

Bureau of Radiological Health Registration for Veterinary Radiation Machines

Please send the following items:

- Complete application
- Nonrefundable fee in a <u>check or money order</u> payable to lowa Department of Health and Human Services (IDHHS)
- Completed equipment information
- The date(s) of your last equipment calibration

Mailing Address:

Iowa Department of Health and Human Services, Bureau of Radiological Health Lucas State Office Building, 5th Floor 321 East 12th Street Des Moines, IA 5031

You can also complete the application online at <a href="https://hhs.iowa.gov/public-health/radiological-health/radiological-health/radiological-health-radiological-h

Customer Support Phone: (855) 824-4357 Email: radhealthia@idph.iowa.gov

FACILITY INFORMAT	ION (Type or	print the	informati	on below)	
Facility Name:					
Facility Contact Person: _				Street Address:	
City:	States			Phone Number:	
Email: Require	d			Registration Number: <u>VET</u>	
EIN/SSN			□this	is a new registration \Box this is a renewal application	
Туре	Price (\$)	Unit(s)	Total	Registration/Renewal: Please submit your	
X-Ray (Itube)	60			application approximately 45 days before your	
Fluoro (Itube)	60			registration expired.	
Rad/Fuoro (2 tubes)	120			If your registration is 30 days past due, please <u>add</u>	
СТ	60				
C-arm	60			\$25 per month late fee to the total fees due.	
Cone Beam CT	60			Max fee for Veterinary is \$2,000	
Intraoral Dental	60				
Late Registration	\$25/month				
Final Total Due					

AFFIRMATION QUESTIONS (Required)

Sole proprietor applicants must answer these questions only. If you answer, "Yes" to any of the qu	
below (I) attach a signed letter explaining the details of the incident, including date(s), location(s), status, re	eason, etc.
(2) attach a copy of any court ordered evaluations, showing completion & recommendations, and/or (3)	
Attach a letter from a physician or treatment program for any medical condition(s).	
(New) Do you have (Renewal) During the previous licensing period, did you develop a medical condition, which in any way currently impairs or limits your ability to perform the duties Of this profession? Medical Condition means any physiological, mental, or psychological condition, impairment, or disorder, including drug addiction and alcoholism.	☐ Yes ☐ No
(New) Have you, within the past 5 years (Renewal) During the previous licensing period, did you engage in illegal or improper use of drugs or other chemical substances?	☐ Yes ☐ No

(New) Have you ever been (Renewal) During the previous licensing period, were you convicted of a misdemeanor or felony crime? (You do not need to answer yes if your sole conviction or convictions are for minor traffic violations with fines under \$250). In answering this question, note that a conviction means a finding, plea, or verdict of guilt made or returned in a criminal proceeding, even if the adjudication of guilt is deferred, withheld, or not entered. This means you must answer yes if a finding or verdict of guilt was returned against you in a criminal proceeding or if you plead guilty, entered a plea of nolo contendere, or entered an Alford plea in a criminal proceeding, even if the court expunged the matter or the court-deferred judgment. You must submit the complaint and judgment of conviction for each offense.				
<u>All applicants</u> must answer these questions. If you answer, "Yes" to any of the questions below attach a signed letter explaining the details of the incident, including date(s), location(s), status, reason (2) attach a copy of any court ordered evaluations, showing completion & recommendations,		. and/or		
(New) Has (Renewal) During the previous licensing period, did any state or other jurisdiction of the United States or any other nation limit, restrict, warn, censure place on probation, suspend, revoke, or otherwise discipline a professional license, permit, registration, or certification issued to you or your organization?		Yes No		
(New) Have there ever been (Renewal) During the previous licensing period, were there judgments or settlements paid on your or your organization behalf as a result of a professional liability case?		Yes No		
(New) Have you ever had (Renewal) During the previous licensing period, did a license, permit, registration, or certification denied, suspended, revoked, or otherwise disciplined by a certification body?				
FACILITY DETAILS (Required)				
Do you have a Radiation Protection Program that meets the parameters as outlined in IDPH guidance?	□Ye	s 🗆 No		
Is dosimetry issued to operators? If yes, Dosimetry Vendor name:				
If no dosimetry is issued, I have documentation from a medical physicist or other personnel qualified to make the determination that no staff will exceed 10% of the annual 5 rem dose limit.				
This facility has been previously registered to use radiation-emitting equipment.		s 🗆 No		
The licensed practitioner is the only operator of this x-ray equipment.		s 🗆 No		
All radiation equipment operators have an lowa permit to operate the equipment.		s 🗆 No		
All radiation equipment operators are trained in safe operating procedures and are competent in the safe use of the radiation machine.				
The facility has a method to log all x-ray exposures with the required information.				
The facility will periodically review the exposure log for repeat trends and reinstruct staff accordingly.				
Leaded aprons and gloves are available for use during x-ray procedures.				
Are facility familiar with Image Gently/Image Wisely campaign advisements specific to the types of equipment your facility operates?				

FLUORO QUESTIONS: (Complete only if you have a fluoroscopy machine)

All fluoroscopic procedures are supervised by an individual who meets the requirements in IAC 641-		
41.1(6)n		
Leaded aprons and gloves and/or portable shields are available for use during fluoroscopy	□Yes □ No	
procedures.		
Facility has a process to maintain records of cumulative fluoroscopic exposure time used and the	□Yes □ No	
number of spot films for each examination.		
Equipment has a dose area product monitor capable of recording the total radiation dose received	□Yes □ No	
by the patient.		
Patient doses are logged in the patient chart for each exam.	□Yes □ No	
Processes in place to review adult doses exceeding 300 rad and child doses (under 18) exceeding 100	□Yes □ No	
rad		

EQUIPMENT INFORMATION

Mark the boxes and fill in your equipment information below. If you are including copies of your most recent calibration reports and the information on the reports is accurate, you do not need to complete this section.

$\Box X$ -Ray (I tube)	□ Rad/Flı	uoro (2 tubes)	\square Fluoro (1 tube)	
□Cone Beam CT	□СТ	☐ C-Arm	☐ Intraoral Dental	
Is this a Mobile Unit? ☐ Yes ☐	☐ Yes ☐ No			
Machine Manufacture: Machine Serial #:				
Machine Model:		Room		
Manufacture Date:		Installa		
Date of current calibration or se	ervice evalua	ation report		
□X-Ray (I tube)	□ Rad/Flι	uoro (2 tubes)	☐ Fluoro (I tube)	
□Cone Beam CT	□СТ	☐ C-Arm	\square Intraoral Dental	
Is this a Mobile Unit? ☐ Yes ☐ No				
Machine Manufacture:		Machine Serial #:		
Machine Model:	Room	Room ID:		
Manufacture Date: Installation Date:				
Date of current calibration or se	ervice evalua	ation report:		
□X-Ray (I tube) □ Rad/Fluoro (2 tubes) □ Fluoro (I tube)				
□X-Ray (1 tube)	□ Rad/Flu	Joro (2 tubes)	☐ Fluoro (I tube)	
□X-Ray (I tube) □Cone Beam CT	□ Rad/Flu	uoro (2 tubes)	☐ Fluoro (I tube) ☐ Intraoral Dental	
• • • • • • • • • • • • • • • • • • • •	□СТ	□ C-Arm	, ,	☐ Yes ☐ No
□Cone Beam CT	□СТ	☐ C-Arm	☐ Intraoral Dental	□ Yes □ No
□Cone Beam CT Is this a Mobile Unit? □ Yes □	□СТ	☐ C-Arm	☐ Intraoral Dental ed outside your facility? ne Serial #:	□ Yes □ No
□ Cone Beam CT Is this a Mobile Unit? □ Yes □ Machine Manufacture:	□СТ	□ C-Arm Is this unit us Machi Room	☐ Intraoral Dental ed outside your facility? ne Serial #:	☐ Yes ☐ No
□Cone Beam CT Is this a Mobile Unit? □ Yes □ Machine Manufacture: Machine Model:	□ CT No	Is this unit us Machi Room	□ Intraoral Dental ed outside your facility? ne Serial #: ID:	☐ Yes ☐ No
□Cone Beam CT Is this a Mobile Unit? □ Yes □ Machine Manufacture: Machine Model: Manufacture Date:	□ CT No	Is this unit us Machi Room	□ Intraoral Dental ed outside your facility? ne Serial #: ID:	☐ Yes ☐ No
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□ Cone Beam CT Is this a Mobile Unit? □ Yes □ Machine Manufacture: Machine Model: Manufacture Date: Date of current calibration or se	□ CT No ervice evalua □ Rad/Flu □ CT	□ C-Arm Is this unit use Machi Room Installa ation report: uoro (2 tubes) □ C-Arm	□ Intraoral Dental ed outside your facility? ne Serial #: n ID: ation Date: □ Fluoro (I tube)	
□ Cone Beam CT Is this a Mobile Unit? □ Yes □ Machine Manufacture: Machine Model: Manufacture Date: Date of current calibration or se □ X-Ray (I tube) □ Cone Beam CT	□ CT No ervice evalua □ Rad/Flu □ CT	□ C-Arm Is this unit use Machi Room Installa ation report: uoro (2 tubes) □ C-Arm Is this unit use	□ Intraoral Dental ed outside your facility? ne Serial #: n ID: ation Date: □ Fluoro (I tube) □ Intraoral Dental	
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DUPLICATE THIS PAGE AS NEEDED

MOBILE SITE INFORMATION: (Complete only if you have mobile equipment used outs	ide of the registered facility.)
Site Name:	
Address, City, State, Zip	
Typical Schedule	
Equipment Description	
MOBILE SITE INFORMATION: (Complete only if you have mobile equipment used outside	de of the registered facility.)
Site Name:	
Address, City, State, Zip	
Typical Schedule	
Equipment Description	
Privacy Act Notice: Disclosure of your social security number on this application is red [13] and Iowa Code § 252J.8 (I). The number will be used in connection with the obligations and as an internal means to accurately identify licensees, and may be share allowed by law including Iowa Code § 421.18. NOTE: This does not apply to facilities only to facilities under a Sole Proprietorship.	collection of child support ed with taxing authorities as
am authorized to complete this application on behalf of the organization.	
As representative of the organization, I hereby certify and declare under penalty of perovided in this document, including any attachments, is true and correct. As organization, I am responsible for the accuracy of the information provided regard submits the application. I understand that providing false and misleading information may be cause for disciplinary action, denial, revocation, and/or criminal prochat a representative of the organization is responsible to update information submitted the information changes.	said representative of the less of who completes and tion in or concerning this osecution. I also understand
n submitting this application, the organization agrees to any reasonable inquiry that m clarify the information provided on or in conjunction with this application.	ay be necessary to verify or
understand this information is a public record in accordance with lowa Code chap nformation is public information, subject to the exceptions contained in lowa law.	eter 22 and that application
have read the Administrative Rules governing this license, permit, registration, or employees aware as required and will comply with those provisions.	certification and will make
Required	
Signature of Organizational Representative	Date

rev 18-Dec-23