	Licensee: New Haven Heal	h Demrtment	
E	vent Description: Stoten Lead Ar	alvzer	
License		19999000/ MLER-RI: RI 201	4-012
Event D	1011110	12/17/13 HQ Ops Event #: 44640	
1.	REPORTING REQUIREMENT	<u></u>	
	10 CFR 20.1906 Package Contamination	10 CFR 30.50 Report	
	X 10 CFR 20.2201 Theft or Loss	10 CFR 35.3045 Medi	cal Event
	10 CFR 20.2203 30 Day Report	License Condition	
	Other		
2.	REGION I RESPONSE		
	Immediate Site Inspection	Inspector/Date	
	X Special Inspection		12014
	Telephone Inquiry	Inspector/Date	12011
	Preliminary Notification/Report	Daily Report	
	Information Entered in RI Log	Review at Next Inspection	
	Report Referred To:		
3.	REPORT EVALUATION		
0.		Compating Antione	
	Description of Event	Corrective Actions	
	Levels of RAM Involved	Calculations Adequate	
	MANAGEMENT DIRECTIVE 8.3 EVALUATION	Additional Information Requested from Licensee	,
\$.			
	Release w/Exposure > Limits	Deliberate Misuse w/Exposure > Limits	
	Repeated Inadequate Control	Pkging Failure>10 rads/hr or Contamination>10	
	Exposure 5x Limits	Large# Indivs w/Exp>Limits or Medical Determin	
	Potential Fatality	Unique Circumstances or Safeguards Concerns	
	If any of the above are involved:		
	Considered Need for IIT Decision/Made By/Date:	BLANK WELLINC 12/17/13	
			S .
5.	MANAGEMENT DIRECTIVE 8.10 EVALUATION (additional evaluation for medical events only)		
		s (5 days for overdose / 10 days for underdose)	
	Medical Consultant Used-Name of Consultant/Date of Report:		
	Medical Consultant Determined Event Direct		
	Device Failure with Possible Adverse Gene	ic Implications	
	HQ or Contractor Support Required to Evaluate Consequences		
5.	SPECIAL INSTRUCTIONS OR COMMENTS		1 0+
	I tem recovered report receiv	O DUNRC. INSpersion Nomplete. se	ut insolo Co

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Rev. 09/12/13