NRC FORM 591M PART 1
(06-2010)
10 CFR 2.201

U.S NUCLEAR REGULATORY COMMISSION

SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE/LOCATION INSPECTED:	2. NRC/REGIONAL OFFICE
Clarian Arnett Health System, Inc. 5165 McCarty Lane Lafayette, IN 47905 REPORT NUMBER(S) 2010-001	U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4351

3. DOCKET NUMBER(S)	4. LICENSEE NUMBER(S)	5. DATE(S) OF INSPECTION
030-37189	13-32535-02	~~December~~ November 30, 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) 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. This form is a NOTICE OF VIOLATION, which may be subject to posting in accordance with 10 CFR 19.11

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	Deborah A. Piskura	Deborah A Piskura	~~12/30~~ 11/30/2010
Branch Chief	Tamara E. Bloomer	TE Bloomer	12/9/10

NRC FORM 591 M PART 3
(06-2010)
10 CFR 2.201

U.S. NUCLEAR REGULATORY COMMISSION

Docket File Information
SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE	2. NRC/REGIONAL OFFICE
Clarian Arnett Health System, Inc. 5165 McCarty Lane Lafayette, IN 47905 REPORT NUMBER(S) 2010-001	U.S. Nuclear Regulatory Commission Region III 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4351

3. DOCKET NUMBER(S)	4. LICENSE NUMBER(S)	5. DATE(S) OF INSPECTION
030-37189	13-32535-02	Nov. 30, 2010

6. INSPECTION PROCEDURES	7. INSPECTION FOCUS AREAS
87130 & 87131	03.01-03.08

SUPPLEMENTAL INSPECTION INFORMATION

1.PROGRAM	2. PRIORITY	3. LICENSEE CONTACT	4. TELEPHONE NUMBER
02200	3	Rodney A. Dunseath, D.O., RSO	765-448-8122

⊘ Main Office Inspection Next Inspection Date: Nov. 2013
Field Office Inspection
☐ Temporary Job Site Inspection _____

PROGRAM SCOPE

This licensee was a medical institution (150+ bed hospital) with authorization for materials in Sections 35.100, 35.200, and 35.300. The licensee's consulting RSO audited the radiation safety program on a quarterly basis.

The nuclear medicine department was staffed with 4 technologists who performed approximately 250-300+ diagnostic nuclear medicine procedures per month; this work included a full spectrum of diagnostic imaging studies. The licensee received unit doses and bulk Tc-99m from a local radiopharamacy. The department maintained an active therapy program and administered numerous I-131 dosages for CA, whole body follow up studies, and hyperthyroidism (capsules only).

This inspection consisted of interviews with select licensee personnel; a review of select records; tour of the nuclear medicine department; and independent measurements. The inspector observed the administration of several diagnostic nuclear medicine procedures. The inspection included observations of dose calibrator QA checks, security of byproduct material, use of personnel monitoring, package receipts/returns, and surveys.