

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED: Rochester Medical Center, P.C. 543 N. Main Street Rochester, MI 48308 REPORT NUMBER(S) 12-01		2. NRC/REGIONAL OFFICE Region III U. S. Nuclear Regulatory Commission 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352	
3. DOCKET NUMBER(S) 030-32081	4. LICENSE NUMBER(S) 21-26287-01	5. DATE(S) OF INSPECTION 10/2/12	

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:

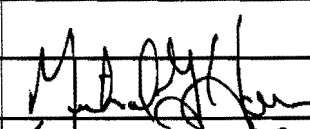
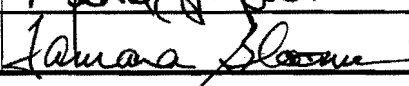
- 1. Based on the inspection findings, no violations were identified.
- 2. Previous violation(s) closed.
- 3. The violation(s), specifically described to you by the inspector as non-cited violations, are not being cited because they were self-identified, non-repetitive, and corrective action was or is being taken, and the remaining criteria in the NRC Enforcement Policy, 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 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	Michael G. Herr, CHP		10-17-12
BRANCH CHIEF	Tamara E. Bloomer		10-18/12

Docket File Information
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6. INSPECTION PROCEDURES USED 87130	7. INSPECTION FOCUS AREAS 03.01-03.08
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SUPPLEMENTAL INSPECTION INFORMATION

1. PROGRAM CODE(S) 02201	2. PRIORITY 5	3. LICENSEE CONTACT Won Chae, MD	4. TELEPHONE NUMBER (248) 651-9200
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Main Office Inspection Next Inspection Date: October 2017

Field Office Inspection

Temporary Job Site Inspection

PROGRAM SCOPE

On October 2, 2012, the U.S. Nuclear Regulatory Commission (NRC) conducted a routine inspection at Rochester Medical Center, P.C. located at 543 N. Main Street, Rochester, Michigan. The other location where licensed materials are authorized to be used, 601 N. Main Street, Rochester, Michigan was not open at the time of the inspections. License activities on occur on Monday and Fridays at the 601 N. Main Street location.

The licensee is small diagnostic clinic located in Rochester, Michigan, authorized to administer materials under Sections 35.100 and 35.200. The licensee employees two full time nuclear medicine technologists and one part-time technologist. The licensee's nuclear medicine technologists typically perform approximately 30 diagnostic procedures per week; with the majority of scans being cardiac studies. The licensee receives unit doses as needed from a local licensed nuclear pharmacy.

The nuclear medicine technologist was observed while preparing and injecting a patient with Tc-99m for a resting cardiac imaging. These technologist conducted these procedure using safe work practices which included the use of syringe shields and gloves. Labels on the syringe and the packaging for the dose were also observed. The waste was properly disposed of in a container for decay in storage.

The nuclear medicine technologist demonstrated package receipt procedures, daily checks performed on instrumentation, and waste disposal surveys. Response to spills, package contamination, and action levels were also discussed with the technologist. Her responses to scenarios were adequate and acceptable.

Selected records that were reviewed by the inspector included; dosimetry, audits, waste disposal, source inventory, and survey instrument calibration.

No violations of regulatory requirements were found.