						EGULATORY COMMISSION		
(06-2010) 10 CFR 2.201								
SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATION INSPECTED:				2. NRC/REGIONAL OFFICE				
Medi-Physics, I d/b/a GE Health			U.S. Nuclear Regulatory Commission Region III					
1623 Lotsie Blvd.			2443 Warrenville Road, Suite 210					
Overland, MO 6	3132		Lisle, IL 60532-4351					
REPORT NUMBER(S) 2010-001								
3. DOCKET NUMBER(S 030-36453			5. DATE(S) OF INSPECTION November 2, 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	he inspection findings, no	violations were identified.						
2. Previous vi	olation(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) we	re discussed involving the	following red	quirement(s):				
4 During this	inspection certain of your	activities, as described belo	ow and/or a	ttached were in violati	on of NPC			
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								
			,					
		f,						
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	F	Printed Name		Signa	ture	Date		
LICENSEE'S REPRESENTATIVE								
NRC INSPECTOR	Deborah A. Pisk	ura		DARKU	ue	11/2/2010		
Branch Chief	Tamara E. Bloon	ner		Itsless	Juse	11/10/10		

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					
Medi-Physics, Inc.			U.S. Nuclear Regulatory Commission					
d/b/a GE Healthcare			Region III					
1623 Lotsie Blvd.			2443 Warrenville Road, Suite 210					
Overland, MO 63132			Lisle, IL 60532-4351					
REPORT NUMBER(S) 2010-001								
3. DOCKET NUMBER(S)		4. LICENSE NUMBER(S)		5. DATE(S) OF INSPECTION				
030-36453		24-32462-01MD		Nov. 2, 2010				
6. INSPECTION PROCEDURES 7		7. INSPECTION FOCUS AREAS						
87127		03.01 - 03.08						
SUPPLEMENTAL INSPECTION INFORMATION								
1.PROGRAM	2. PRIORITY	3. LICENSEE CONTACT		4. TELEPHONE NUMBER				
02500	2	Quent Besing, R.Ph., RSO		609-514-6647				
☐ Field Office Inspection Temporary Job Site Inspection								

PROGRAM SCOPE

This pharmacy employed four ANPs, two pharmacy technicians, and six drivers. The pharmacy served approximately 20 customers located in the St. Louis area and distributed approximately 300-350 doses daily. The licensee received three Mo99/Tc99^m generators each week. The pharmacy re-distributed I-131 therapy capsules and liquid; the pharmacy compounded capsules for I-131 diagnostic studies and hyperthyroid treatments. Beta-emitting materials were used by this pharmacy. The licensee's corporate office conducted annual audits of the pharmacy radiation safety program (last April 2010).

This inspection consisted of interviews with licensee personnel, a review of selected records, tour of the radiopharmacy, and independent measurements. During this inspection, the inspector observed early and mid-morning runs. These observations included dose calibrator QC/QA tests, generator elutions, drawing and assaying doses, thyroid bioassays, receiving packages, packaging doses for shipment, and conducting surveys for compliance with NRC and DOT requirements.

The maximum whole body and extremity exposures were reported (in millirem) as follows:

	<u>YID 2010</u>	2009
Whole body	115	125
Extremity	8,913	12,094