NRC FORM 591M PART 1 U.S. NUCLEAR REGULATORY COMMISSION								
10 CFR 2.201 SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATION INSPECTED:			2. NRC/REGIONAL OFFICE					
Dr. Anil K. Goel			Region III					
1915 E. 14 Mile Road			U. S. Nuclear Regulatory Commission					
Birmingham, Mic			2443 Warrenville Road, Suite 210					
Diffingum, Mongun 40007			Lisle, IL 60532-4352					
REPORT NUMBER(S	3) 2012-001							
		4. LICENSE NUMBER	R(S)	5. DATE(S) OF INSPECTION	1			
030-37802		21-32683-01		Sept. 10, 2012				
LICENSEE:		•	· -	•				
The inspection was a	n examination of the activities conduct							
	on (NRC) rules and regulations and the sentative records, interviews with pers							
	the inspection findings, no violations v			-				
2. Previous								
non-repet	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 discuss	sed involving the foll	owing requirement(s):					
		•	· (,,					
			•					
5			and the control of th	CNDO	al a see to de se			
	s inspection, certain of your activities, cordance with NRC Enforcement Poli							
with 10 Cl	FR 19.11.	-,	· · · · · · · · · · · · · · · · · · ·	.,,				
(Violations	s 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	Geoffrey Warren		Dy Nu		9/10/12			
BRANCH CHIEF	Tamara Bloomer		MIM	Fu-	9/20/12			

NRC FORM 591M PART 3			U.S. N	UCLEAR REGULATORY COMMISSION				
(07-2012) 10 CFR 2.201		Docket File Inf	ormation					
SAFE	ETY INSPEC	CTION REPORT AN	D COMPLIANCE IN	ISPECTION				
1. LICENSEE/LOCATION INSPEC	TED:		2. NRC/REGIONAL OFFICE					
Dr. Anil K. Goel 1915 E. 14 Mile Road Birmingham, Michigan 48009			Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352					
REPORT NUMBER(S) 2012- 3. DOCKET NUMBER(S)	.001	A LICENSE MIMBED	/e\	5. DATE(S) OF INSPECTION				
3. DOCKET NUMBER(S) 030-37802		4. LICENSE NUMBER: 21-32683-01	(8)	September 10, 2012				
6. INSPECTION PROCEDURES US	3ED	7. INSPECTION FOCU	7. INSPECTION FOCUS AREAS					
87130		03.01 - 03.07	03.01 - 03.07					
SUPPLEMENTAL INSPECTION INFORMATION								
1. PROGRAM CODE(S)	2. PRIORITY	3. LICENSEE CONTAI		4. TELEPHONE NUMBER				
02220	3	Dr. Anil Goel, N	M.D.	(248) 723-4777				
✓ Main Office Inspe	ction	Next Inspection	n Date: Sept. 2	015				
Field Office Inspection								
Temporary Job S	ite Inspection		7.700	No. company of the Control of the Co				
PROGRAM SCOPE								
The licensee was a cardiology clinic located in Birmingham, Michigan, with authorization to use byproduct materials in Sections 35.100 and 35.200. Licensed activities were conducted only at the location indicated on the license. While authorized to perform work at temporary job sites, the licensee had never done so and had no plans to begin in the near future. Licensed activities were conducted one day each week, alternating Mondays and Wednesdays. The nuclear medicine department was staffed with one contract nuclear medicine technologist who also performed work for other licensees. The technologist typically administered 10 cardiac doses daily. The clinic received unit doses as needed from a licensed nuclear pharmacy. All waste was either held for decay-in-storage (DIS) or returned to the nuclear pharmacy.								
Performance Observations								
The inspector observed two diagnostic administrations of licensed material, including dose preparation and disposal. The technologist demonstrated survey meter and wipe counter QC, package receipt and return surveys and wipes, dose calibrator constancy, and daily contamination surveys and wipes, and described spill procedures and other activities. The inspector noted no concerns with these activities. Interviews with licensee personnel indicated adequate knowledge of radiation safety concepts and procedures. Review of dosimetry records indicated no doses of regulatory concern. The inspector performed independent and confirmatory radiation measurements which indicated results consistent with licensee survey records and postings.								