RI - DNMS Licensee Event Report Disposition

Licensee:		Dept of Health & Human Services, Bethesda, MD								
Eve	nt Descri	iptio <u>n:</u>	Missing	Material	in a S	Source	Vial			
License No: 19 01			296-10	Docket No:		0300	77£6	MLER-RI:	2005-023	
Event Dat	te:	Ľ,	3/10/05	Report Date:		4/6/0	05	HQ Ops Event #	:	
1.	REPOF	RTING F	REQUIREMENT							
		10 CF	R 20.1906 Packs	age Contamina	tion		1	0 CFR 30.50 Re	port	
	XXX 10 CFR 20.2201 Theft or Loss					10 CFR 35.3045 Medical Event				
	10 CFR 20.2203 30 Day Report					License Condition				
		Other			<u> </u>					
2.	2. REGION I RESPONSE									
	Immediate Site Inspection					Inspector/Date				
	Special Inspection				Inspector/Date					
	Telephone Inquiry					Inspector/Date				
		Prelimi	nary Notification	/Report		- 1		Daily Report		
	X	Informa	ation Entered in	RI Log		•	X	Review at Next	Inspection	
i		Report	Referred To:							
3.	REPO	RT EVA	LUATION							
	X	Descri	otion of Event		$\left[\times \right]$	Correcti	ive Actic	ons		
	X	Levels	of RAM Involved	i	X	Calcula	tions Ad	lequate		
		Cause	of Event -	Com		Addition	nal Infon	mation Requeste	ed from Licensee	
4.	MANAGEMENT DIRECTIVE 8.3 EVALUATION									
	NA	Release w/Exposure > Limits Deliberate Misuse w/Exposure > Limits								
		Repeat	ted Inadequate (Control		Pkging	Failure	>10 rads/hr or C	ontamination>1000x Limits	
	Ш	Exposure 5x Limits Large# Indivs w/Exp>Limits or Medical Deterministic Effect							Medical Deterministic Effects	
		Potential Fatality Unique Circumstances or Safeguards Concerns								
		If any c	of the above are	involved:	\perp	1				
		Consid	ered Need for II	Γ		Consid	ered Ne	ed for AIT		
		Decisio	on/Made By/Date	:						
5.ੁ	MANA	GEME	NT DIRECTIVE 8	3.10 EVALUAT	ION (add	litional ev	valuatio	n for medical eve	ents only)	
	Timeliness - Inspection Meets Requirements (5 days for overdose / 10 days for underdose)									
	Medical Consultant Used-Name of Consultant/Date of Report:									
	Medical Consultant Determined Event Directly Contributed to Fatality									
	Device Failure with Possible Adverse Generic Implications									
	HQ or Contractor Support Required to Evaluate Consequences									
6.	SPECIAL INSTRUCTIONS OR COMMENTS									
	F	alla	w-we a	t nexi	to in	spech	<u>ه</u>			
□ Public			Inspector Signa	uture:	Long	y Co	n		Date: 5/25/05	
Non-Put	olic		Branch Chief In	itials:	\mathcal{L}	J		· · · <u> </u>	Date: 6/8/05	
· \		\Refere	nce\Blank Forms		RM.wpd				Rev. 02/01/05	

DEPARTMENT OF HEALTH & HUMAN SERVICES





National Institutes of Health Bethesda, Maryland 20892

APR - 7 2005

Administrator, Region I U.S. Nuclear Regulatory Commission 475 Allendale Road King of Prussia, PA 19406-1415

Re: License 19-00296-10

Dear Sir or Madam:

This written report is submitted in accordance with 10 CFR 20.2201(b). On 4/6/05, I made a telephone notification to the NRC Operations Center in accordance with 10 CFR 20.2201(a)(ii) regarding the incident described below.

On 3/10/05, we determined that 245 µCi (9.1 MBq) of ¹⁵³Gd (0.5N HCl) in a source vial was missing. The original activity was 1 mCi (37MBq) on 4/12/2004, and the missing quantity, corrected for radioactive decay, represented the remaining activity after several labeling experiments.

Following this discovery, the Authorized User interviewed all researchers who had access to her laboratory. None recalled disposing of the vial. In addition, a health physicist contacted a number of the researchers and performed an independent survey of this and adjacent laboratories with a photonsensitive survey meter. The original lead pig was found in a box where other lead pigs were stored awaiting disposal. However, the source vial was not in the lead pig.

In addition to searching the laboratory areas, we checked the records of both radioactive waste pick ups and other monitored waste from the building in which this laboratory is located. There was no indication of ¹⁵³Gd in any of the records or surveys.

At this time, the disposition of the material is unknown. Considering that the exposure rate for an unshielded point source of 245 µCi of ¹⁵³Gd is only 0.04 mR/hr, the missing material would not represent a significant external exposure risk. The approximately 69 nanograms of material missing would not represent a significant toxic hazard either.

Corrective actions have been implemented by suspending the Authorized User's ability to order more radioactive material. With her pending departure from NIH, we do not feel any further actions will be warranted in this situation.

Should you wish to contact me, my telephone number is 301-496-2254

Radiation Safety Office

cc: Dr. Michael Gottesman, Deputy Director for Intramural Research, NIH

Dr. Lance Liotta, Chairman, NIH Radiation Safety Committee

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