NRC FORM 591M PART 1 (10-2011) 10 CFR 2.201 U.S. NUCLEAR REGULATORY COMMISSION SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION									
1. LICENSEE/LOCATION INSPECTED: 2. NRC/REGIONAL OFFICE									
Missouri Delta Medical Center 1008 North Main Street Sikeston, MO 63801-5099			Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352						
REPORT NUMBER(3. DOCKET NUMBER(4. LICENSE NUMBER	(0)	5. DATE(S) OF INSPECTION					
030-02377)	24-12876-02		September 19, 2013					
LICENSEE: The inspection was an examination of the activities conducted under your license as they relate to radiation safety and to compliance with the Nuclear Regulatory Commission (NRC) rules and regulations and the conditions of your license. The inspection consisted of selective examinations of procedures and representative records, interviews with personnel, and observations by the inspector. The inspection findings are as follows: 1. Based on the inspection findings, no violations were identified. 2. Previous violation(s) closed.									
3. The viola non-repe									
	Non-cited violation(s) were discuss	ed involving the follo	wing requirement(s):						
 During this inspection, certain of your activities, as described below and/or attached, were in violation of NRC requirements and are being cited in accordance with NRC Enforcement Policy. This form is a NOTICE OF VIOLATION, which may be subject to posting in accordance with 10 CFR 19.11. (Violations and Corrective Actions) 									
Statement of Corrective Actions I hereby state that, within 30 days, the actions described by me to the Inspector will be taken to correct the violations identified. This statement of corrective actions is made in accordance with the requirements of 10 CFR 2.201 (corrective steps already taken, corrective steps which will be taken,									
date when full compliance will be achieved). I understand that no further written response to NRC will be required, unless specifically requested.									
TITLE LICENSEE'S REPRESENTATIVE	PRINTED NAME		SIGNATURE						
NRC INSPECTOR	Robert P. Hays	C	Zh QJ	9/19/13					
BRANCH CHIEF	Aaron T. McCraw		AT M	10/2/13					
IRC FORM 591M PART	1 (10-2011)		1110						

U.S. NUCLEAR REGULATORY COMMISSION (10-2011) 10 CFR 2.201 Docket File Information SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATION INSPECTE				ONAL OFFICE				
Missouri Delta Medical C 1008 North Main Street Sikeston, MO 63801-5099 REPORT NUMBER(S) 13-01			Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352					
3. DOCKET NUMBER(S)		4. LICENSE NUMBER(S)	;	5. DATE(S) OF INSPECTION			
030-02377	24-12876-02			September 19, 2013				
6. INSPECTION PROCEDURES USE	7. INSPECTION FOCUS	7. INSPECTION FOCUS AREAS						
87131		03.01-03.07	03.01-03.07					
SUPPLEMENTAL INSPECTION INFORMATION								
1. PROGRAM CODE(S)	2. PRIORITY	3. LICENSEE CONTAC	3. LICENSEE CONTACT		I. TELEPHONE NUMBER			
02121	5	L. Carleton, M.	D., RSO		(573) 472-7340			
✓ Main Office InspectionField Office Inspection		Next Inspection	Date:	09/19/2018	8			
Temporary Job Sit	e Inspection							
		PROGRAM SC	OPE					

The licensee was a medical institution authorized by the license to use any byproduct material as needed, for any study permitted by 10 CFR 35.100, and 35.200 at the location specified on the license.

The nuclear medicine department was staffed with one full-time nuclear medicine technologist (NMT). The NMT administered an average of 2 cardiac studies and 1-2 other studies per day. Xe-133 is used for lung studies and I-123 is used for uptake studies. The nuclear medicine department received unit doses only from a nearby Paducah, KY nuclear pharmacy. All waste was either held for decay-in-storage (DIS) or returned to the nuclear pharmacy as limited quantity shipments.

Performance Observations

The licensee's NMT (Bill Howard) demonstrated/discussed: (1) survey meter use and calibrations; (2) package checkin procedures; (3) unit dosage prep and safe use; (4) wipe test counting; (5) waste handling; (6) sealed source inventories and leak tests; (7) routine security of licensed material; (8) dose calibrator tests; (9) safety program audits; (10) any contamination events (none since previous inspection); (11) HAZMAT refresher training (Jan. 2013); (12) daily surveys and weekly wipe tests; and (13) dosimetry < 125 mrem DDE and <450 mrem SDE for 2012 and < 50 mrem DDE and <250 mrem SDE for 2013 (thru June 2013).

The inspector performed independent and confirmatory radiation measurements which indicated results consistent with licensee survey records and postings.