the primary x-ray beam is not intercepted by a detector device during operation (C) available to all individuals operating the RGD. (2) Operators shall not do the following while operating an RGD: (A) point the primary beam at any individual including him or herself; (B) allow their hand to approach the primary beam: or (C) hold a sample. If a sample is small and it is necessary to hold the sample while operating and emergency procedures shall be paleed in a shielded sample enclosure. (f) Operating and emergency procedures shall be available for review by the individual responsible for radiation protection agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the radiation safety officer india responsible for radiation protection and manufacturer of the RGD. (i) Safety Devices (i) Testing (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD testoms are established and approved in writing by the radiation safety officer india responsible for radiation protection. (B) Testing records shall include the date test was performed, the list of safety devices to the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. (C) Records of the testing shall be retained by the registrant for a	1	(3)	instructions on reporting to the radiaton safety officer individual responsible for radiation protection
(e) Open beam and portable handheld RGDs (1) Registrants shall have operating procedures developed to ensure radiation protective measure (A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation where the primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation of CO available to all individuals operating the RGD. (2) Operators shall not do the following while operating an RGD: (3) allow their hand to approach the primary beam; or (4) Bind a sample. If a sample is small and it is necessary to hold the sample while operating and emergence procedures shall be placed in a shielded sample enclosure. (b) Operating and emergence procedures shall be available for review by the individual responsible for radiation agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the-radiation-safety-officer-individual responsible for radiation protection, and manufacturer of the RGD. (i) Safety Devices (I) Testing (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD be taken out of service until corrective action is performed or temporary administ controls are established and approved in writing by the radiation-safety-officer-individual that performed the test, and any corrective action for a failed test. (C) Records of the testing shall be retained by the registrant for agency review of inspection. (C) Records of the testing shall be incorporated into the radiation protection properation bypassing a safety device shall be incorporated into the radiation protection programs.	2		and agency of actual or suspected accidental exposure or other radiation safety concerns, such as
(1) Registrants shall have operating procedures developed to ensure radiation protective measure (A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation (C) available to all individuals operating the RGD. (C) operators shall not do the following while operating an RGD: (A) point the primary beam at any individual including him or herself; (B) allow their hand to approach the primary beam; or (C) hold a sample, If a sample is small and it is necessary to hold the sample while operating and emergency procedures shall be paralled in a shielded sample enclosure. (f) Operating and emergency procedures shall be available for review by the individual responsible for radiation agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the radiation safety officer individual responsible for radiation protection and manufacturer of the RGD. (i) Safety Devices (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD be taken out of service until corrective action is performed or temporary administ controls are established and approved in writing by the radiation-safety officer individual responsible for radiation protection. (B) Testing records shall include the date test was performed, the list of safety devices in the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. (C) Records of the testing shall be retained by the registrant for agency review of inspection. (A) No individual shall bypass a safety device unless the person has obtained approval the radiation protection pro	3		any unusual occurrence or malfunction that may involve exposure to radiation.
(A) provided to meet the requirements of Rule _1601(a)(15) of this Chapter; (B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in where the primary x-ray beam is not intercepted by a detector device during operation (C) available to all individuals operating the RGD. (C) available to all individuals operating the RGD. (A) point the primary beam at any individual including him or herself; (B) allow their hand to approach the primary beam; or (C) hold a sample. If a sample is small and it is necessary to hold the sample while operating and emergency procedures shall be available for review by the individual responsible for rad protection agency review during inspection. (G) Operating and emergency procedures shall be available for review by the individual responsible for rad protection agency review during inspection. (g) Alignment procedures shall be performed as recommended by the RGD manufacturer. (h) Special alignment procedures shall only be used when approved by the radiation safety officer individes responsible for radiation protection and manufacturer of the RGD. (i) Safety Devices (ii) Safety Devices (A) Safety devices including interlocks, shutters, and warning lights shall be tested annually for proper operation on all RGDs in use. If any safety device fails, the RGD exponsible for radiation protection are established and approved in writing by the radiation safety officer individes are esponsible for radiation protection. (B) Testing records shall include the date test was performed, the list of safety devices to the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. (C) Records of the testing shall be retained by the registrant for agency review of inspection. (B) Procedure the safety officer individual responsible for radiation protection procedures to phypassing a safety device shall be incorporated into the radiation protection programas	4	(e) Open beam a	and portable handheld RGDs
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26 responsible for radiation protection. 27 (B) Testing records shall include the date test was performed, the list of safety devices to the survey instrument used, the calibration date, the results of the test, the name of individual that performed the test, and any corrective actions for a failed test. 29 individual that performed the test, and any corrective actions for a failed test. 30 (C) Records of the testing shall be retained by the registrant for agency review of inspection. 31 inspection. 32 (2) Bypassing 33 (A) No individual shall bypass a safety device unless the person has obtained approval the radiation safety officer individual responsible for radiation protection. Procedure bypassing a safety device shall be incorporated into the radiation protection program.	24		be taken out of service until corrective action is performed or temporary administrative
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31 <u>inspection.</u> 32 (2) <u>Bypassing</u> 33 (A) <u>No individual shall bypass a safety device unless the person has obtained approval 34 <u>the radiation safety officer individual responsible for radiation protection. Procedur</u> 35 <u>bypassing a safety device shall be incorporated into the radiation protection progra</u></u>	29		individual that performed the test, and any corrective actions for a failed test.
32 (2) Bypassing (A) No individual shall bypass a safety device unless the person has obtained approval the radiation safety officer individual responsible for radiation protection. Procedur bypassing a safety device shall be incorporated into the radiation protection progra	30		(C) Records of the testing shall be retained by the registrant for agency review during
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the radiation safety officer individual responsible for radiation protection. Procedur bypassing a safety device shall be incorporated into the radiation protection progra	32	(2)	Bypassing
bypassing a safety device shall be incorporated into the radiation protection progra	33		(A) No individual shall bypass a safety device unless the person has obtained approval from
	34		the radiation safety officer individual responsible for radiation protection. Procedures for
36 <u>the radiation safety officer individual responsible for radiation protection, as set fo</u>	35		bypassing a safety device shall be incorporated into the radiation protection program by

1		Rule .1601 of this Chapter, and the operating procedures as set forth in Paragraph (c) of
2		this Rule.
3		(B) The written approval, as granted by the individual responsible for radiation protection, shall
4		include the start and end date of approval.
5		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
6		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
7		ray source housing and the control panel during the bypassing period.
8	<u>(j) <mark>Prior to an ir</mark></u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
9	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
10	<u>(1)</u>	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
11	(2)	shutters;
12	(3)	collimators; or
13	<u>(4)</u>	beam stops.
14		
15	History Note:	Authority G.S. 104E-7(a)(2);
16		Eff. February 1, 1980;
17		Amended Eff. January 1, 1994;
18		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
19		Amended Eff. October 1, 2015;
20		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
21		2019. <u>2019:</u>
22		Amended Eff. November 1, 2024.

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov>

Sent: Thursday, October 10, 2024 12:26 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon,

In .0804, at the end of line 29, please add "as defined in "10A NCAC .0802(17)."

In .0803, line 23, please add ", as defined in 10A NCAC 15 .0104(130)," after "registrant".

Please reply with the revisions and I'll do a final review.

Thanks,

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings

Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10A NCAC 15 .0802.docx; 10A NCAC 15 .0804.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Thursday, October 10, 2024 9:31 AM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good morning,

We would like to keep the phrase "individual responsible for radiation protection" in .0804. Hence, we added that definition to .0802.

Please let us know your thoughts.

Thanks for your continued assistance.

10A NCAC 15 .0802 is amended with changes as published in 38:19 NCR 1219-1229 as follows:

10A NCAC 15.0802 DEFINITIONS

- In addition to terms found in Rule .0104 of this Chapter Chapter, the following definitions shall apply to this Section:
- 5 (1) "Accredited bomb squad" means a law enforcement agency utilizing certified bomb technicians.
 - (2) "Accessible surface" means the external or outside surface of the enclosure or housing provided by the manufacturer or designer of the RGD. This includes the high-voltage generator, doors, access panels, latches, control knobs, and other permanently mounted hardware, and including the plane across the exterior edge of any opening.
 - (2)(3) "Analytical RGD equipment" means equipment that uses electronic means to generate ionizing radiation for the purpose of examining the microstructure of materials, materials i.e. using direct x-ray transmission, x-ray diffraction, x-ray fluorescence, and x-ray spectroscopy.
 - (3)(4) "Analytical RGD system" means a group of local and remote components utilizing x-rays to determine the elemental composition or to examine the microstructure of materials.
 - (4) "Bomb detection RGDs" means RGDs used for the sole purpose of remotely detecting explosive devices.
 - (5) "Certified bomb technician" means a member of an accredited bomb squad who has successfully completed the FBI Hazardous Devices School. Information pertaining to this program can be found on the school website at http://www.fbi.gov/about-us/cirg/hazardous-devices.
 - (6) "Certifiable cabinet x-ray system" means an existing uncertified RGD that has been modified to meet the certification requirements specified in 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (7) "Certified cabinet x-ray system" means an RGD utilized in an enclosed, interlocked cabinet, such that the radiation machine will not operate unless all openings are securely closed. These systems shall be certified in accordance with 21 CFR 1010.2 1010.2, as incorporated by reference in Rule .0117 of this Chapter, as being manufactured and assembled pursuant to the provisions of 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (8) "Collimator" means a device or mechanism by which the x-ray beam is restricted in size.
 - (9) "Control panel" means that the part of the x-ray control where the switches, knobs, pushbuttons, and other hardware are, that are necessary located for manually setting the technique factors.
 - (10) "Electron Beam Device" means any device using electrons below 1MeV to heat, join, or otherwise irradiate materials.
 - (11) "Enclosed beam RGD" means an RGD with all possible x-ray beam paths contained in a chamber, coupled chambers, or other beam-path-confinement devices devices, to prevent any part of the body from intercepting the beam during normal operations. Normal access to the primary beam path, such as a sample chamber door, shall be interlocked with the high voltage of the x-ray tube or the shutter

1		for the beam to be considered "enclosed." An open-beam device placed in an interlocked enclosure
2		is considered an "enclosed beam" unless there are provisions for routine bypassing of the interlocks.
3	<u>(12)</u>	"Emergency procedure" means the written pre-planned steps to be taken in the event of actual or
4		suspected radiation exposure of an individual exceeding administrative or regulatory limits. limits
5		found in 10A NCAC 15.1601(a)(8) and .1601(a)(15). This procedure shall include the names and
6		telephone numbers of individuals to be contacted contacted, as well as directives for processing
7		individual monitoring devices.
8	(12) (13) "Fail-safe characteristics" means a design feature that causes the radiation beam to terminate, port
9		shutters to close, or otherwise prevents emergence of the primary beam, beam upon the failure of a
10		safety or warning device. For example, if an "X-ray On" light indicator or light indicator, shutter
11		indicator shutter indicator, or interlock fails, the radiation beam shall terminate.
12	<u>(14)</u>	"Gauging device" means a mechanism containing a source of ionizing radiation that is designed and
13		manufactured for the purpose of determining or controlling thickness, density, level, interface
14		location, or qualitative or quantitative composition of materials. It may include components such as
15		radiation shields, useful-beam controls, and other safety features in order to meet the requirements
16		or specifications. specifications of the device.
17	<u>(15)</u>	"General-use system" means a security screening system that delivers an effective dose of 25
18		microrem (0.25 microSv) or less per screening.
19	(13) (16) "Hand-held x-ray system" means any device or equipment that is portable and used for similar
20		purposes as analytical RGD equipment.
21	(14)	"Hybrid gauge" means an x-ray gauge device utilizing both x-ray and radioactive sources.
22	(15)	"Industrial radiography" means RGDs used to make radiographic images to examine the structure
23		of materials by nondestructive methods. These RGDs shall not be contained in a cabinet and are not
24		permanent installations.
25	<u>(17)</u>	"Individual responsible for radiation protection" means a person who has the knowledge and
26		responsibility to apply appropriate radiation rules, for persons registered with the agency in
27		accordance with Section .0200 of this Chapter, commensurate with the scope of the activities
28		authorized by the registrant.
29	(17) (18	"Inspection Zone" means the area established for the purpose of controlling access where screening
30		is performed. Areas controlled due to the presence of radiation shall include areas of ingress, egress,
31		gates, portals, and traffic paths. The area outside of the inspection zone shall not exceed the limits
32		of Rule .1601(a)(13) of this Section.
33	(18) (19	"Interlock" means a feature designed to prevent access to an area of radiation hazard by preventing
34		entry or by automatically removing the hazard.
35	(16)<u>(20</u>	"Ion implantation equipment, low-energy" means any enclosed device operating below 1MeV used
36		to accelerate elemental ions and implant them in other materials.
37	(17)<mark>(21</mark>	Theakage radiation" means radiation emanating from the source assembly housing except for:

l	(A) the primary beam;
2	(B) scatter radiation emanating from other components (e.g., shutter or collimator); and
3	(C) radiation produced when the beam on "beam on" switch or timer is not activated.
4	(21)(22) "Limited-use system" means a screening system that is capable of delivering an effective dose
5	greater than 25 microrem (0.25 microSv) per screening screening, but shall not exceed an effective
6	dose of 1 mrem (10 microSv) per screening, screening,
7	(18)(23) "Local components" means part of an RGD x-ray system and include areas that are struck by x-rays
8	x rays, such as radiation source housings, port and shutter assemblies, collimators, sample holders,
9	cameras, goniometers, detectors, and shielding, but do not include power supplies, transformers,
10	amplifiers, readout devices, and control panels.
11	(19)(24) "Mobile RGD" means RGD equipment mounted on a permanent base with wheels or casters for
12	moving while completely assembled.
13	(20)(25) "Normal operating procedures" means step-by-step instructions necessary to accomplish a task.
14	These procedures shall include sample insertion and manipulation, equipment alignment, routine
15	maintenance by the registrant, and data recording procedures that are related to radiation safety.
16	(21)(26) "Open-beam RGD" means a device or system designed in such a way that the primary beam is not
17	completely enclosed during normal operation operation, and when used for analysis, gauging, or
18	imaging imaging, in which an individual could accidentally place some part of their body in the
19	primary beam or stray radiation path during normal operation.
20	(22) "Permanent radiographic installation" means an RGD utilized in an enclosed shielded room, cell, or
21	vault that allows entry when the RGD is not energized.
22	(23)(27) "Portable RGD" means RGD equipment designed to be carried by hand.
23	(24)(28) "Primary beam" means radiation that passes through an aperture of the source assembly housing by
24	a direct path from the radiation source.
25	(25)(29) "Radiation generating device (RGD)" means any system, device, subsystem, or machine component
26	that may generate generate, by electronic means means, x-rays or particle radiation above 5 keV,
27	but below 1 MeV, and not used for healing arts parts on humans or animals. Examples of RGDs are
28	the following may be used as a:
29	(A) analytical RGD equipment mobile RGD;
30	(B) certified and certifiable cabinet x-ray systems portable RGD; or
31	(C) gauging devices using x-ray sources; stationary RGD.
32	(D) hybrid gauging devices;
33	(E) e beam welders;
34	(F) baggage scanners;
35	(G) industrial radiography RGDs; and
36	(H) permanent radiographic installations.

1	(26)<u>(30)</u>	"Remote components" means parts of an RGD x-ray system that are not struck by x-rays x-rays,
2		such as power supplies, transformers, amplifiers, readout devices, and control panels.
3	(30) (31)	"Safety Device" means a device, interlock or system that prevents the entry of any portion of an
4		individual's body into the primary x-ray beam or that will cause the beam to shut off upon entry into
5		its path.
6	(27)<mark>(32)</mark>	"Scattered radiation" means radiation, other than leakage radiation, that during passage through
7		matter, has been deviated in direction or has been modified by a decrease in energy.
8	(32) (33)	"Screening" means the sum of scans necessary for a security screening system to image concealed
9		objects as intended by the system design under normal operating conditions.
10	(33) (34)	"Security screening device" means a non-human use open-beam device designed for the detection
11		of contraband or weapons concealed in baggage, mail, packages, or other structures. These devices
12		include bomb detection devices used for the sole purpose of detecting explosive devices.
13	(34) (35)	"Security screening system" means a system specifically designed to detect contraband and weapons
14		concealed on a person and is used for the sole purpose of public safety and security evaluation by
15		law enforcement.
16	(28)<mark>(36)</mark>	"Shutter" means an adjustable device, generally made of lead or other high atomic number material,
17		fixed to a source assembly housing to intercept, block, or collimate the primary beam.
18	(29)<mark>(37)</mark>	"Source" means the point of origin of the radiation, such as the focal spot of an x-ray tube.
19	(30)<mark>(38)</mark>	"Stationary RGD" means RGD equipment that is installed or placed in a fixed location.
20	(31)<mark>(39)</mark>	"Stray radiation" means the sum of leakage and scatter radiation emanating from the source
21		assembly or other components components, except for the primary beam, and radiation produced
22		when the beam on switch or timer is not activated.
23	(39) (40)	"Warning device" means an audible or visible signal that warns individuals of a potential radiation
24		<u>hazard.</u>
25	(32)<mark>(41)</mark>	"X-ray generator" means the part of an x-ray system that provides the accelerating (high) voltage
26		and current for the x-ray tube.
27	(33)	"X ray gauge" means an x ray producing device designed and manufactured for the purpose of
28		detecting, measuring, gauging, or controlling thickness, density, level, or interface location of
29		manufactured products.
30	(41) (42)	"X-ray source housing" means the portion of an RGD system which contains the x-ray tube and
31		emitting target. The housing often contains radiation shielding material or inherently provides
32		shielding.
33		
34	History Note:	Authority G.S. 104E-7;
35		Eff. February 1, 1980;
36		Transferred and Recodified from 15A NCAC 11 .0802 Eff. February 1, 2015;
37		Amended Eff. October 1, 2015;

1	Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22
2	2019. <u>2019;</u>
3	Amended Eff. November 1, 2024

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15.0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer individual responsible
29	for radiation protection. as defined in Rule.0104 of this Chapter.
30	
31	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
32	all operators and support staff for review during the use of an RGD.
33	(d) Normal operating and emergency procedures shall include the following:
34	(1) safe use of the RGD;
35	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
36	<u>and</u>

1	<u>(3)</u> i	nstructions on reporting to the radiaton safety officer individual responsible for radiation protection
2	a	and agency of actual or suspected accidental exposure or other radiation safety concerns, such as
3	<u>a</u>	any unusual occurrence or malfunction that may involve exposure to radiation.
4	(e) Open beam and	d portable handheld RGDs
5	<u>(1)</u> F	Registrants shall have operating procedures developed to ensure radiation protective measures are:
6	<u>(</u>	A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter;
7	<u>(</u>	B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in cases
8		where the primary x-ray beam is not intercepted by a detector device during operation; and
9	<u>(</u>	(C) available to all individuals operating the RGD.
10	<u>(2)</u>	Operators shall not do the following while operating an RGD:
11	<u>(</u>	(A) point the primary beam at any individual including him or herself:
12	<u>(</u>	(B) allow their hand to approach the primary beam; or
13	<u>(</u>	(C) hold a sample. If a sample is small and it is necessary to hold the sample while operating
14		the RGD, the sample shall be placed in a shielded sample enclosure.
15	(f) Operating and	emergency procedures shall be available for review by the individual responsible for radiation
16	protection agency 1	review during inspection.
17	(g) Alignment pro	cedures shall be performed as recommended by the RGD manufacturer.
18	(h) Special align	ment procedures shall only be used when approved by the radiation safety officer individual
19	responsible for rad	iation protection and manufacturer of the RGD.
20	(i) Safety Devices	
21	<u>(1)</u>	<u> Testing</u>
22	<u>(</u>	(A) Safety devices including interlocks, shutters, and warning lights shall be tested once
23		annually for proper operation on all RGDs in use. If any safety device fails, the RGD shall
24		be taken out of service until corrective action is performed or temporary administrative
25		controls are established and approved in writing by the radiation safety officer individual
26		responsible for radiation protection.
27	<u>(</u>	B) Testing records shall include the date test was performed, the list of safety devices tested,
28		the survey instrument used, the calibration date, the results of the test, the name of the
29		individual that performed the test, and any corrective actions for a failed test.
30	<u>(</u>	(C) Records of the testing shall be retained by the registrant for agency review during
31		inspection.
	<u>(2)</u> <u>H</u>	Bypassing
32	(<u>yypassing</u>
32 33	-	A) No individual shall bypass a safety device unless the person has obtained approval from
	-	
33	•	(A) No individual shall bypass a safety device unless the person has obtained approval from

1		Rule .1601 of this Chapter, and the operating procedures as set forth in Paragraph (c) of
2		this Rule.
3		(B) The written approval, as granted by the individual responsible for radiation protection, shall
4		include the start and end date of approval.
5		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
6		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
7		ray source housing and the control panel during the bypassing period.
8	<u>(j) <mark>Prior to an ir</mark></u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
9	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
10	<u>(1)</u>	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
11	(2)	shutters;
12	(3)	collimators; or
13	<u>(4)</u>	beam stops.
14		
15	History Note:	Authority G.S. 104E-7(a)(2);
16		Eff. February 1, 1980;
17		Amended Eff. January 1, 1994;
18		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
19		Amended Eff. October 1, 2015;
20		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
21		2019. <u>2019:</u>
22		Amended Eff. November 1, 2024.

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov> **Sent:** Wednesday, October 9, 2024 5:16 PM

To: Kissinger, Regina <regina.kissinger@dhhs.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon, Ms. Kissinger. It's unclear to me who "the registrant" is in this context. I understand you're using the term as is done in .0803, but the Rule needs to

be specific about what "the registrant" is registering for. I'm fine with using the term once this detail is added.

Continued best wishes for you.

Thanks,

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

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Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov> **Sent:** Wednesday, October 9, 2024 5:16 PM

To: Kissinger, Regina <regina.kissinger@dhhs.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

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Continued best wishes for you.

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Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings Telephone: 984-236-1929

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Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10A NCAC 15 .0804 (004).docx

From: Kissinger, Regina < regina.kissinger@dhhs.nc.gov>

Sent: Wednesday, October 9, 2024 4:53 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon,

I hope each of you is doing well and thank you for the recent well wishes. It sure is different here than it was two weeks ago, and the extent of damage was never in our wildest dreams...

First, I want to thank all of you for your part in and time spent reviewing this rule.

Mr. Wiggs, thank you for recommending the finalized rules for approval. I do have a question for you. I contacted members of the Radiation Protection Commission, and they decided to use the wording "registrant" rather than "an authorized representative of the Radiation Protection Section." Hopefully, that will be acceptable. I have attached the document with the change. Please let me know at your convenience if that is acceptable.

Thank you again,

Regina Kissinger, R.T.(R) (M) ARRT Health Physicist Healing Arts Supervisor Division of Health Service Regulation, Radiation Protection NC Department of Health and Human Services

Office <u>919-814-2335</u> regina.kissinger@dhhs.nc.gov

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1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer the registrant.
29	defined in Rule.0104 of this Chapter.
30	
31	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
32	all operators and support staff for review during the use of an RGD.
33	(d) Normal operating and emergency procedures shall include the following:
34	(1) safe use of the RGD;
35	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
36	<u>and</u>

1	(3)	instruct	<u>ions on reporting to <mark>the</mark> radiaton safety officer <mark>individual responsible for radiation protection</mark></u>
2		and age	ney an authorized representative of the Radiation Protection Section of actual or suspected
3		acciden	tal exposure or other radiation safety concerns, such as any unusual occurrence or
4		malfunc	ction that may involve exposure to radiation.
5	(e) Open beam a	and portal	ble handheld RGDs
6	<u>(1)</u>	Registra	ants shall have operating procedures developed to ensure radiation protective measures are:
7		(A)	provided to meet the requirements of Rule .1601(a)(15) of this Chapter;
8		(B)	taken to avoid exposure to any individual from the transmitted primary x-ray beam in cases
9			where the primary x-ray beam is not intercepted by a detector device during operation; and
10		(C)	available to all individuals operating the RGD.
11	(2)	Operato	ors shall not do the following while operating an RGD:
12		(A)	point the primary beam at any individual including him or herself;
13		<u>(B)</u>	allow their hand to approach the primary beam; or
14		<u>(C)</u>	hold a sample. If a sample is small and it is necessary to hold the sample while operating
15			the RGD, the sample shall be placed in a shielded sample enclosure.
16	(f) Operating and	d emerge	ncy procedures shall be available for <mark>review by an authorized representative of the Radiation</mark>
17	Protection Section	<u>n <mark>agency</mark></u>	review during inspection.
18	(g) Alignment p	rocedure	s shall be performed as recommended by the RGD manufacturer.
19	(h) Special alig	gnment p	rocedures shall only be used when approved by the radiation safety officer individual
20	responsible for re	adiation p	rotection an authorized representative of the Radiation Protection Section and manufacturer
21	of the RGD.		
22	(i) Safety Device	<u>es</u>	
23	(1)	Testing	
24		(A)	Safety devices including interlocks, shutters, and warning lights shall be tested once
25			annually for proper operation on all RGDs in use. If any safety device fails, the RGD shall
26			be taken out of service until corrective action is performed or temporary administrative
27			controls are established and approved in writing by the radiation safety officer individual
28			responsible for radiation protection an authorized representative of the Radiation
29			Protection Section.
30		(B)	Testing records shall include the date test was performed, the list of safety devices tested,
31			the survey instrument used, the calibration date, the results of the test, the name of the
32			individual that performed the test, and any corrective actions for a failed test.
33		(C)	Records of the testing shall be retained by the registrant for agency review during
34			inspection.
35	(2)	Bypassi	ng
36		<u>(A)</u>	No individual shall bypass a safety device unless the person has obtained approval from
37			the radiation safety officer individual responsible for radiation protection, an authorized

I		representative of the Radiation Protection Section. Procedures for bypassing a safety
2		device shall be incorporated into the radiation protection program by the radiation safety
3		officer individual responsible for radiation protection, an authorized representative of the
4		Radiation Protection Section, as set forth in Rule .1601 of this Chapter, and the operating
5		procedures as set forth in Paragraph (c) of this Rule.
6		(B) The written approval, as granted by the individual responsible for radiation protection, an
7		authorized representative of the Radiation Protection Section, shall include the start and
8		end date of approval.
9		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
10		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
11		ray source housing and the control panel during the bypassing period.
12	<u>(j) <mark>Prior to an ir</mark></u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
13	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
14	(1)	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
15	(2)	shutters;
16	(3)	collimators; or
17	(4)	beam stops.
18		
19	History Note:	Authority G.S. 104E-7(a)(2);
20		Eff. February 1, 1980;
21		Amended Eff. January 1, 1994;
22		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
23		Amended Eff. October 1, 2015;
24		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
25		2019. <u>2019:</u>
26		Amended Eff. November 1, 2024.

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Black, Shanah <shanah.black@dhhs.nc.gov> Sent: Wednesday, October 9, 2024 6:01 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: Re: Radiation Protection Commission Rules-October 2024

Thank you for the clarification. We will get back to you hopefully tomorrow with revisions

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From: Wiggs, Travis C < travis.wiggs@oah.nc.gov>
Sent: Wednesday, October 9, 2024 5:15:58 PM

To: Kissinger, Regina <regina.kissinger@dhhs.nc.gov>; Black, Shanah <shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon, Ms. Kissinger. It's unclear to me who "the registrant" is in this context. I understand you're using the term as is done in .0803, but the Rule needs to

be specific about what "the registrant" is registering for. I'm fine with using the term once this detail is added.

Continued best wishes for you.

Thanks,

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings

Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

From: Kissinger, Regina < regina.kissinger@dhhs.nc.gov>

Sent: Wednesday, October 9, 2024 4:53 PM

To: Wiggs, Travis C < travis.wiggs@oah.nc.gov>; Black, Shanah < shanah.black@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon,

I hope each of you is doing well and thank you for the recent well wishes. It sure is different here than it was two weeks ago, and the extent of damage was never in our wildest dreams...

First, I want to thank all of you for your part in and time spent reviewing this rule.

Mr. Wiggs, thank you for recommending the finalized rules for approval. I do have a question for you. I contacted members of the Radiation Protection Commission, and they decided to use the wording "registrant" rather than "an authorized representative of the Radiation Protection Section." Hopefully, that will be acceptable. I have attached the document with the change. Please let me know at your convenience if that is acceptable.

Thank you again,

Regina Kissinger, R.T.(R) (M) ARRT Health Physicist Healing Arts Supervisor <u>Division of Health Service Regulation</u>, <u>Radiation Protection</u> NC Department of Health and Human Services

Office <u>919-814-2335</u> regina.kissinger@dhhs.nc.gov

5505 Creedmoor Road, Suite 100, One Crosspointe Plaza 1645 Mail Service Center Raleigh NC 27699-1600

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From: Wiggs, Travis C < travis.wiggs@oah.nc.gov>
Sent: Wednesday, October 9, 2024 3:59 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Thank you both for your quick responses.

I intend to recommend to the RRC that all the final revised rules be approved at the October 30th meeting. Please submit all revised rules via email to <u>oah.rules@oah.nc.gov</u> no later than 5pm on October 18, 2024. The electronic copy must be saved as the official rule name (XX NCAC XXXX). Please include me on the email.

Thank you.

Travis C. Wiggs
Rules Review Commission Counsel
Office of Administrative Hearings

Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10A NCAC 15 .0804.docx; 10A NCAC 15 .0801.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Wednesday, October 9, 2024 2:29 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Hey Travis,

Thank you so much for your help on these rules.

Please see the 2 rules you advised revisions on and let us know your thoughts.

Thanks again!

1	10A NCAC 15	.0801 is amended with changes as published in 38:19 NCR 1219-1229 as follows:		
2	SECTION .0800 - REQUIREMENTS FOR NON-HUMAN USE OF RADIATION GENERATING			
3		DEVICES		
4				
5	10A NCAC 15	.0801 PURPOSE AND SCOPE		
6	(a) This Section	n provides special additional requirements for use of ionizing radiation generating devices (RGDs)		
7	operating above	five thousand electron volts (5 keV), but below one million electron volts (1 MeV) that are in addition		
8	to (1 MeV). Th	e requirements in of this Section are in addition to the provisions in other sections Sections .0100,		
9	.0200, .1000, ar	d .1600 of this Chapter.		
10	(b) This Section	n does not pertain to radiation safety requirements for x-ray equipment that is covered in other sections		
11	industrial radio	graphic machines for non-human use that are covered in Section .0500 of this Chapter Chapter, (e.g.,		
12	x-rays in the he	aling arts in Section .0600 of this Chapter, and particle accelerators in Section .0900 of this Chapter).		
13	Chapter.			
14	(c) RGDs used	for the purpose of elemental analysis, microstructural analysis, quality assurance, quality control,		
15	research and dev	velopment, gauging and measurement, or other nondestructive testing and evaluation RGDs addressed		
16	in this Section i	nelude: in this Section includes:		
17	<u>(1)</u>	analytical RGDs;		
18	(2)	cabinet x-ray systems;		
19	(3)	electron beam devices operating below 1MeV;		
20	<u>(4)</u>	electron microscopes;		
21	<u>(5)</u>	ion implantation equipment, low energy;		
22	<u>(6)</u>	gauging devices;		
23	<u>(7)</u>	radiographic and radioscopic non-healing arts x-ray equipment; and		
24	<u>(8)</u>	security screening devices and systems for government use only.		
25				
26	History Note:	Authority G.S. 104E-7;		
27		Eff. February 1, 1980;		
28		Transferred and Recodified from 15A NCAC 11 .0801 Eff. February 1, 2015;		
29		Amended Eff. October 1, 2015;		
30		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,		
31		2019. <u>2019:</u>		
32		Amended Eff. November 1, 2024.		

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	40.4 NG 4 G 47, 000.4
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer an authorized
29	representative of the Radiation Protection Section. as defined in Rule.0104 of this Chapter.
30	
31	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
32	all operators and support staff for review during the use of an RGD.
33	(d) Normal operating and emergency procedures shall include the following:
34	(1) safe use of the RGD;
35	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
36	<u>and</u>

1	(3)	instruct	<u>ions on reporting to <mark>the</mark> radiaton safety officer <mark>individual responsible for radiation protection</mark></u>
2		and age	ney an authorized representative of the Radiation Protection Section of actual or suspected
3		acciden	tal exposure or other radiation safety concerns, such as any unusual occurrence or
4		malfunc	ction that may involve exposure to radiation.
5	(e) Open beam a	and portal	ble handheld RGDs
6	<u>(1)</u>	Registra	ants shall have operating procedures developed to ensure radiation protective measures are:
7		(A)	provided to meet the requirements of Rule .1601(a)(15) of this Chapter;
8		(B)	taken to avoid exposure to any individual from the transmitted primary x-ray beam in cases
9			where the primary x-ray beam is not intercepted by a detector device during operation; and
10		(C)	available to all individuals operating the RGD.
11	(2)	Operato	ors shall not do the following while operating an RGD:
12		(A)	point the primary beam at any individual including him or herself;
13		<u>(B)</u>	allow their hand to approach the primary beam; or
14		<u>(C)</u>	hold a sample. If a sample is small and it is necessary to hold the sample while operating
15			the RGD, the sample shall be placed in a shielded sample enclosure.
16	(f) Operating and	d emerge	ncy procedures shall be available for <mark>review by an authorized representative of the Radiation</mark>
17	Protection Section	<u>n <mark>agency</mark></u>	review during inspection.
18	(g) Alignment p	rocedure	s shall be performed as recommended by the RGD manufacturer.
19	(h) Special alig	gnment p	rocedures shall only be used when approved by the radiation safety officer individual
20	responsible for re	adiation p	rotection an authorized representative of the Radiation Protection Section and manufacturer
21	of the RGD.		
22	(i) Safety Device	<u>es</u>	
23	(1)	Testing	
24		(A)	Safety devices including interlocks, shutters, and warning lights shall be tested once
25			annually for proper operation on all RGDs in use. If any safety device fails, the RGD shall
26			be taken out of service until corrective action is performed or temporary administrative
27			controls are established and approved in writing by the radiation safety officer individual
28			responsible for radiation protection an authorized representative of the Radiation
29			Protection Section.
30		(B)	Testing records shall include the date test was performed, the list of safety devices tested,
31			the survey instrument used, the calibration date, the results of the test, the name of the
32			individual that performed the test, and any corrective actions for a failed test.
33		<u>(C)</u>	Records of the testing shall be retained by the registrant for agency review during
34			inspection.
35	<u>(2)</u>	Bypassi	ng
36		(A)	No individual shall bypass a safety device unless the person has obtained approval from
37			the radiation safety officer individual responsible for radiation protection, an authorized

I		representative of the Radiation Protection Section. Procedures for bypassing a safety
2		device shall be incorporated into the radiation protection program by the radiation safety
3		officer individual responsible for radiation protection, an authorized representative of the
4		Radiation Protection Section, as set forth in Rule .1601 of this Chapter, and the operating
5		procedures as set forth in Paragraph (c) of this Rule.
6		(B) The written approval, as granted by the individual responsible for radiation protection, an
7		authorized representative of the Radiation Protection Section, shall include the start and
8		end date of approval.
9		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
10		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
11		ray source housing and the control panel during the bypassing period.
12	<u>(j) <mark>Prior to an ir</mark></u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
13	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
14	(1)	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
15	(2)	shutters;
16	(3)	collimators; or
17	(4)	beam stops.
18		
19	History Note:	Authority G.S. 104E-7(a)(2);
20		Eff. February 1, 1980;
21		Amended Eff. January 1, 1994;
22		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
23		Amended Eff. October 1, 2015;
24		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
25		2019. <u>2019:</u>
26		Amended Eff. November 1, 2024.

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov> Sent: Wednesday, October 9, 2024 12:55 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon,

In .0801, now line 16, please capitalize "Section."

- In .0804(c), it seems the key difference between using "individual responsible for radiation protection" in this Rule, versus 10A NCAC 15 .0203(a)(2), is .0203 requires the responsible individual to be listed on an application for the person to be vetted during the application process. Here, the "individual responsible for radiation protection" is not currently defined and it is unclear how to identify this person. Your agency may want to consider how this person will be qualified to approve use of an RGD outside of normal operating procedures.
- You can replace "individual responsible for radiation protection" with "an authorized representative of the Radiation Protection Section." Or, add the definition you intend to use in Phase 7, in Rule 10A NCAC 15 .0802, during this phase of rules.

I'm otherwise satisfied with the changes. Please reply at your earliest convenience. Thank you.

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings

Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Subject:

Attachments: Second revisions RPC phase 6.docx; 10A NCAC 15 .0801.docx; 10A NCAC 15 .0802.docx;

FW: Radiation Protection Commission Rules-October 2024

10A NCAC 15 .0803.docx; 10A NCAC 15 .0804.docx; 10A NCAC 15 .0805.docx; 10A

NCAC 15 .0806.docx; 10A NCAC 15 .0807.docx; 10A NCAC 15 .0808.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Wednesday, October 9, 2024 8:52 AM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good morning,

I just got off the phone with Regina. She is working today but this past weekend was just spent getting water and food to her elderly parents and helping others. It is truly devastating in Western NC.

Thankfully, she got power yesterday. We have revisions for all of the rules you cited, she is still struggling with one rule.

Please see attached and let us know your comments. Any advice you can give for .804 is appreciated.

Thanks for your assistance and patience.

.0801:

• In (c), my suggested change was to make this subsection one complete sentence ("RGDs used...and evaluation in this Section include:"). I suggested the change since complete sentences were used in (a) and (b). The way it has been changed leaves the first sentence as incomplete. I'm satisfied either way you want to do it.

Response: Item addressed in the text.

.0802:

- In (20)(C), based on your response, consider putting quotation marks around "beam on", similar to how "X-ray On" or "enclosed beam" was done in other sections of this Rule.

 Response: Item addressed in the text.
- In (21), line 2, add a comma after "screening". It appears that some change was made to "screening" on line 3 after my initial request to change.

Response: Item addressed in the text.

.0803:

• On pg. 3, line 11, you indicated this process will be defined in "upcoming rule 10A NCAC 15 .0104(b)(1)". What is your intended timeline to amend this exisiting Rule to include the definition? How will the regulated public know how to "demonstrate an understanding in safe operating procedures and use" if the upcoming rule is not codified prior to codification of .0803?

Response: Item addressed, changed from individual to persons operating. This is the way it has been described in existing rules. It is found in Rule 104(e)-5(11).

• My third request for this Rule was intended for a different rule. Please disregard.

.0804:

• For the phrase "individual responsible for radiation protection", you indicated this is planning to be defined in forthcoming Rule 10A NCAC 15. 0104(b)(4). How will the regulated public know who how to identify this individual if .0104(b)(4) is not codified prior to codification of this Rule? 10A NCAC 15.0203(a) says unregistered radiation machines or facilities shall "designate on the application form an individual who shall be responsible for radiation protection." How does this Rule, and any other exisiting rules, ensure the "individual responsible for radiation protection" is qualified to approve an individual to "operate an RGD in any manner other than that specified in the operating procedures"?

Response: She is struggling with this. Can it be defined in more than one section? These

were never defined in the past, it is coming out in phases. Phase 7 has it defined as Individual responsible for radiation protection" means a person who has the knowledge and responsibility to apply appropriate radiation rules, for persons registered with the agency in accordance with Section .0200 of this Chapter, commensurate with the scope of the activities authorized by the registrant.

 On pg. 2, line 15 and 29, consider saying "shall be available for review by an authorized respresentative of the Radiation Protection Section during inspection."

Response: Item addressed in the text.

.0805:

• On pg. 3, lines 5-6, consider saying "shall be available for review by an authorized representative of the Radiation Protection Section during inspection."

Response: Item addressed in the text.

.0806:

• In (c), lines 22-23, in the added language, add a comma after "requirements" and capitalize "rule".

Response: Item addressed in the text.

.0807:

- In line 33, what is meant by "shall be provided"? Is the operator providing this or someone else? Or is the operator required to "implement" some additional measure of protection?

 Response: Item addressed in the text.
- In (11), lines 3-4, consider saying "shall be available for review by an authorized representative of the Radiation Protection Section during inspection."
 Response: Item addressed in the text.

<u>:8080.</u>

• In (a), lines 22-23, it appears an extra "review" was added during the revision. The request would have it read as "used prior to the agency completing a review of information...."

Response: Item addressed in the text.

1	10A NCAC 15	.0801 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	SECTIO	N .0800 - REQUIREMENTS FOR NON-HUMAN USE OF RADIATION GENERATING
3		DEVICES
4		
5	10A NCAC 15	.0801 PURPOSE AND SCOPE
6	(a) This Section	n provides special additional requirements for use of ionizing radiation generating devices (RGDs)
7	operating above	five thousand electron volts (5 keV), but below one million electron volts (1 MeV) that are in addition
8	to (1 MeV). Th	e requirements in of this Section are in addition to the provisions in other sections Sections .0100,
9	.0200, .1000, ar	ad .1600 of this Chapter.
10	(b) This Section	n does not pertain to radiation safety requirements for x-ray equipment that is covered in other sections
11	industrial radio	graphic machines for non-human use that are covered in Section .0500 of this Chapter Chapter, (e.g.,
12	x-rays in the he	aling arts in Section .0600 of this Chapter, and particle accelerators in Section .0900 of this Chapter).
13	Chapter.	
14	(c) RGDs used	I for the purpose of elemental analysis, microstructural analysis, quality assurance, quality control,
15	research and de	velopment, gauging and measurement, or other nondestructive testing and evaluation RGDs addressed
16	in this Section i	nelude: in this section includes:
17	(1)	analytical RGDs;
18	<u>(2)</u>	cabinet x-ray systems;
19	(3)	electron beam devices operating below 1MeV;
20	<u>(4)</u>	electron microscopes;
21	<u>(5)</u>	ion implantation equipment, low energy;
22	<u>(6)</u>	gauging devices;
23	<u>(7)</u>	radiographic and radioscopic non-healing arts x-ray equipment; and
24	(8)	security screening devices and systems for government use only.
25		
26	History Note:	Authority G.S. 104E-7;
27		Eff. February 1, 1980;
28		Transferred and Recodified from 15A NCAC 11 .0801 Eff. February 1, 2015;
29		Amended Eff. October 1, 2015;
30		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
31		2019. <u>2019;</u>
32		Amended Eff. November 1, 2024.

10A NCAC 15 .0802 is amended with changes as published in 38:19 NCR 1219-1229 as follows:

10A NCAC 15.0802 DEFINITIONS

- In addition to terms found in Rule .0104 of this Chapter Chapter, the following definitions shall apply to this Section:
- 5 (1) "Accredited bomb squad" means a law enforcement agency utilizing certified bomb technicians.
 - (2) "Accessible surface" means the external or outside surface of the enclosure or housing provided by the manufacturer or designer of the RGD. This includes the high-voltage generator, doors, access panels, latches, control knobs, and other permanently mounted hardware, and including the plane across the exterior edge of any opening.
 - (2)(3) "Analytical RGD equipment" means equipment that uses electronic means to generate ionizing radiation for the purpose of examining the microstructure of materials, materials i.e. using direct x-ray transmission, x-ray diffraction, x-ray fluorescence, and x-ray spectroscopy.
 - (3)(4) "Analytical RGD system" means a group of local and remote components utilizing x-rays to determine the elemental composition or to examine the microstructure of materials.
 - (4) "Bomb detection RGDs" means RGDs used for the sole purpose of remotely detecting explosive devices.
 - (5) "Certified bomb technician" means a member of an accredited bomb squad who has successfully completed the FBI Hazardous Devices School. Information pertaining to this program can be found on the school website at http://www.fbi.gov/about-us/cirg/hazardous-devices.
 - (6) "Certifiable cabinet x-ray system" means an existing uncertified RGD that has been modified to meet the certification requirements specified in 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (7) "Certified cabinet x-ray system" means an RGD utilized in an enclosed, interlocked cabinet, such that the radiation machine will not operate unless all openings are securely closed. These systems shall be certified in accordance with 21 CFR 1010.2 1010.2, as incorporated by reference in Rule .0117 of this Chapter, as being manufactured and assembled pursuant to the provisions of 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (8) "Collimator" means a device or mechanism by which the x-ray beam is restricted in size.
 - (9) "Control panel" means that the part of the x-ray control where the switches, knobs, pushbuttons, and other hardware are, that are necessary located for manually setting the technique factors.
 - (10) "Electron Beam Device" means any device using electrons below 1MeV to heat, join, or otherwise irradiate materials.
 - (11) "Enclosed beam RGD" means an RGD with all possible x-ray beam paths contained in a chamber, coupled chambers, or other beam-path-confinement devices devices, to prevent any part of the body from intercepting the beam during normal operations. Normal access to the primary beam path, such as a sample chamber door, shall be interlocked with the high voltage of the x-ray tube or the shutter

1		for the beam to be considered "enclosed." An open-beam device placed in an interlocked enclosure
2		is considered an "enclosed beam" unless there are provisions for routine bypassing of the interlocks.
3	<u>(12)</u>	"Emergency procedure" means the written pre-planned steps to be taken in the event of actual or
4		suspected radiation exposure of an individual exceeding administrative or regulatory limits-limits
5		found in 10A NCAC 15 .1601(a)(8) and .1601(a)(15). This procedure shall include the names and
6		telephone numbers of individuals to be contacted contacted, as well as directives for processing
7		individual monitoring devices.
8	(12) (13) "Fail-safe characteristics" means a design feature that causes the radiation beam to terminate, port
9		shutters to close, or otherwise prevents emergence of the primary beam, beam upon the failure of a
10		safety or warning device. For example, if an "X-ray On" light indicator or light indicator, shutter
11		indicator shutter indicator, or interlock fails, the radiation beam shall terminate.
12	<u>(14)</u>	"Gauging device" means a mechanism containing a source of ionizing radiation that is designed and
13		manufactured for the purpose of determining or controlling thickness, density, level, interface
14		location, or qualitative or quantitative composition of materials. It may include components such as
15		radiation shields, useful-beam controls, and other safety features in order to meet the requirements
16		or specifications. specifications of the device.
17	<u>(15)</u>	"General-use system" means a security screening system that delivers an effective dose of 25
18		microrem (0.25 microSv) or less per screening.
19	(13) (16) "Hand-held x-ray system" means any device or equipment that is portable and used for similar
20		purposes as analytical RGD equipment.
21	(14)	"Hybrid gauge" means an x-ray gauge device utilizing both x-ray and radioactive sources.
22	(15)	"Industrial radiography" means RGDs used to make radiographic images to examine the structure
23		of materials by nondestructive methods. These RGDs shall not be contained in a cabinet and are not
24		permanent installations.
25	<u>(17)</u>	"Inspection Zone" means the area established for the purpose of controlling access where screening
26		is performed. Areas controlled due to the presence of radiation shall include areas of ingress, egress,
27		gates, portals, and traffic paths. The area outside of the inspection zone shall not exceed the limits
28		of Rule .1601(a)(13) of this Section.
29	<u>(18)</u>	"Interlock" means a feature designed to prevent access to an area of radiation hazard by preventing
30		entry or by automatically removing the hazard.
31	(16) (19) "Ion implantation equipment, low-energy" means any enclosed device operating below 1MeV used
32		to accelerate elemental ions and implant them in other materials.
33	(17) (20) "Leakage radiation" means radiation emanating from the source assembly housing except for:
34		(A) the primary beam;
35		(B) scatter radiation emanating from other components (e.g., shutter or collimator); and
36		(C) radiation produced when the beam on "beam on" switch or timer is not activated.

1	<u>(21)</u>	"Limited-use system" means a screening system that is capable of delivering an effective dose
2		greater than 25 microrem (0.25 microSv) per screening, but shall not exceed an effective
3		dose of 1 mrem (10 microSv) per sereening. screening.
4	(18) (22)	"Local components" means part of an RGD x-ray system and include areas that are struck by x-rays
5		<u>x rays.</u> such as radiation source housings, port and shutter assemblies, collimators, sample holders,
6		cameras, goniometers, detectors, and shielding, but do not include power supplies, transformers,
7		amplifiers, readout devices, and control panels.
8	(19) (23)	"Mobile RGD" means RGD equipment mounted on a permanent base with wheels or casters for
9		moving while <u>completely</u> assembled.
10	(20) (24)	"Normal operating procedures" means step-by-step instructions necessary to accomplish a task.
11		These procedures shall include sample insertion and manipulation, equipment alignment, routine
12		maintenance by the registrant, and data recording procedures that are related to radiation safety.
13	(21) (25)	"Open-beam RGD" means a device or system designed in such a way that the primary beam is not
14		completely enclosed during normal operation operation, and when used for analysis, gauging, or
15		imaging imaging, in which an individual could accidentally place some part of their body in the
16		primary beam or stray radiation path during normal operation.
17	(22)	"Permanent radiographic installation" means an RGD utilized in an enclosed shielded room, cell, or
18		vault that allows entry when the RGD is not energized.
19	(23) (26)	"Portable RGD" means RGD equipment designed to be carried of carried by hand.
20	(24) (27)	"Primary beam" means radiation that passes through an aperture of the source assembly housing by
21		a direct path from the radiation source.
22	(25) (28)	"Radiation generating device (RGD)" means any system, device, subsystem, or machine component
23		that may generate generate, by electronic means means, x-rays or particle radiation above 5 keV,
24		but below 1 MeV, and not used for healing arts parts on humans or animals. Examples of RGDs are
25		the following may be used as a:
26		(A) analytical RGD equipment mobile RGD;
27		(B) certified and certifiable cabinet x-ray systems portable RGD; or
28		(C) gauging devices using x-ray sources; stationary RGD.
29		(D) hybrid gauging devices;
30		(E) e beam welders;
31		(F) baggage scanners;
32		(G) industrial radiography RGDs; and
33		(H) permanent radiographic installations.
34	(26) (29)	"Remote components" means parts of an RGD x-ray system that are not struck by $\frac{x - rays}{x}$
35		such as power supplies, transformers, amplifiers, readout devices, and control panels.

I	<u>(30)</u>	"Safety Device" means a device, interlock or system that prevents the entry of any portion of an
2		individual's body into the primary x-ray beam or that will cause the beam to shut off upon entry into
3		its path.
4	(27) (31)	"Scattered radiation" means radiation, other than leakage radiation, that during passage through
5		matter, has been deviated in direction or has been modified by a decrease in energy.
6	<u>(32)</u>	"Screening" means the sum of scans necessary for a security screening system to image concealed
7		objects as intended by the system design under normal operating conditions.
8	<u>(33)</u>	"Security screening device" means a non-human use open-beam device designed for the detection
9		of contraband or weapons concealed in baggage, mail, packages, or other structures. These devices
10		include bomb detection devices used for the sole purpose of detecting explosive devices.
11	<u>(34)</u>	"Security screening system" means a system specifically designed to detect contraband and weapons
12		concealed on a person and is used for the sole purpose of public safety and security evaluation by
13		law enforcement.
14	(28) (35)	"Shutter" means an adjustable device, generally made of lead or other high atomic number material,
15		fixed to a source assembly housing to intercept, block, or collimate the primary beam.
16	(29) (36)	"Source" means the point of origin of the radiation, such as the focal spot of an x-ray tube.
17	(30) (37)	"Stationary RGD" means RGD equipment that is installed or placed in a fixed location.
18	(31) (38)	"Stray radiation" means the sum of leakage and scatter radiation emanating from the source
19		assembly or other ecomponents components, except for the primary beam, and radiation produced
20		when the beam on switch or timer is not activated.
21	<u>(39)</u>	"Warning device" means an audible or visible signal that warns individuals of a potential radiation
22		<u>hazard.</u>
23	(32) (40)	"X-ray generator" means the part of an x-ray system that provides the accelerating (high) voltage
24		and current for the x-ray tube.
25	(33)	"X ray gauge" means an x ray producing device designed and manufactured for the purpose of
26		detecting, measuring, gauging, or controlling thickness, density, level, or interface location of
27		manufactured products.
28	<u>(41)</u>	"X-ray source housing" means the portion of an RGD system which contains the x-ray tube and
29		emitting target. The housing often contains radiation shielding material or inherently provides
30		shielding.
31		
32	History Note:	Authority G.S. 104E-7;
33		Eff. February 1, 1980;
34		Transferred and Recodified from 15A NCAC 11 .0802 Eff. February 1, 2015;
35		Amended Eff. October 1, 2015;
36		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
37		2019. <u>2019;</u>

Amended Eff. November 1, 2024.

1

1	10A NCAC 15 .0803 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0803 EQUIPMENT REQUIREMENTS PERSONNEL REQUIREMENTS
4	(a) Certified cabinet x ray systems shall meet the requirements of 21 CFR 1020.40 as incorporated by reference in
5	Rule .0117(a)(3) of this Chapter.
6	(b) All certified and certifiable cabinet x-ray systems shall:
7	(1) be constructed so that, the radiation emitted from the system shall not exceed an exposure of 0.5
8	milliroentgen (mR) in one hour at any point five centimeters outside the external surface; and
9	(2) have a fail safe interlock that prevents irradiation when the cabinet, chamber, or coupled chambers
10	are open.
11	(c) Open beam analytical RGD systems shall be equipped with a safety device that prevents the entry of any portion
12	of an individual's body into the primary x-ray beam path that causes the beam to be shut off upon entry into its path.
13	(d) Open beam analytical RGDs shall be provided with a visible and legible indication of:
14	(1) x ray tube status (ON OFF) located near the radiation source housing, if the primary beam is
15	controlled in this manner; or
16	(2) shutter status (OPEN CLOSED) or beam status (ON OFF) located near each port on the radiation
17	source housing, if the primary beam is controlled in this manner.
18	(e) Warning devices on open beam analytical RGDs shall be labeled so that their purpose is identified. On open beam
19	analytical RGDs installed after February 1, 1980, warning devices and lights shall have fail safe characteristics.
20	(f) Unused ports on radiation source housings for open-beam RGDs shall be secured in the closed position in a manner
21	that will prevent unintended opening.
22	(g) Each port on the radiation source housing on open beam analytical RGDs installed after February 1, 1980 and
23	designed to accommodate interchangeable components shall be equipped with a shutter that cannot be opened unless
24	a collimator or a component coupling is connected to the port.
25	(h) Portable open beam analytical RGDs that shall be manufactured to be used hand held without safety devices are
26	exempt from the requirements of Paragraph (c) of this Rule and shall be constructed according to International
27	Standard IEC 62495 that is incorporated by reference and includes subsequent amendments. This standard can be
28	downloaded for one hundred twenty one dollars (\$121.00) at the following website:
29	http://webstore.ansi.org/FindStandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx.aspx.aspx.aspx.aspx.aspx.aspx.asp
30	geNum=0&SearchTermsArray=null%7cIEC+62495+Ed.+1.0+en%3a2011%7cnull.
31	(i) A registrant may apply to the agency, as defined in Rule .0104 of this Chapter, for an exemption from the
32	requirement of a safety device. This request shall include:
33	(1) a description of the safety devices;
34	(2) the reason safety devices cannot be used; and
35	(3) a description of the alternative methods that will be employed to minimize the possibility of an
36	accidental exposure, including procedures to assure that operators and others in the area will be
37	informed of the absence of safety devices.

1	(j) Analytical RGDs shall be provided with a visible and legible label(s) bearing the radiation symbol and the words:
2	(1) "CAUTION HIGH INTENSITY X RAY BEAM," or words having a similar meaning, near the
3	exit port to identify the location of the beam; and
4	(2) "CAUTION RADIATION THIS EQUIPMENT PRODUCES RADIATION WHEN
5	ENERGIZED", or words having a similar meaning, near any switch that energizes an x-ray tube, if
6	the radiation source is an x ray tube.
7	(k) Warning lights labeled with the words "X RAYS ON," or other words having similar meaning, shall be located:
8	(1) near any switch that activates the high voltage to energize an x-ray tube; or
9	(2) in a conspicuous location near the radiation source housing and radiation beam(s) and visible from
10	all instrument access areas.
11	(l) Warning lights shall activate when the x-ray tube is energized.
12	(m) Each x ray tube housing shall be:
13	(1) constructed that when all shutters are closed the leakage radiation measured at a distance of five
14	centimeters from its surface is not capable of producing an exposure in excess of 2.5 millirem
15	(mrem)/ (25 microsieverts μSv) in one hour; and
16	(2) if the tube housing is the primary shielding for the x ray tube, does not produce x rays when the
17	housing is opened or disassembled.
18	(n) Each x ray generator shall be supplied with a protection cabinet which limits leakage radiation measured at a
19	distance of five centimeters from its surface such that it is not capable of producing an exposure in excess of 0.25
20	mrem/2.5μSv in one hour.
21	(o) Permanent radiographic installations and industrial radiography RGDs shall comply with the requirements of Rule
22	.0807 of this Section.
23	(a) The registrant shall document the scope of training and instruction required for the RGD in use.
24	(b) No individual shall be permitted to operate or maintain RGDs unless the individual has received instruction in the
25	basic principles of radiation protection, training specific to the manufacturer's recommendations for safe operation
26	and unique features of the RGD in use, and instruction in the operating and emergency procedures. Instruction and
27	training shall include:
28	(1) Basic principles of radiation protection:
29	(A) radiation fundamentals;
30	(B) source and magnitude of common sources of radiation exposure;
31	(C) units of radiation dose and measurements;
32	(D) potential hazards, biological effects of ionizing radiation, and recognition of symptoms of
33	an acute localized exposure;
34	(E) ALARA (As Low As Reasonably Achievable) principles for radiation protection concepts
35	of time, distance, and shielding to minimize radiation exposure;
36	(F) declared pregnancy policy;
37	(G) occupational, embryo/fetus, and public dose limits; and

1		(H) proper use of individual monitoring devices and survey instruments.
2	(2)	Device specific training for each RGD:
3		(A) hands-on training for proper use;
4		(B) radiation hazards associated with use;
5		(C) precautions to take or measures required to minimize radiation exposure;
6		(D) procedures to prevent unauthorized use; and
7		(E) agency rules regarding use.
8	(3)	Operating and emergency procedure requirements of Rule .0804 in this Section.
9	(c) Records of i	nstruction and training for each individual operating RGDs, documenting that the requirements of this
10	Rule have been	met, shall be maintained and available for agency review during inspection.
11	<u>(d) <mark>Individuals</mark></u>	Persons who will be operating the RGD shall be able to demonstrate an understanding in safe operating
12	procedures and	use of the RGD according to the manufacturer's specifications and to an authorized representative of
13	the Radiation P	rotection Section.
14	(e) Each registr	ant shall provide ring or wrist individual monitoring devices to individuals:
15	<u>(1)</u>	operating open-beam RGDs; and
16	(2)	performing maintenance on an RDG, if the maintenance procedures require the presence of a
17		primary x-ray beam when any local component in the RGD is disassembled or removed.
18		
19	History Note:	Authority G.S. 104E-7;
20		Eff. February 1, 1980;
21		Transferred and Recodified from 15A NCAC 11 .0803 Eff. February 1, 2015;
22		Amended Eff. October 1, 2015;
23		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
24		2019. <u>2019:</u>
25		<u>Amended Eff.</u> <mark>Oetober </mark>

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer individual responsible
29	for radiation protection as defined in Rule.0104 of this Chapter
30	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
31	all operators and support staff for review during the use of an RGD.
32	(d) Normal operating and emergency procedures shall include the following:
33	(1) safe use of the RGD;
34	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
35	<u>and</u>

1	(3)	instructions on reporting to the radiaton safety officer individual responsible for radiation protection
2		and agency of actual or suspected accidental exposure or other radiation safety concerns, such as
3		any unusual occurrence or malfunction that may involve exposure to radiation.
4	(e) Open beam a	and portable handheld RGDs
5	(1)	Registrants shall have operating procedures developed to ensure radiation protective measures are:
6		(A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter;
7		(B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in cases
8		where the primary x-ray beam is not intercepted by a detector device during operation; and
9		(C) available to all individuals operating the RGD.
10	<u>(2)</u>	Operators shall not do the following while operating an RGD:
11		(A) point the primary beam at any individual including him or herself:
12		(B) allow their hand to approach the primary beam; or
13		(C) hold a sample. If a sample is small and it is necessary to hold the sample while operating
14		the RGD, the sample shall be placed in a shielded sample enclosure.
15	(f) Operating and	d emergency procedures shall be available for review by an authorized representative of the Radiation
16	Protection Section	<u>n agency review</u> during inspection.
17	(g) Alignment p	rocedures shall be performed as recommended by the RGD manufacturer.
18	(h) Special alig	nment procedures shall only be used when approved by the radiation safety officer individual
19	responsible for ra	adiation protection and manufacturer of the RGD.
20	(i) Safety Device	e <u>s</u>
21	(1)	Testing
22		(A) Safety devices including interlocks, shutters, and warning lights shall be tested once
23		annually for proper operation on all RGDs in use. If any safety device fails, the RGD shall
24		be taken out of service until corrective action is performed or temporary administrative
25		controls are established and approved in writing by the radiation safety officer individual
26		responsible for radiation protection.
27		(B) Testing records shall include the date test was performed, the list of safety devices tested,
28		the survey instrument used, the calibration date, the results of the test, the name of the
29		individual that performed the test, and any corrective actions for a failed test.
30		(C) Records of the testing shall be retained by the registrant for agency review during
31		inspection.
32	(2)	Bypassing
33		(A) No individual shall bypass a safety device unless the person has obtained approval from
34		the radiation safety officer individual responsible for radiation protection. Procedures for
35		bypassing a safety device shall be incorporated into the radiation protection program by
36		the radiation safety officer individual responsible for radiation protection, as set forth in

1		Rule .1601 of this Chapter, and the operating procedures as set forth in Paragraph (c) of
2		this Rule.
3		(B) The written approval, as granted by the individual responsible for radiation protection, shall
4		include the start and end date of approval.
5		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
6		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
7		ray source housing and the control panel during the bypassing period.
8	<u>(j) <mark>Prior to an i</mark>i</u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
9	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
10	<u>(1)</u>	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
11	(2)	shutters;
12	(3)	collimators; or
13	<u>(4)</u>	beam stops.
14		
15	History Note:	Authority G.S. 104E-7(a)(2);
16		Eff. February 1, 1980;
17		Amended Eff. January 1, 1994;
18		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
19		Amended Eff. October 1, 2015;
20		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
21		2019. <u>2019;</u>
22		Amended Eff. November 1, 2024.

1	10A NCAC 15.	.0805 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2		
3	10A NCAC 15	.0805 OPERATING REQUIREMENTS AREA REQUIREMENTS
4	(a) RGDs shall l	be operated by individuals that have completed the training requirements of Rule .0806 of this Section.
5	(b) Normal ope	rating procedures shall be written and available to all RGD operators and support staff.
6	(c) No individ	ual shall be permitted to operate RGDs in any manner other than that specified in the operating
7	procedures unles	ss the person has obtained written approval of the individual responsible for radiation or the Radiation
8	Safety Officer (l	RSO) as defined in Rule .0104 of this Chapter.
9	(d) No individu	nal shall bypass a safety device unless the person has obtained the approval of the person responsible
10	for radiation saf	ety or the RSO. This process shall be incorporated into the radiation protection program by the RSO,
11	as set forth in R	ule .1603(a), and the operating procedures as set forth in Rule .0603(a)(1)(B). The written approval,
12	as granted by the	e RSO, shall include an expiration date. When a safety device has been bypassed, a legible sign bearing
13	the words "SAF	ETY DEVICE NOT WORKING" or words having a similar meaning shall be placed on the radiation
14	source housing a	and the control panel during the bypassing period.
15	(e) Prior to an i	ndividual modifying the:
16	(1)	x ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
17	(2)	— shutters;
18	(3)	— collimators; or
19	(4)	— beam stops
20	the individual sl	nall determine the tube is off and will remain off until safe conditions have been restored.
21	(f) Safety devic	es including interlocks, shutters, and warning lights shall be tested once annually for proper operation
22	on all RGDs in o	operation. Records of the testing shall be retained by the registrant for three years.
23	(g) No individu	nal shall hold a sample or object while it is being irradiated.
24	(a) Each radiati	on area, as defined in Rule .1601(a)(3) of this Chapter, containing RGDs shall be:
25	(1)	conspicuously posted with caution signs, in accordance with the requirements of Rule .1601(a)(34)
26		of this Chapter, bearing the words "CAUTION - RADIATION AREA", or words having a similar
27		meaning; and
28	<u>(2)</u>	supervised continuously during operation of the RDG or shall utilize one or more of the following:
29		(A) door interlocks;
30		(B) entry monitors; or
31		(C) engineering controls.
32	(b) Access to ea	ach restricted area where an individual may receive a dose equivalent exceeding 100 mrem in any year,
33	but does not exc	eed levels of a radiation area, shall be designated as a controlled area. The area shall be controlled by:
34	<u>(1)</u>	visibly separating adjacent uncontrolled areas so doses do not exceed the limits of Rule .1601(a)(15)
35		of this Chapter; and
36	(2)	posting a sign bearing the words "Warning: X-rays in Use", or words having a similar meaning.

I	(c) The local co	omponents of RGDs shall be located and arranged to include sufficient shielding or access control to
2	ensure no radia	tion levels exist in any area surrounding the local components that result in a dose to an individual
3	present in exces	s of the dose limits in Rule .1601(a)(15) of this Chapter.
4	(d) Surveys sha	all be performed for each RGD, as set forth in Rule .1601(a)(23) of this Chapter, to show compliance
5	with Paragraph	(c) of this Rule.
6	<u>(1)</u>	Radiation survey instruments shall be:
7		(A) capable of measuring the radiation energies of the RGD surveyed; and
8		(B) calibrated annually when a frequency is not recommended by the manufacturer.
9	(2)	Equipment surveys shall confirm radiation levels do not exceed the requirements of Rule
10		.0806(c)(7); (d)(3); .0806(d)(3); and (h)(2) .0806(h)(2) of this Section. Surveys shall be performed:
11		(A) prior to initial use and include testing of warning and safety devices;
12		(B) prior to use following any change in the initial arrangement, including the number or type
13		of local components in the system or x-ray tube source housing;
14		(C) prior to use following any maintenance requiring the disassembly or removal of a local
15		component in the system or x-ray tube source housing that could affect the radiation
16		exposure to personnel; and
17		(D) during the performance of calibration, maintenance, or any alignment procedure if the
18		presence of a primary x-ray beam is required while any local component in the system is
19		disassembled or removed.
20	(3)	A registrant may apply to the agency for approval of procedures differing from those in
21		Subparagraph (d) of this Rule, provided that the registrant demonstrates satisfactory compliance
22		with Paragraph (c) of this Rule.
23	<u>(4)</u>	Records shall be available for agency review during inspection.
24	(e) RGDs in Ru	ale .0806(i) and .0807(2) of this Section, installed after the effective date of this Rule, shall ensure the
25	following provis	sions are met:
26	<u>(1)</u>	A floor plan with equipment arrangement shall be submitted to the agency for review and
27		acknowledgement prior to installation of the system. The floor plan shall include:
28		(A) the proposed location of the system;
29		(B) direction of the useful beam;
30		(C) adjacent areas; and
31		(D) location of the operator.
32	(2)	An area radiation survey shall be performed prior to initial use to show compliance with dose limits
33		of the rules in Section .1600 of this Chapter. The survey shall include:
34		(A) a drawing of the room indicating the location of the x-ray tube and orientation of the useful
35		beam;
36		(B) radiation levels at the operator location and adjacent areas;
37		(C) survey instrument used; and

	(D) name of the service provider that is registered registered, in accordance with Rule .0205 of
	this Chapter Chapter, and date the survey was performed.
(3)	Modifications to the room, RGD, or adjacent areas that may increase the radiation dose to any
	individual shall require a new survey.
(4)	Records of the floor plan with equipment arrangement and survey shall be available for review by
	authorized representative of the Radiation Protection Section agency review during inspection.
History Note:	Authority G.S. 104E-7; 104E-12;
	Eff. February 1, 1980;
	Transferred and Recodified from 15A NCAC 11 .0805 Eff. February 1, 2015;
	Amended Eff. January 1, 2016; October 1, 2015;
	Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
	2019. <u>2019;</u>
	Amended Eff. November 1, 2024.
	<u>(4)</u>

1	10A NCAC 15 .0806 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0806 PERSONNEL REQUIREMENTS EQUIPMENT REQUIREMENTS
4	(a) Personnel operating or maintaining RGDs shall comply with the following:
5	(1) No person shall be permitted to operate or maintain RGDs unless the person has received instruction
6	in the operating and emergency procedures for the RGD and instruction that is in accordance with
7	Rule .1003 of this Chapter.
8	(2) Each registrant operating or maintaining RGDs shall maintain, for inspection by the agency, record
9	of training that demonstrate the requirements of this Rule have been satisfied.
10	(b) The registrant shall provide ring or wrist personnel monitoring equipment to:
11	(1) individuals using open beam RGDs not equipped with a safety device; and
12	(2) individuals maintaining RGDs if the maintenance procedures require the presence of a primary x
13	ray beam when any local component in the RGD is disassembled or removed.
14	(a) Certified and certifiable cabinet x-ray systems shall comply with the following provisions of 21 CFR 1020.40
15	which are hereby incorporated by reference including subsequent amendments and editions.
16	(1) 21 CFR 1020.40(a) Applicability:
17	(2) 21 CFR 1020.40(b) Definitions;
18	(3) 21 CFR 1020.40(c) Requirements; and
19	(4) 21 CFR 1020.40(d) Modifications of a certified system.
20	(b) The regulations cited in Paragraph (a) of this Rule are available free of charge a
21	https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/CFRSearch.cfm?FR=1020.40.
22	(c) All RGD's shall meet the following requirements: requirements, except certified and certifiable cabinet x-ra
23	systems in paragraph (a) of this Rule:
24	(1) Warning devices shall be labeled so the purpose is easily identified.
25	(2) Warning lights of a fail-safe design labeled with the words "X-RAY ON", or words having a similar
26	meaning, shall be located:
27	(A) within sight of any switch that energizes an x-ray tube;
28	(B) in a conspicuous location near the x-ray tube source housing and x-ray beam, and
29	(C) visible from all instrument access areas.
30	(3) Warning lights shall activate when the x-ray tube is energized.
31	(4) Each shutter shall be equipped with a "shutter open" warning light or device of a fail-safe design.
32	(5) A readily visible and legible label bearing the radiation symbol and the words "CAUTION -
33	RADIATION: THIS EQUIPMENT PRODUCES RADIATION WHEN ENERGIZED", or word
34	having a similar meaning, shall be located near any switch that energizes an x-ray tube.
35	(6) Systems containing an x-ray tube shall be equipped with a fail-safe interlock that will shut off high
36	voltage to the tube if the x-ray tube source housing is disassembled or if the tube is removed.

I	<u>(7)</u>	High voltage generator enclosures or any accessible area 5 centimeters from the RGD shall not
2		exceed a dose rate of .25 mrem/hr (.0025 mSv/hr).
3	(d) All open be	eam RGDs shall meet the following additional requirements:
4	<u>(1)</u>	Each beam port of the x-ray tube source housing shall be equipped with a beam shutter interlocked
5		with the x-ray accessory coupling, or collimator, so that the port will not open unless a collimator
6		or a component coupling is in place.
7	<u>(2)</u>	Shutters at unused ports shall be secured in the closed position to prevent unintended opening.
8	(3)	The x-ray tube source housing shall be constructed so that when all shutters are closed, the leakage
9		radiation measured at a distance of five centimeters from the housing surface does not exceed 2.5
10		mrem (25 microSv) in one hour.
11	<u>(4)</u>	A safety device or interlock shall prevent the entry of any portion of an individual's body into the
12		primary x-ray beam or which causes the primary beam to shut off upon entry into its path.
13	<u>(5)</u>	A registrant may apply to the agency, as defined in Rule .0106 of this Chapter, for an exemption
14		from the requirement of a safety device in Subparagraph (d)(3) of this Rule. The request shall
15		include:
16		(A) justification for the use of an open beam system instead of an enclosed beam system;
17		(B) a description of other safety devices that have been evaluated and reason why a safety
18		devices cannot be used; and
19		(C) a description of the alternative methods that will be employed to minimize the possibility
20		of an accidental exposure, including procedures to assure that operators and others in the
21		area will be informed of the absence of safety devices.
22	(e) All enclose	ed beam RGDs shall meet the following additional requirements:
23	(1)	The radiation source, sample or object, detector, and analyzing crystal (if used) shall be enclosed to
24		prevent entry of any portion of the body during normal operation.
25	(2)	All doors and panels shall be equipped with an interlock. The interlock shall be of a fail-safe design.
26	(f) Bimodal be	am RGDs with the ability to override interlocks between enclosed and open beam shall be designed to
27	be engaged wit	h a device or tool and meet the following requirements:
28	(1)	The tool or key shall only be used by designated individuals as outlined in operating procedures.
29	(2)	When the tool or key is in use, it shall be captive in the equipment and removal of the tool or key
30		returns the RGD to enclosed beam mode.
31	(3)	System use requirements must follow the current use mode.
32	(g) Portable x-1	ray fluorescence analyzers manufactured to be used in a hand-held configuration without safety devices
33	are exempt fron	n the requirements of Subparagraph (d)(4) of this Rule. The following additional requirements shall be
34	provided on the	e analyzer:
35	<u>(1)</u>	A power switch with the power logo: I/O.
36	(2)	A label with the words "CAUTION: THIS EQUIPMENT PRODUCES X-RAYS WHEN
37		OPERATED", or words with similar meaning.

1	(3)	Indicators visible to operators when x-rays are on. The indicator shall be in the form of a light and
2		a warning symbol or text with the words "X-RAY ON", or words with similar meaning.
3	<u>(4)</u>	Warning labels near each beam port that bear a radiation symbol and the words "WARNING HIGH
4		INTENSITY X-RAYS – DO NOT EXPOSE ANY PART OF BODY TO BEAM", or words having
5		a similar meaning.
6	(h) All gauging	devices shall meet the following additional requirements:
7	<u>(1)</u>	The RGD shall be designed to restrict access to the x-ray beam by personnel who are not trained in
8		accordance with Rule .0803 of this Section.
9	<u>(2)</u>	A useful beam control system shall be provided whenever the useful beam is accessible, and the
10		radiation levels exceed one hundred mrem per hour (100 mrem/hr) (1 mSv/hr) at five centimeters
11		from any accessible surface or five mrem per hour (5 mrem/h) (.05 mSv/h) at thirty centimeters (30
12		cm). The useful beam controls may include a moving shutter, a moving source, or a high voltage
13		power supply.
14	<u>(3)</u>	On-Off indicators shall be marked with symbols or wording clarifying the status of the device.
15	<u>(4)</u>	Each indicating system for automatic beam controls shall consist of at least one "ON" indicating
16		signal, and one "OFF" indicating signal. If lights are used, green indicates the "OFF" and red
17		indicates any other condition of the useful beam control.
18	<u>(5)</u>	Indicators for RGDs high voltage control shall be a yellow or amber warning light with the words
19		"HIGH VOLTAGE ON" and shall be located on the control panel and near the x-ray tube source
20		housing. The warning light shall illuminate only when power is applied to the RGD.
21	<u>(6)</u>	Interlocks shall be used to prevent accidental exposure to high voltage and ionizing radiation.
22	<u>(7)</u>	The RGD shall be conspicuously marked with a label permanently affixed to the device with the
23		following information:
24		(A) ANSI device classification;
25		(B) name of manufacturer;
26		(C) model; and
27		(D) serial number.
28	<u>(8)</u>	Radiation safety labels shall provide instructions and precautions for safe operation. If space is
29		limited on the RGD, operating or service manuals may be referenced for the information.
30	(i) Radiographi	c and radioscopic non-healing arts x-ray equipment operating below energies of 1 MeV designed for
31	non-medical x-r	ray shall comply with the following additional requirements:
32	<u>(1)</u>	Written instructions shall be supplied by the manufacturer or supplier at the time of sale or transfer
33		to the first user. When the manufacturer or supplier does not provide services to the RGD.
34		installation instructions shall describe:
35		(A) radiation safety pertaining to each unit or accessory;
36		(B) instruction for assembly operations when assembly not performed by manufacturer;
37		(C) interconnections instructions of interlocks, warning lights and audible alarms systems;

1		(D) test instructions to determine if the RGD and accessory components are properly operating;
2		<u>and</u>
3		(E) if the x-ray tube assembly is shielded or non-shielded.
4	<u>(2)</u>	Operating instructions shall be supplied by the manufacturer or supplier, at the time of sale or
5		transfer to the first user, in accordance with operating requirements of Rule .0804 of this Section.
6	<u>(3)</u>	The controls shall be:
7		(A) clearly marked with for the "on-off" position of the component disconnecting the power;
8		<u>and</u>
9		(B) equipped with a means to prevent production of x-rays when in the "off" position, such as
10		a key or password. When a key is used, the RGD shall be manufactured so it may only be
11		removed when the key is in the "off" position.
12	<u>(4)</u>	The "X-ray On" indicator control shall be:
13		(A) yellow or amber in color;
14		(B) be of a fail-safe design; and
15		(C) have two indicators viewable from the control panel indicating when x-rays are being
16		produced in a period of greater than 0.5 seconds.
17	<u>(5)</u>	The "X-ray Off" indicators shall be:
18		(A) red in color; and
19		(B) permanently marked.
20	<u>(6)</u>	Shutters devices that control emission of the primary beam shall activate two visual indicators of
21		contrasting colors from the operator's station. One shall activate when shutters are fully closed and
22		the other shall activate when the shutters are not fully closed.
23	<u>(7)</u>	Selection indicators shall indicate which tube assembly or focal spot has been selected if more than
24		one x-ray tube assembly(s) assembly or focal spot can be operated from the control panel.
25	(8)	Warning Device: A red warning lamp or audible device shall be provided on or near the tube
26		assembly in an open beam, for non-permanent installations.
27	(j) All RGDs	shall be secured to prevent access and operation of the device by any individual not meeting the
28	requirements of	Rule .0803 of this Section.
29		
30	History Note:	Authority G.S. 104E-7; 104E-11; 104E-12;
31		Eff. February 1, 1980;
32		Transferred and Recodified from 15A NCAC 11 .0806 Eff. February 1, 2015;
33		Amended Eff. October 1, 2015;
34		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
35		2019. <u>2019;</u>
36		Amended Eff. November 1, 2024.

1	10A NCAC 15 .0807 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0807 PERMANENT RADIOGRAPHIC INSTALLATIONS AND INDUSTRIAL
4	RADIOGRAPHY RGDS SECURITY SCREENING EQUIPMENT REQUIREMENTS FOR
5	GOVERNMENT USE ONLY
6	(a) Permanent radiographic installations and industrial radiography RGDs are exempt from the requirements of t
7	rules of this Section except Rule .0802 and Rule .0804(a), (b)(1)(A), (b)(1)(C), (b)(2), and (b)(3).
8	(b) Permanent radiographic installations and industrial radiography RGDs shall comply with the following rules
9	this Chapter:
10	(1) .0501;
11	(2) .0502;
12	(3) .0506;
13	(4) .0509 .0520;
14	(5) .0522;
15	(6) .0523(a)(1);
16	(7) .0523(a)(3);
17	(8) .0523(a)(6) .0523(a)(15);
18	(9) .0523(b)(1) .0523(b)(4);
19	(10) .0523(b)(6) .0523(b)(7);
20	(11) .0523(b)(9) .0523(b)(12);
21	(12) .0523(c); and
22	(13) .0525.
23	(a) All security screening devices shall meet the following additional requirements:
24	(1) Security screening RGDs shall only be utilized by accredited bomb squads, certified bor
25	technicians, law enforcement agencies, or forensic investigators.
26	(2) The operator must be present and maintain access control during operation of the RGD. If the RG
27	is not operated in a restricted area and the RGD is capable of producing a radiation area, the opera
28	shall:
29	(A) establish a visible barrier;
30	(B) perform a visual check of the controlled area to ensure all unauthorized individuals a
31	removed prior to activating or initiating the RGD; and
32	(C) if the operator is unable to maintain visual control of the area during operation of the RG
33	an additional means to control the area shall be provided, the operator is required
34	implement additional means to control the area so no one can access the radiation area.
35	(3) Utilization logs shall be maintained each time the RGD is used and accurately include the following
36	(A) date and time of use;
37	(B) location of use; and

1		(C) operator of the RGD.
2	<u>(4)</u>	Records of utilization logs shall be available for agency review during inspection.
3	(b) All security	screening systems shall meet the following additional requirements:
4	(1)	Security screening systems shall only be utilized in a correctional institution, detention center, jail,
5		or prison for public safety and security screening purposes.
6	(2)	No individual shall be exposed to the useful beam unless authorized by a law enforcement agency
7		representative.
8	<u>(3)</u>	No individual shall be exposed to the useful beam for demonstration or training purposes.
9	<u>(4)</u>	Screening of staff for training purposes is prohibited.
10	<u>(5)</u>	Policies and procedures shall be established for screening of minors and pregnant individuals.
11	(6)	An inspection zone shall be:
12		(A) established around the system where bystanders are prohibited during operation;
13		(B) visibly marked; and
14		(C) the ambient dose equivalent outside the inspection zone shall not exceed 2 mrem (20
15		microSv) in any 1 hour.
16	<u>(7)</u>	The system shall be stationary, and the exposure switch shall be located in a manner requiring the
17		operator to remain behind a protected barrier during the entire exposure while able to view the
18		<u>following:</u>
19		(A) the individual being scanned;
20		(B) the inspection zone; and
21		(C) any access areas.
22	<u>(8)</u>	Equipment surveys shall be conducted to verify compliance with reference effective dose limits, the
23		inspection zone, and manufacturer specified parameters. Surveys shall be performed:
24		(A) upon installation;
25		(B) every 12 months; and
26		(C) after maintenance that may affect the system's shielding or x-ray beam.
27	<u>(9)</u>	Reference effective dose limits shall be met as follows:
28		(A) General-use systems reference effective dose shall not exceed 25 microrem (.25 microSv)
29		per screening.
30		(B) Limited-use systems reference effective dose shall not exceed 1 mrem (10 microSv) per
31		screening.
32		(C) The reference effective dose received by an individual shall not exceed 25 mrem (250
33		microSv) in a 12-month period for both general use and limited-use systems.
34	(10)	Compliance to reference effective dose limits shall be demonstrated by the registrant maintaining
35		records of each individual screened. Records shall show one of the following:
36		(A) the number of screenings each individual received, for General-use systems, does not
37		exceed 1,000 in a 12-month period; or

1		(B) the reference effective dose multiplied by the number of screenings, for both General-use
2		and Limited-use systems, does not exceed 25 mrem (250 microSv) in a 12-month period.
3	<u>(11)</u>	Records of each individual scanned at the same facility shall be maintained for available for review
4		by an authorized representative of the Radiation Protection Section agency review during inspection.
5	(12)	Each individual being screened shall be informed the system emits radiation and be provided with
6		the following prior to scanning:
7		(A) the estimated effective dose from one (1) screening;
8		(B) an example to compare the dose to a commonly known source of radiation; and
9		(C) confirmation the screening complies with the reference effective dose limits in
10		Subparagraph (b)(9) of this Rule.
11		
12	History Note:	Authority G.S. 104E-7;
13		Eff. October 1, 2015. <u>2015:</u>
14		Amended Eff. November 1, 2024.

```
1
      10A NCAC 15 .0808 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
 2
 3
      10A NCAC 15.0808
                                APPLICABLE RULES FOR BOMB DETECTION RGDS OTHER EQUIPMENT
 4
                                REQUIREMENTS
 5
      Bomb detection RGDs utilized by accredited bomb squads and certified bomb technicians shall comply with the
 6
      following rules of this Chapter:
 7
                      .0501;
               (1)
 8
               (2)
                      .0502;
 9
                       .0509;
               (3)
                       .0511 .0520 except for the requirements for a direct reading pocket dosimeter and operating alarm
10
11
                       ratemeter in .0512(a);
                       .0522:
12
               (5)
13
                       .0523(a)(1);
14
                       .0523(a)(3);
               (7)
15
               (8)
                       .0523(a)(6) .0523(a)(15);
                     .0523(b)(1) .0523(b)(4);
16
                       .0523(b)(6) .0523(b)(7);
17
18
               (11) .0523(b)(9) .0523(b)(12);
19
               (12)
                       .0523(c); and
               (13) .0525.
20
21
      (a) RGD's not listed in Rule .0801 of this Section, or that are not able to meet the equipment requirements of
22
      either Rule .0806 or .0807 of this Section, shall not be sold, installed, or used prior teeto the agency completing review
23
      of information regarding the RGD and determining if use of the RGD is allowed. The user or manufacturer of the
24
      RGD shall submit the following information to the agency for review:
25
                       equipment form for application;
               (1)
26
               (2)
                       manufacturer manual;
27
               (3)
                       description of use;
28
               (4)
                       operator training;
29
                       a survey in accordance with Rule.0805(d) of this Section;
               (5)
               <u>(6)</u>
30
                       an area survey in accordance with Rule.0805(e)(2) of this Section;
31
                       the hazard level associated with use of the RGD; and
               (7)
32
               (8)
                       means to achieve radiation protection equivalent to the Rules rules of this Section.
33
      (b) After receiving the information in Paragraph (a) of this Rule, the agency will respond to the applicant in writing
34
      within 30 days. Upon review, the agency may require additional information if use of the RGD is allowed.
35
36
                       Authority G.S. 104E-7;
      History Note:
37
                       Eff. October 1, 2015. 2015;
```

Amended Eff. November 1, 2024.

1

Burgos, Alexander N

Subject:

FW: Radiation Protection Commission Rules-October 2024

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Thursday, October 3, 2024 8:30 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: Re: Radiation Protection Commission Rules-October 2024

Thank you Travis. We will get back to you with the revisions.

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From: Wiggs, Travis C < travis.wiggs@oah.nc.gov>
Sent: Thursday, October 3, 2024 8:25:36 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N < alexander.burgos@oah.nc.gov >

Subject: RE: Radiation Protection Commission Rules-October 2024

Thank you both for the quick response and, Regina, godspeed to you in Asheville. Please see my comments and requests below.

.0801:

• In (c), my suggested change was to make this subsection one complete sentence ("RGDs used...and evaluation in this Section include:"). I suggested the change since complete sentences were used in (a) and (b). The way it has been changed leaves the first sentence as incomplete. I'm satisfied either way you want to do it.

.0802:

- In (20)(C), based on your response, consider putting quotation marks around "beam on", similar to how "X-ray On" or "enclosed beam" was done in other sections of this Rule.
- In (21), line 2, add a comma after "screening". It appears that some change was made to "screening" on line 3 after my initial request to change.

.0803:

- On pg. 3, line 11, you indicated this process will be defined in "upcoming rule 10A NCAC 15 .0104(b)(1)". What is your intended timeline to amend this exisiting Rule to include the definition? How will the regulated public know how to "demonstrate an understanding in safe operating procedures and use" if the upcoming rule is not codified prior to codification of .0803?
- My third request for this Rule was intended for a different rule. Please disregard.

.0804:

• For the phrase "individual responsible for radiation protection", you indicated this is planning to be defined in forthcoming Rule 10A NCAC 15. 0104(b)(4). How will the regulated public know who how to identify this individual if .0104(b)(4) is not codified prior to codification of this Rule? 10A NCAC 15.0203(a) says unregistered radiation machines or facilities shall "designate on the application form an individual who shall be responsible for radiation protection." How does this Rule, and any other exisiting rules, ensure the

- "individual responsible for radiation protection" is qualified to approve an individual to "operate an RGD in any manner other than that specified in the operating procedures"?
- On pg. 2, line 15 and 29, consider saying "shall be available for review by an authorized respresentative of the Radiation Protection Section during inspection."

.0805:

• On pg. 3, lines 5-6, consider saying "shall be available for review by an authorized representative of the Radiation Protection Section during inspection."

.0806:

• In (c), lines 22-23, in the added language, add a comma after "requirements" and capitalize "rule".

.0807:

- In line 33, what is meant by "shall be provided"? Is the operator providing this or someone else? Or is the operator required to "implement" some additional measure of protection?
- In (11), lines 3-4, consider saying "shall be available for review by an authorized representative of the Radiation Protection Section during inspection."

.0808:

• In (a), lines 22-23, it appears an extra "review" was added during the revision. The request would have it read as "used prior to the agency completing a review of information...."

Let me know if you have any questions.

Thanks,

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

Burgos, Alexander N

Subject: FW: Radiation Protection Commission Rules-October 2024

Attachments: 10-2024_Radiation Protection Commission RTC Responses.docx; 10A NCAC 15

.0801.docx; 10A NCAC 15 .0802.docx; 10A NCAC 15 .0803.docx; 10A NCAC 15 .0804.docx; 10A NCAC 15 .0805.docx; 10A NCAC 15 .0806.docx; 10A NCAC 15

.0807.docx; 10A NCAC 15 .0808.docx

From: Black, Shanah <shanah.black@dhhs.nc.gov>

Sent: Wednesday, October 2, 2024 2:33 PM

To: Wiggs, Travis C <travis.wiggs@oah.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov>

Subject: RE: Radiation Protection Commission Rules-October 2024

Good afternoon,

I am attaching responses to your Request for Changes and the revised rules. Regina, author of these rules, is in Asheville and has been working on the revisions despite the hardships around her.

Please let us know if there are further revisions, explanations we need to make.

Thank you for your assistance.

Request for Changes Pursuant to N.C. Gen. Stat. § 150B-21.10

Staff reviewed these Rules to ensure that each Rule is within the agency's statutory authority, reasonably necessary, clear and unambiguous, and adopted in accordance with Part 2 of the North Carolina Administrative Procedure Act. Following review, staff has issued this document that may request changes pursuant to G.S. 150B-21.10 from your agency or ask clarifying questions.

The imposition of a question implies that the rule as written is unclear or there is some ambiguity. If the request includes questions and you do not understand the question, please contact the reviewing attorney to discuss. Failure to respond may result in a staff opinion recommending objection.

Staff may suggest the agency "consider" an idea or language in this document. This is in no way a formal request that the agency adopt the idea or language but rather is offered merely for consideration which the agency may find preferable and clarifying.

To properly submit rewritten rules, please refer to the following Rules in the NC Administrative Code:

- Rule 26 NCAC 02C .0108 The Rule addresses general formatting.
- Rule 26 NCAC 02C .0404 The Rule addresses changing the introductory statement.
- Rule 26 NCAC 02C .0405 The Rule addresses properly formatting changes made after publication in the NC Register.

Note the following general instructions:

- 1. You must submit the revised rule via email to oah.rules@oah.nc.gov. The electronic copy must be saved as the official rule name (XX NCAC XXXX).
- 2. For rules longer than one page, insert a page number.
- 3. Use line numbers; if the rule spans more than one page, have the line numbers reset at one for each page.
- 4. Do not use track changes. Make all changes using manual strikethroughs, underlines and highlighting.
- 5. You cannot change just one part of a word. For example:
 - Wrong: "aAssociation"
 - Right: "association Association"
- 6. Treat punctuation as part of a word. For example:
 - Wrong: "day; and"
 - Right: "day, day; and"
- 7. Formatting instructions and examples may be found at: www.ncoah.com/rules/examples.html.

If you have any questions regarding proper formatting of edits after reviewing the rules and examples, please contact the reviewing attorney.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0801

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 15: Consider deleting, ". RGDs addressed" as it appears unnecessary.

Response: Item addressed in the text.

Line 19: Add "s" to the end of "device" and considering adding "operating" before "below".

Response: Item addressed in text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0802

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 17: Consider adding "successfully" before "completed".

Response: Item addressed in the text.

Lines 18-19: Consider deleting "can be found on the school website" as it appears superfluous.

Response: Item addressed in the text.

Line 29: Consider replacing "that" with "the" before "part".

Response: Item addressed in the text.

Line 30: Consider replacing ", that are necessary" with "located".

Response: Item addressed in text.

On pg. 2, Line 4: "Exposure" to what? What are the "administrative or regulatory limits" and where are they located? Consider a cross-reference to clarify what the limits are for your regulated public.

Response: Item addressed in the text.

Line 5: Add a comma after "contacted".

Response: Item addressed in the text.

Line 8: Consider deleting the comma after "beam".

Response: Item addressed in the text.

Line 9: Add a comma after "indicator", delete "or", and add a comma after "indicator".

Response: Item addressed in the text.

Line 15: Consider adding "of the device." after "specifications".

Response: Item addressed in the text.

Line 34: Per the OAH Style Guide, don't abbreviate using "e.g."

Response: Item addressed in the text.

Line 35: Add "the" before "switch".

Response: Adding "the" changes the whole meaning of the phrase. Beam on, means the beam is on, not describing what the beam is on

On pg. 3, Line 2: add a comma after "screening".

Response: Item addressed in the text.

Line 4: Add a comma after "x-rays".

Response: Item addressed in the text.

Line 14: Add a comma after "operation", replace "and" with "when", add a comma after "imaging", and delete "in which".

Response: Item addressed in the text.

Line 24: Did you intend to use "parts" instead of "arts"?

Response: Item addressed in the text.

On pg. 4, Line 19: Add a comma after "components".

Response: Item addressed in the text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0803

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

On pg. 2, Line 25: Add "the" before "manufacturer's".

Response: Item addressed in the text.

On pg. 3, Line 11: How does an individual "demonstrate an understanding in safe operating procedures and use"? Is this found in another rule or law?

Response: Item addressed in the text. This will be defined in upcoming rule 10A NCAC 15.0104(b)(1).

Lines 22-23: How does the regulated public determine if their items or devices contain "exempt quantities of byproduct material?" Consider inserting the appropriate source(s), i.e. 10 CFR 30.70.

Please see the explanation for Lines 13-14, above, for the resolution of this comment.

Response: Was this meant for another rule?

Line 24: The earliest this Rule can become effective is November 1, 2024.

Response: Item addressed in text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0804

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 28 and elsewhere: How is the "individual responsible for radiation protection" determined? Is this process or determination found in a rule?

Response: This term is currently used in Rule 10A NCAC 15.0203(a)(2). It has never been defined and is being defined in the phase 7 RPC rules, 10A NCAC 15.0104(b)(4). Just trying to use the same verbiage used in other RP rules.

On pg. 2, Line 15 and elsewhere: What "agency" will be conducting the inspection? Please be specific.

Response: Item addressed in the text.

Line 27: Add "the" before "survey".

Response: Item addressed in the text.

Line 28: Add "a" before "failed".

Response: Item addressed in the text.

On pg. 3, (j), Lines 6-7: Consider restructuring this sentence for a more active voice. "An individual shall determine tube is off, and will remain off until safe conditions have been restored, prior to an individual modifying the following:"

Response: Item addressed in the text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0805

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

On pg. 2, Line 3: Consider deleting "present".

Response: Item addressed in the text.

Line 10: Add "806" before "(d)" and "(h)". Response: Item addressed in the text.

On pg. 3, Line 1: Add a comma after "registered" and after "Chapter".

Response: Item addressed in the text.

Line 5: What "agency" will be conducting the inspection? Please be specific.

Response: Item addressed in the text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0806

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE</u>: This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 21: I was unable to access the regulations through the linked website. How can you ensure the regulations can be accessed free of charge by the regulated public? Response: This website does come up. Did you try it with the period?

On pg. 3, (h)(4), Lines 14-15: Are the colors uniform for said devices? Do all manufacturers use the same colors?

Response: Yes, it is an ANSI standard, and it is explained in the fiscal note.

On pg. 4, Line 22: Delete "(s)" at the end of "assembly".

Response: Item addressed in the text.

Line 24: Consider adding "for" before "non-permanent".

Response: Item addressed in the text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0807

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 33: Who provides "an additional means to control the area"? How is this accomplished?

Response: Item addressed in the text

On pg. 3, Line 3: What "agency" will be conducting the inspection? Please be specific. Response: Item addressed in the text.

AGENCY: Radiation Protection Commission

RULE CITATION: 10A NCAC 15.0808

DEADLINE FOR RECEIPT: October 11, 2024

<u>PLEASE NOTE:</u> This request may extend to several pages. Please be sure you have reached the end of the document.

The Rules Review Commission staff has completed its review of this Rule prior to the Commission's next meeting. The Commission has not yet reviewed this Rule and therefore there has not been a determination as to whether the Rule will be approved. You may email the reviewing attorney to inquire concerning the staff recommendation.

In reviewing this Rule, the staff recommends the following changes be made:

Line 21: Add a comma after "Section".

Response: Item addressed in the text.

Line 22: Delete the colon after "to" and add "a" before "review".

Response: Item addressed in the text.

Line 32: Don't capitalize "Rules".

Response: Item addressed in the text.

1	10A NCAC 15	.0801 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	SECTIO	N .0800 - REQUIREMENTS FOR NON-HUMAN USE OF RADIATION GENERATING
3		DEVICES
4		
5	10A NCAC 15	.0801 PURPOSE AND SCOPE
6	(a) This Section	on provides special additional requirements for use of ionizing radiation generating devices (RGDs)
7	operating above	five thousand electron volts (5 keV), but below one million electron volts (1 MeV) that are in addition
8	to (1 MeV). Th	requirements in of this Section are in addition to the provisions in other sections Sections .0100,
9	.0200, .1000, ar	ad .1600 of this Chapter.
10	(b) This Section	n does not pertain to radiation safety requirements for x-ray equipment that is covered in other sections
11	industrial radio	graphic machines for non-human use that are covered in Section .0500 of this Chapter Chapter, (e.g.,
12	x-rays in the he	aling arts in Section .0600 of this Chapter, and particle accelerators in Section .0900 of this Chapter).
13	Chapter.	
14	(c) RGDs used	I for the purpose of elemental analysis, microstructural analysis, quality assurance, quality control,
15	research and de	velopment, gauging and measurement, or other nondestructive testing and evaluation. <mark>RGDs addressed</mark>
16	in this Section i	nelude: This section includes:
17	(1)	analytical RGDs;
18	(2)	cabinet x-ray systems;
19	(3)	electron beam devices operating below 1MeV;
20	<u>(4)</u>	electron microscopes;
21	<u>(5)</u>	ion implantation equipment, low energy;
22	<u>(6)</u>	gauging devices;
23	<u>(7)</u>	radiographic and radioscopic non-healing arts x-ray equipment; and
24	(8)	security screening devices and systems for government use only.
25		
26	History Note:	Authority G.S. 104E-7;
27		Eff. February 1, 1980;
28		Transferred and Recodified from 15A NCAC 11 .0801 Eff. February 1, 2015;
29		Amended Eff. October 1, 2015;
30		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
31		2019. <u>2019;</u>
32		Amended Eff. November 1, 2024.

10A NCAC 15 .0802 is amended with changes as published in 38:19 NCR 1219-1229 as follows:

10A NCAC 15.0802 DEFINITIONS

- In addition to terms found in Rule .0104 of this Chapter Chapter, the following definitions shall apply to this Section:
- 5 (1) "Accredited bomb squad" means a law enforcement agency utilizing certified bomb technicians.
 - (2) "Accessible surface" means the external or outside surface of the enclosure or housing provided by the manufacturer or designer of the RGD. This includes the high-voltage generator, doors, access panels, latches, control knobs, and other permanently mounted hardware, and including the plane across the exterior edge of any opening.
 - (2)(3) "Analytical RGD equipment" means equipment that uses electronic means to generate ionizing radiation for the purpose of examining the microstructure of materials, materials i.e. using direct x-ray transmission, x-ray diffraction, x-ray fluorescence, and x-ray spectroscopy.
 - (3)(4) "Analytical RGD system" means a group of local and remote components utilizing x-rays to determine the elemental composition or to examine the microstructure of materials.
 - (4) "Bomb detection RGDs" means RGDs used for the sole purpose of remotely detecting explosive devices.
 - (5) "Certified bomb technician" means a member of an accredited bomb squad who has successfully completed the FBI Hazardous Devices School. Information pertaining to this program can be found on the school website at http://www.fbi.gov/about-us/cirg/hazardous-devices.
 - (6) "Certifiable cabinet x-ray system" means an existing uncertified RGD that has been modified to meet the certification requirements specified in 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (7) "Certified cabinet x-ray system" means an RGD utilized in an enclosed, interlocked cabinet, such that the radiation machine will not operate unless all openings are securely closed. These systems shall be certified in accordance with 21 CFR 1010.2 1010.2, as incorporated by reference in Rule .0117 of this Chapter, as being manufactured and assembled pursuant to the provisions of 21 CFR 1020.40 1020.40, as incorporated by reference in Rule .0117 of this Chapter.
 - (8) "Collimator" means a device or mechanism by which the x-ray beam is restricted in size.
 - (9) "Control panel" means that the part of the x-ray control where the switches, knobs, pushbuttons, and other hardware are, that are necessary located for manually setting the technique factors.
 - (10) "Electron Beam Device" means any device using electrons below 1MeV to heat, join, or otherwise irradiate materials.
 - (11) "Enclosed beam RGD" means an RGD with all possible x-ray beam paths contained in a chamber, coupled chambers, or other beam-path-confinement devices devices, to prevent any part of the body from intercepting the beam during normal operations. Normal access to the primary beam path, such as a sample chamber door, shall be interlocked with the high voltage of the x-ray tube or the shutter

1		for the beam to be considered "enclosed." An open-beam device placed in an interlocked enclosure
2		is considered an "enclosed beam" unless there are provisions for routine bypassing of the interlocks.
3	<u>(12)</u>	"Emergency procedure" means the written pre-planned steps to be taken in the event of actual or
4		suspected radiation exposure of an individual exceeding administrative or regulatory limits-limits
5		found in 10A NCAC 15 .1601(a)(8) and .1601(a)(15). This procedure shall include the names and
6		telephone numbers of individuals to be contacted contacted, as well as directives for processing
7		individual monitoring devices.
8	(12) (13)) "Fail-safe characteristics" means a design feature that causes the radiation beam to terminate, port
9		shutters to close, or otherwise prevents emergence of the primary beam, beam upon the failure of a
10		safety or warning device. For example, if an "X-ray On" light indicator or light indicator, shutter
11		indicator shutter indicator, or interlock fails, the radiation beam shall terminate.
12	<u>(14)</u>	"Gauging device" means a mechanism containing a source of ionizing radiation that is designed and
13		manufactured for the purpose of determining or controlling thickness, density, level, interface
14		location, or qualitative or quantitative composition of materials. It may include components such as
15		radiation shields, useful-beam controls, and other safety features in order to meet the requirements
16		or specifications. specifications of the device.
17	<u>(15)</u>	"General-use system" means a security screening system that delivers an effective dose of 25
18		microrem (0.25 microSv) or less per screening.
19	(13) (16	"Hand-held x-ray system" means any device or equipment that is portable and used for similar
20		purposes as analytical RGD equipment.
21	(14)	"Hybrid gauge" means an x-ray gauge device utilizing both x-ray and radioactive sources.
22	(15)	"Industrial radiography" means RGDs used to make radiographic images to examine the structure
23		of materials by nondestructive methods. These RGDs shall not be contained in a cabinet and are not
24		permanent installations.
25	<u>(17)</u>	"Inspection Zone" means the area established for the purpose of controlling access where screening
26		is performed. Areas controlled due to the presence of radiation shall include areas of ingress, egress,
27		gates, portals, and traffic paths. The area outside of the inspection zone shall not exceed the limits
28		of Rule .1601(a)(13) of this Section.
29	<u>(18)</u>	"Interlock" means a feature designed to prevent access to an area of radiation hazard by preventing
30		entry or by automatically removing the hazard.
31	(16) (19)) "Ion implantation equipment, low-energy" means any enclosed device operating below 1MeV used
32		to accelerate elemental ions and implant them in other materials.
33	(17) (20)	The land the source assembly housing except for:
34		(A) the primary beam;
35		(B) scatter radiation emanating from other components (e.g., shutter or collimator); and
36		(C) radiation produced when the beam on switch or timer is not activated.

1	<u>(21)</u>	"Limited-use system" means a screening system that is capable of delivering an effective dose
2		greater than 25 microrem (0.25 microSv) per screening but shall not exceed an effective dose of 1
3		mrem (10 microSv) per screening. screening.
4	(18) (22)	"Local components" means part of an RGD x-ray system and include areas that are struck by x rays
5		<u>x rays.</u> such as radiation source housings, port and shutter assemblies, collimators, sample holders,
6		cameras, goniometers, detectors, and shielding, but do not include power supplies, transformers,
7		amplifiers, readout devices, and control panels.
8	(19) (23)	"Mobile RGD" means RGD equipment mounted on a permanent base with wheels or casters for
9		moving while <u>completely</u> assembled.
10	(20) (24)	"Normal operating procedures" means step-by-step instructions necessary to accomplish a task.
11		These procedures shall include sample insertion and manipulation, equipment alignment, routine
12		maintenance by the registrant, and data recording procedures that are related to radiation safety.
13	(21) (25)	"Open-beam RGD" means a device or system designed in such a way that the primary beam is not
14		completely enclosed during normal operation operation, and when used for analysis, gauging, or
15		imaging imaging, in which an individual could accidentally place some part of their body in the
16		primary beam or stray radiation path during normal operation.
17	(22)	"Permanent radiographic installation" means an RGD utilized in an enclosed shielded room, cell, or
18		vault that allows entry when the RGD is not energized.
19	(23) (26)	"Portable RGD" means RGD equipment designed to be carried of carried by hand.
20	(24) (27)	"Primary beam" means radiation that passes through an aperture of the source assembly housing by
21		a direct path from the radiation source.
22	(25) (28)	"Radiation generating device (RGD)" means any system, device, subsystem, or machine component
23		that may generate generate, by electronic means means, x-rays or particle radiation above 5 keV,
24		but below 1 MeV, and not used for healing arts parts on humans or animals. Examples of RGDs are
25		the following may be used as a:
26		(A) analytical RGD equipment mobile RGD;
27		(B) certified and certifiable cabinet x-ray systems portable RGD; or
28		(C) gauging devices using x-ray sources; stationary RGD.
29		(D) hybrid gauging devices;
30		(E) e beam welders;
31		(F) baggage scanners;
32		(G) industrial radiography RGDs; and
33		(H) permanent radiographic installations.
34	(26) (29)	"Remote components" means parts of an RGD x-ray system that are not struck by $\frac{x - rays}{x}$
35		such as power supplies, transformers, amplifiers, readout devices, and control panels.

I	<u>(30)</u>	"Safety Device" means a device, interlock or system that prevents the entry of any portion of an
2		individual's body into the primary x-ray beam or that will cause the beam to shut off upon entry into
3		its path.
4	(27) (31)	"Scattered radiation" means radiation, other than leakage radiation, that during passage through
5		matter, has been deviated in direction or has been modified by a decrease in energy.
6	<u>(32)</u>	"Screening" means the sum of scans necessary for a security screening system to image concealed
7		objects as intended by the system design under normal operating conditions.
8	<u>(33)</u>	"Security screening device" means a non-human use open-beam device designed for the detection
9		of contraband or weapons concealed in baggage, mail, packages, or other structures. These devices
10		include bomb detection devices used for the sole purpose of detecting explosive devices.
11	<u>(34)</u>	"Security screening system" means a system specifically designed to detect contraband and weapons
12		concealed on a person and is used for the sole purpose of public safety and security evaluation by
13		law enforcement.
14	(28) (35)	"Shutter" means an adjustable device, generally made of lead or other high atomic number material,
15		fixed to a source assembly housing to intercept, block, or collimate the primary beam.
16	(29) (36)	"Source" means the point of origin of the radiation, such as the focal spot of an x-ray tube.
17	(30) (37)	"Stationary RGD" means RGD equipment that is installed or placed in a fixed location.
18	(31) (38)	"Stray radiation" means the sum of leakage and scatter radiation emanating from the source
19		assembly or other ecomponents components, except for the primary beam, and radiation produced
20		when the beam on switch or timer is not activated.
21	<u>(39)</u>	"Warning device" means an audible or visible signal that warns individuals of a potential radiation
22		<u>hazard.</u>
23	(32) (40)	"X-ray generator" means the part of an x-ray system that provides the accelerating (high) voltage
24		and current for the x-ray tube.
25	(33)	"X ray gauge" means an x ray producing device designed and manufactured for the purpose of
26		detecting, measuring, gauging, or controlling thickness, density, level, or interface location of
27		manufactured products.
28	<u>(41)</u>	"X-ray source housing" means the portion of an RGD system which contains the x-ray tube and
29		emitting target. The housing often contains radiation shielding material or inherently provides
30		shielding.
31		
32	History Note:	Authority G.S. 104E-7;
33		Eff. February 1, 1980;
34		Transferred and Recodified from 15A NCAC 11 .0802 Eff. February 1, 2015;
35		Amended Eff. October 1, 2015;
36		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
37		2019. <u>2019;</u>

Amended Eff. November 1, 2024.

1

1	10A NCAC 15 .0803 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0803 EQUIPMENT REQUIREMENTS PERSONNEL REQUIREMENTS
4	(a) Certified cabinet x ray systems shall meet the requirements of 21 CFR 1020.40 as incorporated by reference in
5	Rule .0117(a)(3) of this Chapter.
6	(b) All certified and certifiable cabinet x-ray systems shall:
7	(1) be constructed so that, the radiation emitted from the system shall not exceed an exposure of 0.5
8	milliroentgen (mR) in one hour at any point five centimeters outside the external surface; and
9	(2) have a fail safe interlock that prevents irradiation when the cabinet, chamber, or coupled chambers
10	are open.
11	(c) Open beam analytical RGD systems shall be equipped with a safety device that prevents the entry of any portion
12	of an individual's body into the primary x-ray beam path that causes the beam to be shut off upon entry into its path.
13	(d) Open beam analytical RGDs shall be provided with a visible and legible indication of:
14	(1) x ray tube status (ON OFF) located near the radiation source housing, if the primary beam is
15	controlled in this manner; or
16	(2) shutter status (OPEN CLOSED) or beam status (ON OFF) located near each port on the radiation
17	source housing, if the primary beam is controlled in this manner.
18	(e) Warning devices on open beam analytical RGDs shall be labeled so that their purpose is identified. On open beam
19	analytical RGDs installed after February 1, 1980, warning devices and lights shall have fail safe characteristics.
20	(f) Unused ports on radiation source housings for open-beam RGDs shall be secured in the closed position in a manner
21	that will prevent unintended opening.
22	(g) Each port on the radiation source housing on open beam analytical RGDs installed after February 1, 1980 and
23	designed to accommodate interchangeable components shall be equipped with a shutter that cannot be opened unless
24	a collimator or a component coupling is connected to the port.
25	(h) Portable open beam analytical RGDs that shall be manufactured to be used hand held without safety devices are
26	exempt from the requirements of Paragraph (c) of this Rule and shall be constructed according to International
27	Standard IEC 62495 that is incorporated by reference and includes subsequent amendments. This standard can be
28	downloaded for one hundred twenty one dollars (\$121.00) at the following website:
29	http://webstore.ansi.org/FindStandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx?SearchString=IEC+62495+Ed.+1.0+en%3a2011&SearchOption=0&Pandards.aspx.aspx.aspx.aspx.aspx.aspx.aspx.asp
30	geNum=0&SearchTermsArray=null%7cIEC+62495+Ed.+1.0+en%3a2011%7cnull.
31	(i) A registrant may apply to the agency, as defined in Rule .0104 of this Chapter, for an exemption from the
32	requirement of a safety device. This request shall include:
33	(1) a description of the safety devices;
34	(2) the reason safety devices cannot be used; and
35	(3) a description of the alternative methods that will be employed to minimize the possibility of an
36	accidental exposure, including procedures to assure that operators and others in the area will be
37	informed of the absence of safety devices.

1	(j) Analytical RGDs shall be provided with a visible and legible label(s) bearing the radiation symbol and the words:
2	(1) "CAUTION HIGH INTENSITY X RAY BEAM," or words having a similar meaning, near the
3	exit port to identify the location of the beam; and
4	(2) "CAUTION RADIATION THIS EQUIPMENT PRODUCES RADIATION WHEN
5	ENERGIZED", or words having a similar meaning, near any switch that energizes an x-ray tube, if
6	the radiation source is an x ray tube.
7	(k) Warning lights labeled with the words "X RAYS ON," or other words having similar meaning, shall be located:
8	(1) near any switch that activates the high voltage to energize an x-ray tube; or
9	(2) in a conspicuous location near the radiation source housing and radiation beam(s) and visible from
10	all instrument access areas.
11	(l) Warning lights shall activate when the x-ray tube is energized.
12	(m) Each x ray tube housing shall be:
13	(1) constructed that when all shutters are closed the leakage radiation measured at a distance of five
14	centimeters from its surface is not capable of producing an exposure in excess of 2.5 millirem
15	(mrem)/ (25 microsieverts μSv) in one hour; and
16	(2) if the tube housing is the primary shielding for the x ray tube, does not produce x rays when the
17	housing is opened or disassembled.
18	(n) Each x ray generator shall be supplied with a protection cabinet which limits leakage radiation measured at a
19	distance of five centimeters from its surface such that it is not capable of producing an exposure in excess of 0.25
20	mrem/2.5μSv in one hour.
21	(o) Permanent radiographic installations and industrial radiography RGDs shall comply with the requirements of Rule
22	.0807 of this Section.
23	(a) The registrant shall document the scope of training and instruction required for the RGD in use.
24	(b) No individual shall be permitted to operate or maintain RGDs unless the individual has received instruction in the
25	basic principles of radiation protection, training specific to the manufacturer's recommendations for safe operation
26	and unique features of the RGD in use, and instruction in the operating and emergency procedures. Instruction and
27	training shall include:
28	(1) Basic principles of radiation protection:
29	(A) radiation fundamentals;
30	(B) source and magnitude of common sources of radiation exposure;
31	(C) units of radiation dose and measurements;
32	(D) potential hazards, biological effects of ionizing radiation, and recognition of symptoms of
33	an acute localized exposure;
34	(E) ALARA (As Low As Reasonably Achievable) principles for radiation protection concepts
35	of time, distance, and shielding to minimize radiation exposure;
36	(F) declared pregnancy policy;
37	(G) occupational, embryo/fetus, and public dose limits; and

1		(H) proper use of individual monitoring devices and survey instruments.
2	<u>(2)</u>	Device specific training for each RGD:
3		(A) hands-on training for proper use;
4		(B) radiation hazards associated with use;
5		(C) precautions to take or measures required to minimize radiation exposure;
6		(D) procedures to prevent unauthorized use; and
7		(E) agency rules regarding use.
8	(3)	Operating and emergency procedure requirements of Rule .0804 in this Section.
9	(c) Records of i	nstruction and training for each individual operating RGDs, documenting that the requirements of this
10	Rule have been	met, shall be maintained and available for agency review during inspection.
11	(d) Individuals	s who will be operating the RGD shall be able to demonstrate an understanding in safe operating
12	procedures and	use of the RGD according to the manufacturer's specifications and to an authorized representative of
13	the Radiation Pr	rotection Section.
14	(e) Each registr	ant shall provide ring or wrist individual monitoring devices to individuals:
15	(1)	operating open-beam RGDs; and
16	(2)	performing maintenance on an RDG, if the maintenance procedures require the presence of a
17		primary x-ray beam when any local component in the RGD is disassembled or removed.
18		
19	History Note:	Authority G.S. 104E-7;
20		Eff. February 1, 1980;
21		Transferred and Recodified from 15A NCAC 11 .0803 Eff. February 1, 2015;
22		Amended Eff. October 1, 2015;
23		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
24		2019. <u>2019;</u>
25		Amended Eff. October November 1, 2024.

1	10A NCAC 15 .0804 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0804 AREA REQUIREMENTS OPERATING REQUIREMENTS
4	(a) The local components of RGDs shall be located and arranged to include sufficient shielding or access control to
5	ensure no radiation levels exist in any area surrounding the local components that could result in a dose to an individual
6	present in excess of the dose limits given in Rule .1611(a) of this Chapter.
7	(b) Survey Requirements
8	(1) Radiation surveys, as set forth in Rule .1613(a) and (b) of this Chapter, of all RGDs sufficient to
9	show compliance with Paragraph (a) of this Rule, shall be performed:
10	(A) within 30 days after initial operation of the device;
11	(B) prior to use following any change in the initial arrangement, including the number or type
12	of local components in the system; and
13	(C) prior to use following any maintenance requiring the disassembly or removal of a local
14	component in the system that could affect the radiation exposure to personnel.
15	(2) A registrant may apply to the agency for approval of procedures differing from those in
16	Subparagraph (b)(1) of this Rule, provided that the registrant demonstrates satisfactory compliance
17	with Paragraph (a) of this Rule.
18	(3) Surveys shall be performed with a radiation survey instrument capable of the following:
19	(A) measuring the radiation energies of the system surveyed;
20	(B) confirming that the radiation limits of this Section are met; and
21	(C) calibrated according to the manufacture's recommended frequency or at least annually
22	when a frequency is not recommended.
23	(c) Each area of use or room containing RGDs shall be conspicuously posted with caution signs in accordance with
24	the requirements of Rule .1623 of this Chapter, bearing the radiation caution symbol and the words "CAUTION X
25	RAY EQUIPMENT," or words having a similar meaning.
26	(a) RGDs shall only be operated by individuals who have completed the requirements in Rule .0803 of this Section.
27	(b) No individual shall be permitted to operate an RGD in any manner other than that specified in the operating
28	procedures, unless the individual has obtained written approval from the radiation safety officer individual responsible
29	for radiation protection as defined in Rule.0104 of this Chapter
30	(c) Normal operating and emergency procedures from the manufacturer or supplier of the RGD shall be available to
31	all operators and support staff for review during the use of an RGD.
32	(d) Normal operating and emergency procedures shall include the following:
33	(1) safe use of the RGD;
34	(2) protocols in the event of device malfunction, emergency, or incident involving radiation exposure;
35	<u>and</u>

1	<u>(3)</u>	instructions on reporting to the radiaton safety officer individual responsible for radiation protection
2		and agency of actual or suspected accidental exposure or other radiation safety concerns, such as
3		any unusual occurrence or malfunction that may involve exposure to radiation.
4	(e) Open beam	and portable handheld RGDs
5	<u>(1)</u>	Registrants shall have operating procedures developed to ensure radiation protective measures are:
6		(A) provided to meet the requirements of Rule .1601(a)(15) of this Chapter;
7		(B) taken to avoid exposure to any individual from the transmitted primary x-ray beam in cases
8		where the primary x-ray beam is not intercepted by a detector device during operation; and
9		(C) available to all individuals operating the RGD.
10	<u>(2)</u>	Operators shall not do the following while operating an RGD:
11		(A) point the primary beam at any individual including him or herself:
12		(B) allow their hand to approach the primary beam; or
13		(C) hold a sample. If a sample is small and it is necessary to hold the sample while operating
14		the RGD, the sample shall be placed in a shielded sample enclosure.
15	(f) Operating as	nd emergency procedures shall be available for authorized representative of the Radiation Protection
16	agency review d	uring inspection.
17	(g) Alignment p	procedures shall be performed as recommended by the RGD manufacturer.
18	(h) Special ali	gnment procedures shall only be used when approved by the radiation safety officer individual
19	responsible for 1	adiation protection and manufacturer of the RGD.
20	(i) Safety Device	<u>ves</u>
21	<u>(1)</u>	Testing
22		(A) Safety devices including interlocks, shutters, and warning lights shall be tested once
23		annually for proper operation on all RGDs in use. If any safety device fails, the RGD shall
24		be taken out of service until corrective action is performed or temporary administrative
25		controls are established and approved in writing by the radiation safety officer individual
26		responsible for radiation protection.
27		(B) Testing records shall include the date test was performed, the list of safety devices tested,
28		the survey instrument used, the calibration date, the results of the test, the name of the
29		individual that performed the test, and any corrective actions for a failed test.
30		(C) Records of the testing shall be retained by the registrant for agency review during
31		inspection.
32	<u>(2)</u>	Bypassing
33		(A) No individual shall bypass a safety device unless the person has obtained approval from
34		the radiation safety officer individual responsible for radiation protection. Procedures for
35		bypassing a safety device shall be incorporated into the radiation protection program by
36		the radiation safety officer individual responsible for radiation protection, as set forth in

1		Rule .1601 of this Chapter, and the operating procedures as set forth in Paragraph (c) of
2		this Rule.
3		(B) The written approval, as granted by the individual responsible for radiation protection, shall
4		include the start and end date of approval.
5		(C) When a safety device has been bypassed, a legible sign bearing the words "SAFETY
6		DEVICE NOT WORKING", or words having a similar meaning, shall be placed on the x-
7		ray source housing and the control panel during the bypassing period.
8	<u>(j) <mark>Prior to an i</mark>i</u>	ndividual modifying the following, the An individual shall determine the tube is off off, and will remain
9	off until safe co	nditions have been restored: restored, prior to an individual modifying the following;
10	<u>(1)</u>	x-ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
11	(2)	shutters;
12	(3)	collimators; or
13	<u>(4)</u>	beam stops.
14		
15	History Note:	Authority G.S. 104E-7(a)(2);
16		Eff. February 1, 1980;
17		Amended Eff. January 1, 1994;
18		Transferred and Recodified from 15A NCAC 11 .0804 Eff. February 1, 2015;
19		Amended Eff. October 1, 2015;
20		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
21		2019. <u>2019;</u>
22		Amended Eff. November 1, 2024.

1	10A NCAC 15.	.0805 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2		
3	10A NCAC 15	.0805 OPERATING REQUIREMENTS AREA REQUIREMENTS
4	(a) RGDs shall l	be operated by individuals that have completed the training requirements of Rule .0806 of this Section.
5	(b) Normal ope	rating procedures shall be written and available to all RGD operators and support staff.
6	(c) No individ	ual shall be permitted to operate RGDs in any manner other than that specified in the operating
7	procedures unles	ss the person has obtained written approval of the individual responsible for radiation or the Radiation
8	Safety Officer (l	RSO) as defined in Rule .0104 of this Chapter.
9	(d) No individu	nal shall bypass a safety device unless the person has obtained the approval of the person responsible
10	for radiation saf	ety or the RSO. This process shall be incorporated into the radiation protection program by the RSO,
11	as set forth in R	ule .1603(a), and the operating procedures as set forth in Rule .0603(a)(1)(B). The written approval,
12	as granted by the	e RSO, shall include an expiration date. When a safety device has been bypassed, a legible sign bearing
13	the words "SAF	ETY DEVICE NOT WORKING" or words having a similar meaning shall be placed on the radiation
14	source housing a	and the control panel during the bypassing period.
15	(e) Prior to an i	ndividual modifying the:
16	(1)	x ray tube system, resulting in the removal of tube housings, covers, or shielding materials;
17	(2)	— shutters;
18	(3)	— collimators; or
19	(4)	— beam stops
20	the individual sl	nall determine the tube is off and will remain off until safe conditions have been restored.
21	(f) Safety devic	es including interlocks, shutters, and warning lights shall be tested once annually for proper operation
22	on all RGDs in o	operation. Records of the testing shall be retained by the registrant for three years.
23	(g) No individu	nal shall hold a sample or object while it is being irradiated.
24	(a) Each radiati	on area, as defined in Rule .1601(a)(3) of this Chapter, containing RGDs shall be:
25	<u>(1)</u>	conspicuously posted with caution signs, in accordance with the requirements of Rule .1601(a)(34)
26		of this Chapter, bearing the words "CAUTION - RADIATION AREA", or words having a similar
27		meaning; and
28	<u>(2)</u>	supervised continuously during operation of the RDG or shall utilize one or more of the following:
29		(A) door interlocks;
30		(B) entry monitors; or
31		(C) engineering controls.
32	(b) Access to ea	ach restricted area where an individual may receive a dose equivalent exceeding 100 mrem in any year,
33	but does not exc	eed levels of a radiation area, shall be designated as a controlled area. The area shall be controlled by:
34	<u>(1)</u>	visibly separating adjacent uncontrolled areas so doses do not exceed the limits of Rule .1601(a)(15)
35		of this Chapter; and
36	(2)	posting a sign bearing the words "Warning: X-rays in Use", or words having a similar meaning.

I	(c) The local co	omponents of RGDs shall be located and arranged to include sufficient shielding or access control to
2	ensure no radia	tion levels exist in any area surrounding the local components that result in a dose to an individual
3	present in exces	s of the dose limits in Rule .1601(a)(15) of this Chapter.
4	(d) Surveys sha	all be performed for each RGD, as set forth in Rule .1601(a)(23) of this Chapter, to show compliance
5	with Paragraph	(c) of this Rule.
6	<u>(1)</u>	Radiation survey instruments shall be:
7		(A) capable of measuring the radiation energies of the RGD surveyed; and
8		(B) calibrated annually when a frequency is not recommended by the manufacturer.
9	(2)	Equipment surveys shall confirm radiation levels do not exceed the requirements of Rule
10		.0806(c)(7); (d)(3); .0806(d)(3); and (h)(2) .0806(h)(2) of this Section. Surveys shall be performed:
11		(A) prior to initial use and include testing of warning and safety devices;
12		(B) prior to use following any change in the initial arrangement, including the number or type
13		of local components in the system or x-ray tube source housing;
14		(C) prior to use following any maintenance requiring the disassembly or removal of a local
15		component in the system or x-ray tube source housing that could affect the radiation
16		exposure to personnel; and
17		(D) during the performance of calibration, maintenance, or any alignment procedure if the
18		presence of a primary x-ray beam is required while any local component in the system is
19		disassembled or removed.
20	(3)	A registrant may apply to the agency for approval of procedures differing from those in
21		Subparagraph (d) of this Rule, provided that the registrant demonstrates satisfactory compliance
22		with Paragraph (c) of this Rule.
23	<u>(4)</u>	Records shall be available for agency review during inspection.
24	(e) RGDs in Ru	ale .0806(i) and .0807(2) of this Section, installed after the effective date of this Rule, shall ensure the
25	following provis	sions are met:
26	<u>(1)</u>	A floor plan with equipment arrangement shall be submitted to the agency for review and
27		acknowledgement prior to installation of the system. The floor plan shall include:
28		(A) the proposed location of the system;
29		(B) direction of the useful beam;
30		(C) adjacent areas; and
31		(D) location of the operator.
32	(2)	An area radiation survey shall be performed prior to initial use to show compliance with dose limits
33		of the rules in Section .1600 of this Chapter. The survey shall include:
34		(A) a drawing of the room indicating the location of the x-ray tube and orientation of the useful
35		beam;
36		(B) radiation levels at the operator location and adjacent areas;
37		(C) survey instrument used; and

1		(D) name of the service provider that is registered registered, in accordance with Rule .0205 of
2		this Chapter Chapter, and date the survey was performed.
3	(3)	Modifications to the room, RGD, or adjacent areas that may increase the radiation dose to any
4		individual shall require a new survey.
5	(4)	Records of the floor plan with equipment arrangement and survey shall be available for authorized
6		representatives of Radiation Protection agency review during inspection.
7		
8	History Note:	Authority G.S. 104E-7; 104E-12;
9		Eff. February 1, 1980;
10		Transferred and Recodified from 15A NCAC 11 .0805 Eff. February 1, 2015;
11		Amended Eff. January 1, 2016; October 1, 2015;
12		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
13		2019. <u>2019;</u>
14		Amended Eff. November 1, 2024.

1	10A NCAC 15 .0806 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2	
3	10A NCAC 15 .0806 PERSONNEL REQUIREMENTS EQUIPMENT REQUIREMENTS
4	(a) Personnel operating or maintaining RGDs shall comply with the following:
5	(1) No person shall be permitted to operate or maintain RGDs unless the person has received instruction
6	in the operating and emergency procedures for the RGD and instruction that is in accordance with
7	Rule .1003 of this Chapter.
8	(2) Each registrant operating or maintaining RGDs shall maintain, for inspection by the agency, record
9	of training that demonstrate the requirements of this Rule have been satisfied.
10	(b) The registrant shall provide ring or wrist personnel monitoring equipment to:
11	(1) individuals using open beam RGDs not equipped with a safety device; and
12	(2) individuals maintaining RGDs if the maintenance procedures require the presence of a primary x
13	ray beam when any local component in the RGD is disassembled or removed.
14	(a) Certified and certifiable cabinet x-ray systems shall comply with the following provisions of 21 CFR 1020.40
15	which are hereby incorporated by reference including subsequent amendments and editions.
16	(1) 21 CFR 1020.40(a) Applicability;
17	(2) 21 CFR 1020.40(b) Definitions;
18	(3) 21 CFR 1020.40(c) Requirements; and
19	(4) 21 CFR 1020.40(d) Modifications of a certified system.
20	(b) The regulations cited in Paragraph (a) of this Rule are available free of charge a
21	https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/CFRSearch.cfm?FR=1020.40.
22	(c) All RGD's shall meet the following requirements: requirements except certified and certifiable cabinet x-ra
23	systems in paragraph (a) of this rule:
24	(1) Warning devices shall be labeled so the purpose is easily identified.
25	(2) Warning lights of a fail-safe design labeled with the words "X-RAY ON", or words having a similar
26	meaning, shall be located:
27	(A) within sight of any switch that energizes an x-ray tube;
28	(B) in a conspicuous location near the x-ray tube source housing and x-ray beam, and
29	(C) visible from all instrument access areas.
30	(3) Warning lights shall activate when the x-ray tube is energized.
31	(4) Each shutter shall be equipped with a "shutter open" warning light or device of a fail-safe design.
32	(5) A readily visible and legible label bearing the radiation symbol and the words "CAUTION
33	RADIATION: THIS EQUIPMENT PRODUCES RADIATION WHEN ENERGIZED", or word
34	having a similar meaning, shall be located near any switch that energizes an x-ray tube.
35	(6) Systems containing an x-ray tube shall be equipped with a fail-safe interlock that will shut off high
36	voltage to the tube if the x-ray tube source housing is disassembled or if the tube is removed.

l	<u>(7)</u>	High voltage generator enclosures or any accessible area 5 centimeters from the RGD shall not
2		exceed a dose rate of .25 mrem/hr (.0025 mSv/hr).
3	(d) All open be	eam RGDs shall meet the following additional requirements:
4	<u>(1)</u>	Each beam port of the x-ray tube source housing shall be equipped with a beam shutter interlocked
5		with the x-ray accessory coupling, or collimator, so that the port will not open unless a collimator
6		or a component coupling is in place.
7	<u>(2)</u>	Shutters at unused ports shall be secured in the closed position to prevent unintended opening.
8	(3)	The x-ray tube source housing shall be constructed so that when all shutters are closed, the leakage
9		radiation measured at a distance of five centimeters from the housing surface does not exceed 2.5
10		mrem (25 microSv) in one hour.
11	<u>(4)</u>	A safety device or interlock shall prevent the entry of any portion of an individual's body into the
12		primary x-ray beam or which causes the primary beam to shut off upon entry into its path.
13	<u>(5)</u>	A registrant may apply to the agency, as defined in Rule .0106 of this Chapter, for an exemption
14		from the requirement of a safety device in Subparagraph (d)(3) of this Rule. The request shall
15		include:
16		(A) justification for the use of an open beam system instead of an enclosed beam system;
17		(B) a description of other safety devices that have been evaluated and reason why a safety
18		devices cannot be used; and
19		(C) a description of the alternative methods that will be employed to minimize the possibility
20		of an accidental exposure, including procedures to assure that operators and others in the
21		area will be informed of the absence of safety devices.
22	(e) All enclose	ed beam RGDs shall meet the following additional requirements:
23	(1)	The radiation source, sample or object, detector, and analyzing crystal (if used) shall be enclosed to
24		prevent entry of any portion of the body during normal operation.
25	(2)	All doors and panels shall be equipped with an interlock. The interlock shall be of a fail-safe design.
26	(f) Bimodal be	am RGDs with the ability to override interlocks between enclosed and open beam shall be designed to
27	be engaged wit	h a device or tool and meet the following requirements:
28	(1)	The tool or key shall only be used by designated individuals as outlined in operating procedures.
29	(2)	When the tool or key is in use, it shall be captive in the equipment and removal of the tool or key
30		returns the RGD to enclosed beam mode.
31	(3)	System use requirements must follow the current use mode.
32	(g) Portable x-1	ray fluorescence analyzers manufactured to be used in a hand-held configuration without safety devices
33	are exempt fron	n the requirements of Subparagraph (d)(4) of this Rule. The following additional requirements shall be
34	provided on the	e analyzer:
35	<u>(1)</u>	A power switch with the power logo: I/O.
36	(2)	A label with the words "CAUTION: THIS EQUIPMENT PRODUCES X-RAYS WHEN
37		OPERATED", or words with similar meaning.

1	(3)	Indicators visible to operators when x-rays are on. The indicator shall be in the form of a light and
2		a warning symbol or text with the words "X-RAY ON", or words with similar meaning.
3	<u>(4)</u>	Warning labels near each beam port that bear a radiation symbol and the words "WARNING HIGH
4		INTENSITY X-RAYS – DO NOT EXPOSE ANY PART OF BODY TO BEAM", or words having
5		a similar meaning.
6	(h) All gauging	g devices shall meet the following additional requirements:
7	<u>(1)</u>	The RGD shall be designed to restrict access to the x-ray beam by personnel who are not trained in
8		accordance with Rule .0803 of this Section.
9	(2)	A useful beam control system shall be provided whenever the useful beam is accessible, and the
10		radiation levels exceed one hundred mrem per hour (100 mrem/hr) (1 mSv/hr) at five centimeters
11		from any accessible surface or five mrem per hour (5 mrem/h) (.05 mSv/h) at thirty centimeters (30
12		cm). The useful beam controls may include a moving shutter, a moving source, or a high voltage
13		power supply.
14	(3)	On-Off indicators shall be marked with symbols or wording clarifying the status of the device.
15	<u>(4)</u>	Each indicating system for automatic beam controls shall consist of at least one "ON" indicating
16		signal, and one "OFF" indicating signal. If lights are used, green indicates the "OFF" and red
17		indicates any other condition of the useful beam control.
18	(5)	Indicators for RGDs high voltage control shall be a yellow or amber warning light with the words
19		"HIGH VOLTAGE ON" and shall be located on the control panel and near the x-ray tube source
20		housing. The warning light shall illuminate only when power is applied to the RGD.
21	<u>(6)</u>	Interlocks shall be used to prevent accidental exposure to high voltage and ionizing radiation.
22	<u>(7)</u>	The RGD shall be conspicuously marked with a label permanently affixed to the device with the
23		following information:
24		(A) ANSI device classification;
25		(B) name of manufacturer;
26		(C) model; and
27		(D) serial number.
28	<u>(8)</u>	Radiation safety labels shall provide instructions and precautions for safe operation. If space is
29		limited on the RGD, operating or service manuals may be referenced for the information.
30	(i) Radiograph	ic and radioscopic non-healing arts x-ray equipment operating below energies of 1 MeV designed for
31	non-medical x-	ray shall comply with the following additional requirements:
32	(1)	Written instructions shall be supplied by the manufacturer or supplier at the time of sale or transfer
33		to the first user. When the manufacturer or supplier does not provide services to the RGD,
34		installation instructions shall describe:
35		(A) radiation safety pertaining to each unit or accessory;
36		(B) instruction for assembly operations when assembly not performed by manufacturer;
37		(C) interconnections instructions of interlocks, warning lights and audible alarms systems;

I		(D) test instructions to determine if the RGD and accessory components are properly operating;
2		<u>and</u>
3		(E) if the x-ray tube assembly is shielded or non-shielded.
4	(2)	Operating instructions shall be supplied by the manufacturer or supplier, at the time of sale or
5		transfer to the first user, in accordance with operating requirements of Rule .0804 of this Section.
6	<u>(3)</u>	The controls shall be:
7		(A) clearly marked with for the "on-off" position of the component disconnecting the power;
8		<u>and</u>
9		(B) equipped with a means to prevent production of x-rays when in the "off" position, such as
10		a key or password. When a key is used, the RGD shall be manufactured so it may only be
11		removed when the key is in the "off" position.
12	(4)	The "X-ray On" indicator control shall be:
13		(A) yellow or amber in color;
14		(B) be of a fail-safe design; and
15		(C) have two indicators viewable from the control panel indicating when x-rays are being
16		produced in a period of greater than 0.5 seconds.
17	(5)	The "X-ray Off" indicators shall be:
18		(A) red in color; and
19		(B) permanently marked.
20	<u>(6)</u>	Shutters devices that control emission of the primary beam shall activate two visual indicators of
21		contrasting colors from the operator's station. One shall activate when shutters are fully closed and
22		the other shall activate when the shutters are not fully closed.
23	<u>(7)</u>	Selection indicators shall indicate which tube assembly or focal spot has been selected if more than
24		one x-ray tube assembly(s) assembly or focal spot can be operated from the control panel.
25	<u>(8)</u>	Warning Device: A red warning lamp or audible device shall be provided on or near the tube
26		assembly in an open beam, for non-permanent installations.
27	(j) All RGDs	shall be secured to prevent access and operation of the device by any individual not meeting the
28	requirements of	Rule .0803 of this Section.
29		
30	History Note:	Authority G.S. 104E-7; 104E-11; 104E-12;
31		Eff. February 1, 1980;
32		Transferred and Recodified from 15A NCAC 11 .0806 Eff. February 1, 2015;
33		Amended Eff. October 1, 2015;
34		Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. June 22,
35		2019. <u>2019:</u>
36		Amended Eff. November 1, 2024.

1	10A NCAC 15 .0	807 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
2		
3	10A NCAC	15 .0807 PERMANENT RADIOGRAPHIC INSTALLATIONS AND INDUSTRIAL
4	RADIO	GRAPHY RGDS SECURITY SCREENING EQUIPMENT REQUIREMENTS FOR
5		GOVERNMENT USE ONLY
6	(a) Permanent ra	diographic installations and industrial radiography RGDs are exempt from the requirements of the
7	rules of this Secti	on except Rule .0802 and Rule .0804(a), (b)(1)(A), (b)(1)(C), (b)(2), and (b)(3).
8	(b) Permanent ra	adiographic installations and industrial radiography RGDs shall comply with the following rules of
9	this Chapter:	
10	(1)	0501;
11	(2)	0502;
12	(3)	0506;
13	(4)	.0509 .0520;
14	(5)	0522;
15	(6)	0523(a)(1);
16	(7)	0523(a)(3);
17	(8)	.0523(a)(6) .0523(a)(15);
18	(9)	0523(b)(1)0523(b)(4);
19	(10)	.0523(b)(6) .0523(b)(7);
20	(11)	.0523(b)(9) .0523(b)(12);
21	(12)	0523(c); and
22	(13)	0525.
23	(a) All security s	creening devices shall meet the following additional requirements:
24	<u>(1)</u>	Security screening RGDs shall only be utilized by accredited bomb squads, certified bomb
25		technicians, law enforcement agencies, or forensic investigators.
26	(2)	The operator must be present and maintain access control during operation of the RGD. If the RGD
27		$\underline{is\ not\ operated\ in\ a\ restricted\ area\ and\ the\ RGD\ is\ capable\ of\ producing\ a\ radiation\ area,\ the\ operator}$
28		shall:
29		(A) establish a visible barrier;
30		(B) perform a visual check of the controlled area to ensure all unauthorized individuals are
31		removed prior to activating or initiating the RGD; and
32		(C) if the operator is unable to maintain visual control of the area during operation of the RGD,
33		an additional means to control the area shall be provided provided so no one can access
34		the radiation area.
35	(3)	Utilization logs shall be maintained each time the RGD is used and accurately include the following:
36		(A) date and time of use;
37		(B) location of use; and

1		(C) operator of the RGD.
2	<u>(4)</u>	Records of utilization logs shall be available for agency review during inspection.
3	(b) All security	screening systems shall meet the following additional requirements:
4	(1)	Security screening systems shall only be utilized in a correctional institution, detention center, jail,
5		or prison for public safety and security screening purposes.
6	(2)	No individual shall be exposed to the useful beam unless authorized by a law enforcement agency
7		representative.
8	(3)	No individual shall be exposed to the useful beam for demonstration or training purposes.
9	<u>(4)</u>	Screening of staff for training purposes is prohibited.
10	<u>(5)</u>	Policies and procedures shall be established for screening of minors and pregnant individuals.
11	<u>(6)</u>	An inspection zone shall be:
12		(A) established around the system where bystanders are prohibited during operation;
13		(B) visibly marked; and
14		(C) the ambient dose equivalent outside the inspection zone shall not exceed 2 mrem (20
15		microSv) in any 1 hour.
16	<u>(7)</u>	The system shall be stationary, and the exposure switch shall be located in a manner requiring the
17		operator to remain behind a protected barrier during the entire exposure while able to view the
18		following:
19		(A) the individual being scanned;
20		(B) the inspection zone; and
21		(C) any access areas.
22	<u>(8)</u>	Equipment surveys shall be conducted to verify compliance with reference effective dose limits, the
23		inspection zone, and manufacturer specified parameters. Surveys shall be performed:
24		(A) upon installation;
25		(B) every 12 months; and
26		(C) after maintenance that may affect the system's shielding or x-ray beam.
27	<u>(9)</u>	Reference effective dose limits shall be met as follows:
28		(A) General-use systems reference effective dose shall not exceed 25 microrem (.25 microSv)
29		per screening.
30		(B) Limited-use systems reference effective dose shall not exceed 1 mrem (10 microSv) per
31		screening.
32		(C) The reference effective dose received by an individual shall not exceed 25 mrem (250
33		microSv) in a 12-month period for both general use and limited-use systems.
34	<u>(10)</u>	Compliance to reference effective dose limits shall be demonstrated by the registrant maintaining
35		records of each individual screened. Records shall show one of the following:
36		(A) the number of screenings each individual received, for General-use systems, does not
37		exceed 1,000 in a 12-month period; or

1		(B) the reference effective dose multiplied by the number of screenings, for both General-use
2		and Limited-use systems, does not exceed 25 mrem (250 microSv) in a 12-month period.
3	<u>(11)</u>	Records of each individual scanned at the same facility shall be maintained for authorized
4		representatives of the Radiation Protection agency review during inspection.
5	<u>(12)</u>	Each individual being screened shall be informed the system emits radiation and be provided with
6		the following prior to scanning:
7		(A) the estimated effective dose from one (1) screening:
8		(B) an example to compare the dose to a commonly known source of radiation; and
9		(C) confirmation the screening complies with the reference effective dose limits in
10		Subparagraph (b)(9) of this Rule.
11		
12	History Note:	Authority G.S. 104E-7;
13		Eff. October 1, 2015. <u>2015:</u>
14		Amended Eff. November 1, 2024.

```
1
      10A NCAC 15 .0808 is amended with changes as published in 38:19 NCR 1219-1229 as follows:
 2
 3
      10A NCAC 15.0808
                                APPLICABLE RULES FOR BOMB DETECTION RGDS OTHER EQUIPMENT
 4
                                REQUIREMENTS
 5
      Bomb detection RGDs utilized by accredited bomb squads and certified bomb technicians shall comply with the
 6
      following rules of this Chapter:
 7
                      .0501;
               (1)
 8
               (2)
                       .0502;
 9
                        .0509;
               (3)
                        .0511 .0520 except for the requirements for a direct reading pocket dosimeter and operating alarm
10
11
                       ratemeter in .0512(a);
                        .0522:
12
               (5)
13
                        .0523(a)(1);
14
                        .0523(a)(3);
               (7)
15
               (8)
                        .0523(a)(6) .0523(a)(15);
                      .0523(b)(1) .0523(b)(4);
16
                       .0523(b)(6) .0523(b)(7);
17
18
               (11) .0523(b)(9) .0523(b)(12);
19
               (12)
                       .0523(c); and
               (13) .0525.
20
21
      (a) RGD's not listed in Rule .0801 of this Section, or that are not able to meet the equipment requirements of
22
      either Rule .0806 or .0807 of this Section, shall not be sold, installed, or used prior testo a review the agency completing
23
      review of information regarding the RGD and determining if use of the RGD is allowed. The user or manufacturer of
24
      the RGD shall submit the following information to the agency for review:
25
                       equipment form for application;
               (1)
26
               (2)
                       manufacturer manual;
27
               (3)
                       description of use;
28
               (4)
                       operator training;
29
                       a survey in accordance with Rule.0805(d) of this Section;
               (5)
               <u>(6)</u>
30
                       an area survey in accordance with Rule.0805(e)(2) of this Section;
31
                       the hazard level associated with use of the RGD; and
               (7)
32
               (8)
                       means to achieve radiation protection equivalent to the Rules rules of this Section.
33
      (b) After receiving the information in Paragraph (a) of this Rule, the agency will respond to the applicant in writing
34
      within 30 days. Upon review, the agency may require additional information if use of the RGD is allowed.
35
36
                       Authority G.S. 104E-7;
      History Note:
37
                       Eff. October 1, 2015. 2015;
```

Amended Eff. November 1, 2024.

1

Burgos, Alexander N

From: Black, Shanah

Sent: Friday, September 27, 2024 1:43 PM **To:** Wiggs, Travis C; Kissinger, Regina

Cc: Burgos, Alexander N

Subject: RE: Radiation Protection Commission Rules-October 2024

Thank you Travis

From: Wiggs, Travis C <travis.wiggs@oah.nc.gov> **Sent:** Friday, September 27, 2024 1:42 PM

To: Black, Shanah <shanah.black@dhhs.nc.gov>; Kissinger, Regina <regina.kissinger@dhhs.nc.gov>

Cc: Burgos, Alexander N <alexander.burgos@oah.nc.gov> **Subject:** Radiation Protection Commission Rules-October 2024

Good afternoon,

I'm the attorney who reviewed the rules submitted by the N.C. Radiation Protection Commission for the October 2024 RRC meeting. The RRC will formally review these rules at its meeting on Wednesday, October 30, 2024, at 10:00 a.m. The meeting will be a hybrid of in-person and WebEx attendance, and an evite should be sent to you as we get close to the meeting. If there are any other representatives from your agency who want to attend virtually, please let me know prior to the meeting, and we will get evites out to them as well.

Attached is the Request for Changes Pursuant to G.S. 150B-21.10. Please submit the revised Rules and forms to me via email, no later than 5 p.m. on October 11, 2024. Let me know if you have any questions.

Thank you.

Travis C. Wiggs Rules Review Commission Counsel Office of Administrative Hearings Telephone: 984-236-1929

Email: travis.wiggs@oah.nc.gov

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