

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED: Wyandotte Internal Medicine Associates, PLLC 1700 Biddle Avenue Wyandotte, MI 48192		2. NRC/REGIONAL OFFICE U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road Suite 210 Lisle, Illinois 60532-4351	
REPORT NUMBER(S)	2010/001		
3. DOCKET NUMBER(S) 030-36514	4. LICENSEE NUMBER(S) 21-32495-01	5. DATE(S) OF INSPECTION June 10, 2010, with continued NRC in office review through 7/6/2010	

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, NUREG-1600, to exercise discretion, were satisfied.

_____ Non-Cited Violation(s) was/were discussed involving the following requirement(s) and Corrective Action(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.
(Violations and Corrective Actions)

Licensee's Statement of Corrective Actions for Item 4, above.

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	E. Kulzer/R. Gattone	<i>E. Kulzer / Robert R. Gattone, Jr.</i>	7/12/2010

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6. INSPECTION PROCEDURES USED 87130	7. INSPECTION FOCUS AREAS 03.01-03.07		

SUPPLEMENTAL INSPECTION INFORMATION

1. PROGRAM CODE(S) 02201	2. PRIORITY 5	3. LICENSEE CONTACT Subrahmanya S. Yellayi, M.D.	4. TELEPHONE NUMBER 313-928-4040
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Main Office Inspection Next Inspection Date: July 2015

Field Office _____

Temporary Job Site Inspection _____

PROGRAM SCOPE

The routine inspection began on June 9, 2010, with in office review through July 8, 2010. The in-office-review included receipt and review of information about how the licensee interpreted removable contamination survey results. The licensee was a medical clinic located in Wyandotte, Michigan, and authorized to use any byproduct material permitted by 10 CFR 35.100 and 35.200. Licensed activities were conducted in the nuclear medicine scanning room at the authorized location. The nuclear medicine department was staffed with one nuclear medicine technologist (NMT). The NMT routinely conducted about 10 cardiac studies per day on Mondays and Thursdays between 6 am and 3 pm. The licensee received unit doses from a local licensed nuclear pharmacy. All waste was held for decay in storage (DIS) or returned to the nuclear pharmacy as a limited quantity shipment. No change in ownership or NMT had occurred since the previous inspection.

Performance Observations

During the inspection, the licensee's NMT demonstrated/discussed : (1) survey meter use; (2) package check-in procedures and wipe test counting; (3) dosimetry; (4) safe handling techniques; (5) security; (6) radiation safety program audits; (7) survey instrument calibration; (8) sealed source inventory; (9) leak tests; (10) radiation surveys; (11) dosage preparation and administration; and (12) dose calibrator constancy checks.