NRC FORM 591M PAR	T 1		U.S. NUCLEAR REGULATORY COMMISSION		
10 CFR 2.201 SAFET	Y INSPECTION F	REPORT AND CO	MPLIANCE INS	SPECTION	8
1. LICENSEE/LOCATION INSPECTED:			2. NRC/REGIONAL OFFICE		
Methodist Hospital of Gary, Inc. 8701 Broadway Merrillville, IN 46410			UNITED STATES NUCLEAR REGULATORY COMMISSION REGION III 2443 WARRENVILLE ROAD, SUITE 210 LISLE, IL 60532-4352		
REPORT  3. DOCKET NUMBER(S)	2006-001	LICENSEE NUMBER(S)	<del></del>	5. DATE(S) OF	INSPECTION
030-1	1234	13-165	5558-01 April 18 2006		2006
LICENSEE:	<u> </u>			1, 10, 11, 12, 1	7000
Nuclear Regulatory Comof procedures and representations of the second se	emission (NRC) rules and a sentative records, interview e inspection findings, no vio lation(s) closed. (s), specifically described to and corrective action was or ion, were satisfied.	regulations and the conditions with personnel, and obstations were identified.	ons of your license. The ervations by the inspect when he keep the	adiation safety and to comple inspection consisted of setor. The inspection findings  CAMPUS,  ot being cited because they all Enforcement Policy, NUR  at(s) and Corrective Action(s)	lective examinations are as follows:  ""  "  were self-identified, EG-1600, to
cited. This form	nspection certain of your actions a NOTICE OF VIOLATION and Corrective Actions)	tivities, as described below DN, which may be subject to	and/or attached, were in posting in accordance	n violation of NRC requirements with 10 CFR 19.11.	nts and are being
1					
	Licensee's	Statement of Correct	ive Actions for Item	4, above.	
corrective actions is mad date when full compliand Title	le in accordance with the re e will be achieved). I under	quirements of 10 CFR 2.20	<ol> <li>(corrective steps alrea response to NRC will be</li> </ol>	the violations identified. This idy taken, corrective steps wi e required, unless specifically Signature	hich will be taken,
LICENSEE'S REPRESENTATIVE					
NRC INSPECTOR	Robert P. I	Hays			4/18/06
NRC FORM 591M PART 1			Very T	<del>~~~7\                                  </del>	14.0/20

NRC FORM 591M PAF (10-2003) 10 CFR 2.201	<b>RT 3</b>	Docket Fi SAFETY INSP AND COMPLIA	U.S. NUCLEAR REGULATORY COMMISSION DCKET FILE Information TY INSPECTION REPORT DMPLIANCE INSPECTION		
1. LICENSEE  Methodist Hospital of Gary, Inc. REPORT NUMBER(S) 2006-001			2. NRC/REGIONAL OFFICE  Region III		
3. DOCKET NUMBER(S)		4. LICENSE NUMBER(S	S)	5. DATE(S) OF INSPECTION	
03011234		13-1	6558-01	April 18, 2006	
6. INSPECTION PROCEDURES USED 87131, 87132		7. INSPECTION FOCUS AREAS  03.01 - 03.07			
07101,	07102		SPECTION INFORMA	TION	
1. PROGRAM CODE(S)  02240	2. PRIORITY	3. LICENSEE CONTAC			
Main Office Inspection			Next Inspec	tion Date: April 2008	
X Field Office Temporary Jo	Northlake Can	npus, 600 Grant S	treet, Gary, IN		
· · ·		PROGI	RAM SCOPE		
with authoriza medical proce stereotactic ra The Northlake	tion by the lice dures under 10 idiosurgery, Gli Campus routin	nse to use any by CFR 35.100, 35.2 aSite® procedure ely conducts an a	product materia 00, 35.300, 35.40 s and investiga average of 2-6 a	ons in Merrillville and Gary, Indiana, als for diagnostic and therapeutic 00, 35.500, HDR afterloader, gamma tional new drug testing. dministrations/scans per day for a staff of 1-2 nuclear medicine	

routine diagnostic, imaging, and therapeutic procedures with a staff of 1-2 nuclear medicine technologists, depending on the number of patients scheduled, who rotate from the Merrillville IN, facility. The licensee receives all licensed material as unit doses and a 50 mCi Tc-99m bulk vial for on call as needed from a local nuclear pharmacy. No thyroid carcinoma therapy is conducted at this authorized location of use. Samarium-153 is administered infrequently.

No brachytherapy procedures are conducted at this facility, although the licensee possesses brachytherapy cesium-137 sources and a strontium-90 eye applicator which have been in storage.

## **Performance Observations**

During the inspection, the licensee's available staff demonstrated/discussed: (1) survey instruments and required surveys; (2) package receipt and check-in procedures; (3) wipe test counting; (4) dosimetry; (5) waste handling; (6) sealed source inventories and leak tests; (7) routine security of licensed material; (8) dose calibrator tests; (9) written directives; and (10) radiation safety committee meetings and radiation safety program audit results.