NRC FORM 591M PART 1 U.S. NUCLEAR REGULATORY COMMISSION								
10 CFR 2.201 SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATIO	ON INSPECTED:	2. NRC/REGIONAL OFFICE	2. NRC/REGIONAL OFFICE					
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The Community 901 MacArthur B	-		•	Region III				
Munster, IN 4632			_	U. S. Nuclear Regulatory Commission				
Munster, IN 46321			Lisle, IL 60532-4352	2443 Warrenville Road, Suite 210				
REPORT NUMBER(S	s) 2016001	Bisic, 11 00332-4332	Bisic, 11 00332-4332					
3. DOCKET NUMBER(S		4. LICENSE NUMB	ER(S)	5. DATE(S) OF INSPECTION	V.			
030-09964		13-15882-01		Oct. 17, 2016				
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	2. Previous violation(s) closed.							
non-repet	3. The violations(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, to exercise discretion, were satisfied.							
	Non-cited violation(s) were discussed involving the following 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	PRINTED NAME		SIGNATURE		DATE			
LICENSEE'S REPRESENTATIVE								
NRC INSPECTOR	Deborah A. Piskura, Senior Heal	th Physicist	Delivered A	A8 Rua	10/17/16			
BRANCH CHIEF	Aaron T. McCraw, Chief, MIB		Mr. K		11/16			

NRC FORM 591M PART 1 (07-2012)

NRC FORM 591M PART 3	-		II S NII	CLEAR REGULATORY COMMISSION				
(07-2012) Docket File Information								
SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATION INSPECT	ED:		2. NRC/REGIONAL OFFICE					
The Community Hospital			Region III					
901 MacArthur Boulevard	1		U. S. Nuclear Regulatory Commission					
Munster, IN 46321			2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352					
REPORT NUMBER(S) 20160	Ω1		Lisic, IL 00332-4332	2				
3. DOCKET NUMBER(S)	01	4. LICENSE NUMBER(S	5. DATE(S) OF INSPECTION					
030-09964		13-15882-01 Oct		Oct. 17, 2016				
6. INSPECTION PROCEDURES US	ED	7. INSPECTION FOCUS AREAS						
87130, 87131, & 87132		03.01 - 03.07						
	SUPPLEM	ENTAL INSPECT	ON INFORMATION					
1. PROGRAM CODE(S)	2. PRIORITY	3. LICENSEE CONTAC	Т	4. TELEPHONE NUMBER				
02240	2	Santosh K. Kar,	M.S., RSO	(219) 836-7351				
✓ Main Office Inspec	ction	Next Inspection	Date: Oct. 17, 20	018				
✓ Field Office Inspec	tion 801 MacArth	nur Blvd, Munster,	IN					
☐ Temporary Job Sit	te Inspection							
PROGRAM SCOPE								
•	_	• •		naterial permitted by 10 CFR				
•		, ,		(Y-90) microspheres. The				
licensee conducted licens								
Collectively, the nuclear								
				ous iodine-131 (I-131) dosages nsee received unit doses and				
bulk technetium-99m(Tc-99m) for imaging studies and I-131 therapy capsules from a licensed radiopharmacy.								
The radiation oncology d	epartment was staffe	ed with three AMP	s, two dosimetrists, and t	hree AUs. The licensee				
administered approximately 8-10 patient cases annually utilizing its HDR. These treatments were for GYN cancer								
cases. All HDR patient treatments were administered by the attending radiation oncologist and the AMP. The licensee								
maintained an inventory of cesium-137 tube sources in secured storage; these sources had not be used for several years.								
Occasionally, the licensee performed permanent prostate implants (2-3 cases/year). The department administered all								
radiopharmacuetical treatments for cancers including; 5-7 Y-90 SIRSpheres treatments/year; 2-3 Xofigo cases/year;								
12-15 I-131 cancer treatments/year; and 1-2 Zevalin treatments/year.								
This inspection consisted of interviews with licensee personnel, a review of selected records (including permanent								
implant treatment plans), a tour of the nuclear medicine and radiation oncology departments, and independent								
measurements. The inspection included observations of dose calibrator QA checks and HDR safety checks, security of								
licensed material, package surveys, inventories of sealed sources, and use of personnel monitoring. The inspector								
observed the licensee personnel prepare, assay, and administer three dosages for diagnostic testing procedures.								
No violations of NRC requirements were identified during this inspection.								
and makes were								