NRC FORM 591M PAR	T 1				U.S NUCLEAR RE	GULATORY COMMISSION
(06-2010) 10 CFR 2.201						
SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION						
1. LICENSEE/LOCATION INSPECTED:			2. NRC/REGIONAL OFFICE			
Bay Regional Medical Center 315 E. Warwick Drive			U.S. Nuclear Regulatory Commission, Region III 2443 Warrenville Road, Suite 210			
Alma, Michigan			Lisle, Illinois 60532			
REPORT NUMBER(S): 11-02			, in the second			
3. DOCKET NUMBER(S) 4. LIG		4. LICENSEE NUMBER(NUMBER(S) 5. DATE			PECTION
030-13900		21-18585-01			November	15,2011
LICENSEE:						
Regulatory Commission	(NRC) rules and regulation	es conducted under your lic ons and the conditions of your and observations by the in	our license.	he inspection consist	ted of selective exami	
1. Based on	the inspection findings	s, no violations were ide	ntified.			
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						
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						,
			•		•	
		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 Name		Sign	ature	Date
LICENSEE'S REPRESENTATIVE						
NRC INSPECTOR	Andrew M. Bran	nnik		Andaly	K Benif	11/15/2011
Branch Chief	Tamara E. Bloor	mer			T Fan	11 25 11

U.S. NUCLEAR REGULATORY COMMISSION NRC FORM 591 M PART 3 (06-2010)10 CFR 2.201 Docket File Information SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION 2. NRC/REGIONAL OFFICE 1. LICENSEE **Bay Regional Medical Center** U.S. Nuclear Regulatory Commission, Region III 2443 Warrenville Road, Suite 210 Lisle, Illinois 60532 REPORT NUMBER(S) 11-02 5. DATE(S) OF INSPECTION 3. DOCKET NUMBER(S) 4. LICENSEE NUMBER(S) November 15, 2011 030-13900 21-18585-01 6. INSPECTION PROCEDURES 7. INSPECTION FOCUS AREAS 87131 03.01 - 03.07SUPPLEMENTAL INSPECTION INFORMATION 4. TELEPHONE NUMBER 1.PROGRAM 3. LICENSEE CONTACT 2. PRIORITY 989-894-3000 2120 Tvre K. Jones, M.D. – RSO Next Inspection Date: October 2014 (unchanged) Main Office Inspection Field Office Inspection 315 E. Warwick Drive, Alma, MI

PROGRAM SCOPE

This was a routine inspection at a second authorized location for Bay Regional Medical Center. An NRC inspector had originally intended to inspect this location during a routine inspection of the licensee in October 2011, but was unable to visit the site. The facility was added to Bay Regional Medical Center's NRC license in May 2011, concurrent with the removal of the facility as a location of use for another NRC licensee (Michigan CardioVascular Institute, NRC license no. 21-26447-01).

This facility was a standalone clinic located in Alma, Michigan, and was authorized only for activities under Title 10 of the Code of Federal Regulations (10 CFR) Section 35.200. One full time nuclear medicine technologist performed five cardiac procedures per day between Monday and Thursday. The licensee obtained licensed material as unit doses from an area nuclear pharmacy, and did not use xenon-133, bulk doses, or molybdenum/technetium generators.

PERFORMANCE OBSERVATIONS

No administrations of byproduct materials were available for observation during the inspection. Interviews of available staff revealed an adequate level of understanding of emergency and material handling procedures and techniques. Dose calibrator constancy checks, package receipt, daily surveys, and waste handling and disposal procedures were successfully demonstrated. An outside consultant performed quarterly program audits that were adequate to oversee the program.

Licensed material was adequately secured and not readily accessible to members of the general public. The licensee possessed a radiation survey meter that was calibrated, operational, and performed well in side-by-side comparison with an NRC instrument. Independent measurements did not indicate readings in excess of 10 CFR Part 20 limits in restricted or unrestricted areas. Personal whole body and extremity dosimetry were observed being worn by the staff during the inspection, and records did not indicate doses in excess of 10 CFR Part 20 limits. Dosimetry records indicated that the highest annual whole body and extremity readings for the year to date were 70 millirem (mrem) and 320 mrem, respectively.

No violations were identified during this inspection.

Temporary Job Site Inspection

