NRC FORM 591M PART 1 (06-2010)

U.S NUCLEAR REGULATORY COMMISSION

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION						
1. LICENSEE/LOC	CATION INSPECTED):	2. NRC/REGIONAL OFF	2. NRC/REGIONAL OFFICE		
Saint Margaret Mercy Healthcare Centers 5454 Hohman Avenue Hammond, IN 46320			Region III 2443 Warrenville R	U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road, Suite 210 Lisle, Illinois 60532-4351		
REPORT NUMBER(the state of the s					
3. DOCKET NUMBE 030-016		4. LICENSEE NUM 13-0	BER(S) 02047-02	5. DATE(S) OF INSP		
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 violation(s), specifically described to you by the inspector as non-cited violations, are not being cited because they were self-identified, non-repetitive, and corrective action was or is being taken, and the remaining criteria in the NRC Enforcement Policy, NUREG-1600, to exercise discretion, were satisfied Non-cited violation(s) were discussed involving the following requirement(s): 4. During this inspection certain of your activities, as described below and/or attached, were in violation of NRC requirements and are being cited. This form is a NOTICE OF VIOLATION, which may be subject to posting in accordance with 10 CFR 19.11						
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	Printed N	Name	Signature		Date	
LICENSEE'S REPRESENTATIVE						
NRC INSPECTOR	Robert P	. Hays	VALER	bur	11/15/10	
Branch Chief	Tamara E. I	Bloomer	Ben Lamber	* for	12/23/10	

NRC FORM 591 M PART 3 (06-2010)			U.S. NUCLEAR REGULATORY COMMISSION				
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	SAFETY IN	SPECTION RE	EPORT AND COMPLIANCE INSP	PECTION			
1. LICENSEE Saint Margar	et Mercy Healtho	care Centers	2. NRC/REGIONAL OFFICE U.S. Nuclear Regulatory Commission				
5454 Hohma	n Avenue		Region III 2443 Warrenville Road, Suite 210				
Hammond, IN 46320			Lisle, Illinois 60532-4351				
REPORT NUMBER(S) 2010-01							
		4. LICENSE NUMBER(S)		5. DATE(S) OF INSPECTION			
03001602 6. INSPECTION PROCEDURES		13-02047-02		November 15, 2010			
87131 (10/24/02)		7. INSPECTION FOCUS AREAS 03.01-03.07					
SUPPLEMENTAL INSPECTION INFORMATION							
1.PROGRAM	2. PRIORITY			4. TELEPHONE NUMBER			
2230	2	Terry Harrigan, Supervisor		219-932-2300, x. 45070			
 Main Office Inspection X Field Office Inspection South Campus, 24 Joliet Street, Dyer, IN ☐ Temporary Job Site Inspection 							
PROCRAM CCORE							

PROGRAM SCOPE

The licensee was a medical institution with two authorized locations in Hammond and Dyer, Indiana, with authorization by the license to use any byproduct materials for diagnostic and therapeutic medical procedures under 10 CFR 35.100, 35.200, 35.300, 35.400, and a HDR afterloader. The North Campus was reviewed during the previous inspection.

The South Campus is authorized by the license to use any byproduct materials for diagnostic and therapeutic medical procedures under 10 CFR 35.100, 35.200, 35.300, 35.400, and in-vitro studies. The licensee routinely conducts an average of 10-15 administrations/scans per day for routine diagnostic, imaging, and therapeutic procedures with a staff of 3 nuclear medicine technologists. The licensee receives all licensed material as unit doses from a local nuclear pharmacy as needed. Iodine-131 procedures average 1-2 cases per month. Dosages up to 150 millicuries are administered to thyroid carcinoma therapy patients who are released in accordance with 10 CFR 35.75, and are contacted at home for several days after the administration to ensure patient is following instructions. The licensee possesses cesium-137 sources for low-dose brachytherapy which have been in storage and not used since the previous inspection. The inspector performed independent and confirmatory radiation measurements which indicated results consistent with licensee survey records and postings.

Performance Observations

During the inspection, the licensee's NMTs (Mike Muceski, Chris Mully, Terry Harrigan) demonstrated/discussed: (1) survey meter use and calibrations; (2) package check-in procedures; (3) dosage prep; (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) quarterly radiation safety meetings and program audits; (10) any contamination events (none since the previous inspection); and (11) dosimetry (< 10% of regulatory limits) for CY 2009: 354mr-DDE; 660mr-finger; and 2010 YTD 2010: 143mr-DDE; 500mr-finger.