

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED:

My Heart Partners, P.C.
3078 Niles Road
St. Joseph, MI 49085

REPORT NUMBER(S) 2012001

2. NRC/REGIONAL OFFICE

Region III
U. S. Nuclear Regulatory Commission
2443 Warrenville Road, Suite 210
Lisle, IL 60532-4352

3. DOCKET NUMBER(S)

030-35952

4. LICENSE NUMBER(S)

21-32375-01

5. DATE(S) OF INSPECTION

September 12, 2012

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.

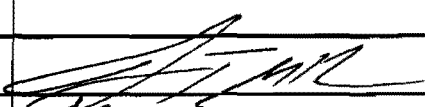
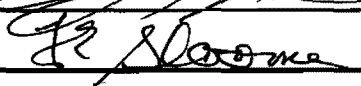
Non-cited violation(s) were discussed involving the following requirement(s):

- ☐ 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	Aaron T. McCraw		9/12/12
BRANCH CHIEF	Tamara E. Bloomer		9/26/12

Docket File Information

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3. DOCKET NUMBER(S) 030-35952	4. LICENSE NUMBER(S) 21-32375-01	5. DATE(S) OF INSPECTION September 12, 2012	
6. INSPECTION PROCEDURES USED IP 87130	7. INSPECTION FOCUS AREAS 03.01-03.08		

SUPPLEMENTAL INSPECTION INFORMATION

1. PROGRAM CODE(S) 02201	2. PRIORITY 5	3. LICENSEE CONTACT Dawn Edwards, RSO	4. TELEPHONE NUMBER (734) 662-3197
<input checked="" type="checkbox"/> Main Office Inspection Next Inspection Date: 09/12/2017			
<input type="checkbox"/> Field Office Inspection			
<input type="checkbox"/> Temporary Job Site Inspection			

PROGRAM SCOPE

This was a routine inspection of a cardiology clinic in St. Joseph, Michigan. The licensee was authorized to use materials permitted by 10 CFR 35.100 and 35.200. The licensee performed studies Monday through Friday between 8:00am and 4:30pm. On average, the licensee served 2-3 patients per day, both resting and stress studies for each patient. The licensee used only unit doses received from a local radiopharmacy. The licensee employed one nuclear medicine technologist. The licensee retained the services of a consulting physicist who serves as the radiation safety officer and reviews the program quarterly.

PERFORMANCE OBSERVATIONS

During this inspection, the inspector interviewed the nuclear medicine technologist; observed package return surveys and wipes; observed the technologist conduct the end of the day surveys; and reviewed select records. The inspector reviewed instrumentation records, quarterly audit reports, training records, waste disposal records, and dosimetry records. Dosimetry records for 2011 showed that the highest doses received were 213 mrem whole body and 1063 mrem extremity. The licensee had completed all studies for the day by the time of the inspection; therefore, the inspector could not observe dose preparation, dose assay, or patient injection. The technologist demonstrated a dose calibrator constancy check and described spill procedures.

No violations were identified during this inspection.