NRC FORM 591M (04-2022)



Materials Inspection Report

1. Licensee/Location Inspected:			2. NRC/Regional Office			
IndyVet Emergency and Specialty Hospital 5425 Victory Dr. Indianapolis, IN 46203			Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352			
Report Number(s) 2	2024001					
3. Docket Number(s)		4. License Number(s)		5. Date(s) of Inspection		
030-37823		13-32698-01		April 17, 2024		
Nuclear Regulatory examinations of pro are as follows: 1. Based of 2. Previous 3. During twere as A. The vious identification were sailly and the sail of the sa	Commission (NRC) rules and regreedures and representative record on the inspection findings, no violates violation(s) closed. This inspection, certain of your active sessed at Severity Level IV, in accordation(s), specifically described to	ulations and the ds, interviews to tions were identified wities, as descrete description with the insertion was or is being the distribution was or is being the distribution with the distribution was or is being the distribution was an experienced with the distribution was or is being the distribution with the distribution was also as the distribution was also as the distribution was also as the distribution with the distribution was also as the distribution with the distribution was also as the distribution was	ne conditions of your license. The inswith personnel, and observations by ntified. ribed below and/or attached, were in the NRC Enforcement Policy. pector as non-cited violations, are ning taken, and the remaining criteria	the inspector. The inspection findings in violation of NRC requirements, and ot being cited because they were self-		
which r (Violati Contrary protectio committe	may be subject to posting in accorons and Corrective Actions) to 10 CFR 20.1101(c), the on program content and implement	dance with 10 licensee fail ementation adiation prot	CFR 19.11. led to periodically (at least and since the last inspection. As tection program review within	corrective action, the licensee		
		Statement of	f Corrective Actions			
actions is made in ac	ccordance with the requirements of 1	me to the Inspec 0 CFR 2.201 (c	ctor will be taken to correct the violations	e steps which will be taken, date when full		
TITLE	PRINTED NAME		SIGNAT	TURE AND DATE		
LICENSEE'S REPRESENTATIVE						
NRC INSPECTOR	Zahid Sulaiman, Health Physi	icist	Zahid M. Sulaiman	Digitally signed by Zahid M. Sulaiman Date: 2024.05.06 13:42:05 -05'00'		
BRANCH CHIEF Rhex Edwards, Chief, MIB		100	Digitally signed by RHEX EDWARDS Date: 2024.05.07 20:01:26 -05'00'			
			<u> </u>			

NRC FORM 592M (10-04-2022) SEAR REGUY					U.S. NI	ICLEAR REGULATORY COMMISSION		
	Mate	erials Insp	ection	Record				
1. Licensee Name:	2. Docket Number(s):			3. Licen:	3. License Number(s)			
IndyVet Emergency and Special	030-37823			13-32	13-32698-01			
4. Report Number(s):	•	5. Date(s) of Inspection:						
2024-001				April 17, 2024				
6. Inspector(s):		7. Program Code(s):		8. Priority:	9. Inspection Guidance Used:			
Zahid Sulaiman, Health Physicis		02400		5	87126			
10. Licensee Contact Name(s):	mail Address:		12. Licensee Telephone Number(s):					
Erin Hunt, RSO erin.hun		nt@nva.com			Work Cell: 317-893-6047			
13. Inspection Type: Initial 14.	Type: Initial 14. Locations Inspected: Hyl			rid 15. Next Inspection Date (MM/DD/YYYY):				
Routine Announced	✓ Main Office Field			04/17/2029		✓ Normal Extended		
Non-Routine			note			Reduced No change		
16. Location(s) Inspected List: Main Hospital: 5425 Victory Dr.,	Indianapoli	s, IN 46203	}					

17. Scope and Observations:

This was an unannounced, routine inspection of a multi-specialty veterinary hospital authorized to use Iodine-131 (I-131) for the treatment of thyroid disease in felines at a facility in Indianapolis, Indiana. The licensee employed three full-time veterinarians who served as authorized users of licensed material. Four veterinary assistants supported the authorized users. The licensee obtained its I-131 in liquid form, dispensed in a unit dose syringe, from a licensed radiopharmacy. Feline treatments were administered subcutaneously (on Mondays or Tuesdays) with unit dosages of I-131 based on the patient's weight (typically 3-5 mCi), with maximum dose of 5 mCi. The veterinary hospital administered 4-5 feline patient treatments per month. After administration of I-131, the feline patients were kept in the hospital, segregated in shielded cages within a secured room for three days and released on the fourth day. The licensee released the feline patients to the owners on Thursday or Fridays of the same week with verbal and written instructions on radiation safety precautions, feeding, care, litter waste management, etc. The hospital returned used syringes to the radiopharmacy, and used decay-in-storage techniques for all other radioactive waste. Performance Observations:

This inspection consisted of interviews with licensee personnel; a review of selected records; a tour of the hot lab, feline patient housing, and waste storage areas; and independent measurements. The inspection included observations of security of licensed material, use of personnel monitoring and protective clothing, area surveys, and packaging of radioactive waste for decay in-storage. The inspector had the staff demonstrate: (1) package surveys and check-in procedures, (2) dose calibration and dose assay, and (3) area surveys and wipe tests. The staff described feline patient dose administration, housing after treatment, feeding the felines, cleaning cages, and performing surveys of the cages. The inspector conducted independent surveys of restricted and unrestricted areas and found no residual contamination or exposure to members of the public over regulatory limits. The inspector found the licensee's staff knowledgeable of radiation protection principles and regulatory requirements through these observations, demonstrations, and other discussions.

The inspector reviewed the following records: sealed sources inventory and leak tests, radiation safety training, survey meter calibration, area surveys and wipe tests, feline patient release measurements, instrument quality

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Materials Inspection Record (Continued)

control, dose administration, waste disposal, and dosimetry records.

The inspector identified a SLIV violation of 10 CFR 20.1101(c), related to the failure to perform an annual radiation protection program audit since the last inspection. The violation is cited in the NRC Form 591M. As corrective action, the licensee committed to complete the 2023 annual radiation protection program review within 60-days and ensured the radiation protection program will be reviewed annually.

Signature and Date - Branch Chief

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Digitally signed by RHEX EDWARDS Date: 2024.05.07 20:01:05 -05'00'

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