

Materials Inspection Report

1. Licensee/Location Inspected:			2. NRC/Regional Office						
Parkland Health Center 1101 West Liberty Farmington, MO 63640 Report Number(s) 2022-001			Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352						
3. Docket Number(s) 4. Lid		4. License Nui	mber(s)	5. Date(s) of Inspection					
030-11341		24-16616-01		October 3, 2022					
Nuclear Regulatory examinations of pro are as follows:	Commission (NRC) rules and reg cedures and representative record	ulations and th ds, interviews v	r your license as they relate to radiation e conditions of your license. The insper vith personnel, and observations by the	ction consisted of selective					
1. Based on the inspection findings, no violations were identified.									
 2. Previous violation(s) closed. 3. During this inspection, certain of your activities, as described below and/or attached, were in violation of NRC requirements, and were assessed at Severity Level IV, in accordance with the NRC Enforcement Policy. 									
A. The violation(s), specifically described to you by the inspector as non-cited violations, are not being cited because they were self-identified, non-repetitive, corrective action was or is being taken, and the remaining criteria in the NRC Enforcement Policy were satisfied. (Non-cited violation(s) was/were discussed involving the following requirement(s)									
which r	lowing violation(s) is/are being cite may be subject to posting in accor ons and Corrective Actions)		ce with NRC Enforcement Policy. This f CFR 19.11.	orm is a NOTICE OF VIOLATION,					
		Statement of	Corrective Actions						
actions is made in ac	ccordance with the requirements of 1	0 CFR 2.201 (c	tor will be taken to correct the violations ide orrective steps already taken, corrective st o NRC will be required, unless specifically re	eps which will be taken, date when full					
TITLE	PRINTED NAME		SIGNATUR	E AND DATE					
LICENSEE'S REPRESENTATIVE									
NRC INSPECTOR	Ryan Craffey		Pof- Cofony	Digitally signed by Ryan J. Craffey Date: 2022.10.19 08:47:55 -04'00'					
BRANCH CHIEF Michael Kunowski		Michael A. Kunowski	Digitally signed by Michael A. Kunowski Date: 2022.10.21 07:49:55 -05'00'						
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NRC FORM 592M				U.S. NU	ICLEAR REGULATORY COMMISSION				
Materials Inspection Record									
1. Licensee Name:	2. Docket Num	2. Docket Number(s):		3. License Number(s)					
Parkland Health Center	030-1134	030-11341		24-16616-01					
4. Report Number(s):	•	5. Date(s) of Inspection:							
2022-001		October 3, 2022							
6. Inspector(s):		7. Program Code(s):		8. Priority:	9. Inspection Guidance Used:				
Ryan Craffey		02120		3	IP 87131				
10. Licensee Contact Name(s): 11. Licens	ee E-mail Address:			12. Licensee 1	Telephone Number(s):				
	.miller@bjc.org @bjc.org			573-760-8042					
13. Inspection Type: Initial 14. Locations In	nspected:		15. Next Inspection Date (MM/DD/YYYY):						
✓ Routine Announced ✓ Main Office Fig		d Office	10/03/2025		✓ Normal Extended				
Non-Routine ✓ Unannounced ☐ Temporary	Job Site Rem	mote		72025	Reduced No change				
16. Scope and Observations: Parkland Health Center was a 78-bed acute care hospital in Farmington, Missouri, authorized to perform diagnostic and therapeutic administrations of radiopharmaceuticals. At the time of the inspection, two full-time nuclear medicine technologists performed 10 or more diagnostic administrations per day on average using unit doses from a licensed radiopharmacy in St. Louis, as well as infrequent therapeutic administrations of I-131 capsules. The licensee retained the services of a medical physics consultant to audit the program quarterly. The inspector toured the hospital in Farmington. All areas were properly posted and all licensed material was adequately secured and accounted for. The inspector conducted independent and confirmatory surveys of the nuclear medicine department and found no residual contamination or exposures to members of the public in excess of regulatory limits. The inspector observed the conduct of a hepatobiliary scan using a unit dose of a To-99m radiopharmaceutical, as well as demonstrations of instrument checks, receipt of packages containing licensed material, area surveys, and decay-in-storage waste handling. The technologists were knowledgeable of radiation protection principles and regulatory requirements, and used adequate ALARA practices, personnel dosimetry, and calibrated and operable radiation detection instruments. The inspector reviewed documentation of the only I-131 therapy administered since the last routine inspection and a selection of records related to the radiation safety program, including consultant audits, dose calibrator and well counter quality control tests, sealed source inventories and leak test results, and personnel dosimetry reports. No violations of NRC requirements were identified as a result of this inspection.									

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