

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED: Harrison County Hospital 1141 Hospital Drive, N. W. Corydon, IN 47112 REPORT NUMBER(S)	2. NRC/REGIONAL OFFICE U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road Suite 210 Lisle, Illinois 60532-4351
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3. DOCKET NUMBER(S) 030-28886	4. LICENSEE NUMBER(S) 13-23555-01	5. DATE(S) OF INSPECTION November 20, 2008
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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:

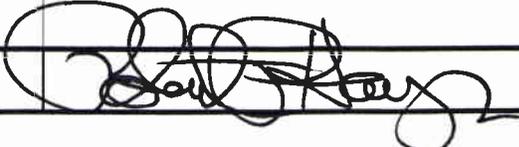
- 1. Based on the inspection findings, no violations were identified.
- 2. Previous violation(s) closed.
- 3. The violation(s), specifically described to you by the inspector as non-cited violations, are not being cited because they were self-identified, non-repetitive, and corrective action was or is 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	Robert P. Hays		11/20/08



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1. LICENSEE Harrison County Hospital REPORT NUMBER(S) 2008-001		2. NRC/REGIONAL OFFICE Region III 2443 Warrenville Road, Suite 210 Lisle, IL 60532	
3. DOCKET NUMBER(S) 03028886	4. LICENSE NUMBER(S) 13-23555-01	5. DATE(S) OF INSPECTION November 20, 2008	
6. INSPECTION PROCEDURES USED 87130 (10/24/02)		7. INSPECTION FOCUS AREAS 03.01-03.07	

SUPPLEMENTAL INSPECTION INFORMATION

1. PROGRAM CODE(S) 02120	2. PRIORITY 3	3. LICENSEE CONTACT Adam Miller, Chief NMT	4. TELEPHONE NUMBER 812-738-7891
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Main Office Inspection Next Inspection Date: **November 2011**

Field Office _____

Temporary Job Site Inspection _____

PROGRAM SCOPE

The licensee was a medical institution located in Corydon, Indiana, with authorization by the license to use any byproduct material as needed permitted by 10 CFR 35.100, 35.200, excluding generators, and 35.300, not to exceed 1 curie, excluding iodine-131 for thyroid carcinoma therapy. Two FT nuclear medicine technologists (NMTs) routinely perform an average of 2 diagnostic and four cardiac studies 0 administrations per week, with the majority of tests being bone scans. Other studies/scans as ordered, including cardiac studies. No I-131 dosages greater than 30 millicuries are administered and average 1 procedure per quarter. Licensed material is received as unit doses from a Louisville, KY, nuclear pharmacy. No change in RSO since the previous inspection. Area surveys of the injection, hot lab work areas, stress test, and scanning areas did not reveal any unusual or elevated readings.

Performance Observations

During the inspection, the licensee's NMTs demonstrated/discussed: (1) survey meter use and calibration; (2) package receiving and check-in procedures; (3) unit dose prep and safe use; (4) package wipe test counting; (5) dosimetry; (6) dose calibrator tests; (7) waste handling; (8) security of licensed material; (9) sealed source inventory; (10) radiation safety program audits; (11) iodine-131 procedures and written directives; and (12) any contamination events – none.