NRC FORM 591M PART 1
(10-2003) 10 CFR 2.201

U.S. NUCLEAR REGULATORY COMMISSION

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED:	2. NRC/REGIONAL OFFICE
American Health Network of Indiana, LLC 10333 N. Meridian Street, Suite 450 Indianapolis, IN 46290 6820 PARKDALE PL., INDIANAPOLIS, IN INSPECTED REPORT NUMBER(S) 2008-001	U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road Suite 210 Lisle, Illinois 60532-4351

3. DOCKET NUMBER(S)	4. LICENSEE NUMBER(S)	5. DATE(S) OF INSPECTION
030-36851	13-32555-01	11/05/08

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:

- [x] 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	S. J. Mulay	S J Mulay	11/05/08

NRC FORM 591M PART 3
(10-2003) 10 CFR 2.201

Docket File Information

U.S. NUCLEAR REGULATORY COMMISSION

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE **American Health Network of Indiana, LLC** REPORT NUMBER(S) **2008-01**	2. NRC/REGIONAL OFFICE **Region III** **2443 Warrenville Road, Suite 210** **Lisle, IL 60532**

3. DOCKET NUMBER(S)	4. LICENSE NUMBER(S)	5. DATE(S) OF INSPECTION
030-36851	**13-32555-01**	**November 05, 2008**

6. INSPECTION PROCEDURES USED	7. INSPECTION FOCUS AREAS
87131	**03.01-03.07**

SUPPLEMENTAL INSPECTION INFORMATION

1. PROGRAM CODE(S)	2. PRIORITY	3. LICENSEE CONTACT	4. TELEPHONE NUMBER
2120	**G3**	**Ed Wroblewski, RSO**	**765-987-0365**

☐ Main Office Inspection — Next Inspection Date: **November 2011**

☒ Field Office **6820 Parkdale Place, Suite 105, Indianapolis, IN**

☐ Temporary Job Site Inspection

PROGRAM SCOPE

This active facility performs about 100 diagnostic PET/CT imaging studies monthly utilizing florine-18 (F-18) and F-18 as FDG in whole body PET scans as well as PET brain and bone imaging. The facility is staffed by two full-time technologists and unit doses are received from an area nuclear pharmacy. This location became operational in June 2007. Nuclear medicine procedures (diagnostic or therapeutic) are not performed at this location. The licensee also maintains an authorized location in Kokomo, IN, which was reviewed during a 2005 inspection and was not reviewed at this time.

Performance Observations

Interviews conducted with available staff revealed an adequate level of understanding of emergency and material handling procedures and techniques. A patient injection for F-18 whole-body imaging was observed with no problems noted. Package surveys and wipes, and dose calibrator constancy checks were demonstrated with no regulatory issues identified. Compliance audits are performed at least quarterly by RSO/consultant which appear to adequately oversee program activities.

Personal dosimetry records were reviewed for 2007 and revealed whole-body readings of 606 mRem and extremity readings of 529 mRem. YTD 2008 records indicated whole-body and extremity measurements of 772 mRem and 527 mRem respectively.

Licensed material was observed well secured and/or under surveillance in the hot-lab area and was not readily accessible to members of the general public. Independent measurements taken in restricted and unrestricted areas of the hot-lab and imaging areas did not indicate readings above expected.