NRC FORM 591M PAR	Т 1			U.S NUCLEAR REGULATORY COMMISSION				
(06-2010) 10 CFR 2.201								
SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION								
1. LICENSEE/LOCATIO			2. NRC/REGIONAL OFFICE					
Midwest Divisio (d/b/a Research	U.S. Nuclear Regulatory Commission Region III							
17065 South 71	2443 Warrenville Road, Suite 210							
Belton, MO 640	Lisle, IL 60532-4351							
REPORT NUMBER(S)	2011-001	•						
3. DOCKET NUMBER(S) 4. LICENSEE NUM			R(S) 5. DATE(S) OF INSPECTION					
030-18606			March /6, 2011					
LICENSEE:			. 1811					
Regulatory Commission	(NRC) rules and regulation	s conducted under your lice	our license. Th	he inspection consist	ed of selective exami			
representative records, interviews with personnel, and observations by the inspector. The inspection findings are as follows:								
名"		violations were identified.						
2. Previous violation(s) closed.								
 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):								
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4. During this i	nspection certain of your	activities, as described belo	ow and/or atta	ached, were in violati	on of NRC			
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								
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		Statement of						
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	iture	Date		
LICENSEE'S REPRESENTATIVE								
NRC INSPECTOR	Deborah A. Pisk	ura		10APs Fu	1a)	3//6/2011		
Branch Chief	Tamara E. Bloor	mer		DARSKU IR DO	owe	3/25/11		

NRC FORM 591 M PART 3 (06-2010)

U.S. NUCLEAR REGULATORY COMMISSION

10 CFR 2.201

Docket File Information SAFETY INSPECTION REPORT AND COMPLIANCE INSPECTION

1. LICENSEE Midwest Division – RBH, LLC (d/b/a Research Belton Hospital) 17065 South 71 Highway Belton, MO 64012 REPORT NUMBER(S) 2011-001			2. NRC/REGIONAL OFFICE 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				
030-18606		24-24405-01		March 16, 2011				
6. INSPECTION PROCEDURES 87130 & 87131		7. INSPECTION FOCUS AREAS 03.01-03.08						
SUPPLEMENTAL INSPECTION INFORMATION								
1.PROGRAM	2. PRIORITY	3. LICENSEE CO	NTACT	4. TELEPHONE NUMBER				
02120	3	Barry A. Gut	oin, M.D., RSO	816-348-1224				

PROGRAM SCOPE

This licensee was a small community hospital and authorized to use licensed material permitted by Sections 35.100, 35.200, and 35.300. The nuclear medicine department was staffed with one full-time and two part-time technologists who performed approximately 100+ diagnostic procedures monthly which included a full spectrum of studies. The licensee received unit doses from a licensed radiopharmacy. The licensee administered no I-131 procedures requiring a written directive since the previous inspection. The hospital retained the services of a consulting physicist who audited the radiation safety program on a quarterly basis (last 1/18/2011).

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