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L L L	U.S. NUCLEAR REGULATORY COMMISSION Amendment No. 66					
MATERIALS LICENSE						
Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 36, 39, 40, and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). This license shall be deemed to contain the conditions specified in Section 183 of the Atomic Energy Act of 1954, as amended, and is subject to all applicable rules, regulations, and orders of the Nuclear Regulatory Commission now or hereafter in effect and to any conditions specified below.						
Licensee		In accordance with	the letter dated			
		March 2, 2013,				
1. Waterbury Hospital Health Center		3. License number 06-02406-01 is amended in its entirety to read as follows:				
	- NR F	Ec.				
2. 64 Robbins Street	CLEAR F	4. Expiration date	December 31, 2015			
Waterbury, Connecticut 06721 🔨	S	5. Docket No. 030	-01251			
4		Reference No. (	06-30218-01			
S			2			
<ol> <li>Byproduct, source, and/or special nuclear material</li> </ol>	7. Chemical and/or	physical form 8	. Maximum amount that licensee may possess at any one time under this license			
<ul> <li>Any byproduct material permitted by 10 CFR 35.100</li> </ul>	A. Any		A. As needed			
<ul> <li>B. Any byproduct material permitted by 10 CFR 35.200</li> </ul>	B. Any	III Share Will E	3. As needed			
C. Any byproduct material permitted by 10 CFR 35.300	C. Any		C. 1 curie			
D. Any byproduct material permitted by 10 CFR 35.500	D. Sealed Source American Scie Model MED 36 Products Labo NES-8412)	ntific, Inc. 01; Isotope	). 2.5 curies			
E. Any byproduct material permitted by 10 CFR 31.11	E. Prepackaged k	Kits E	E. 10 millicuries			
9. Authorized use:						
<ul> <li>A. Any uptake, dilution and excretion study permitted by 10 CFR 35.100.</li> <li>B. Any imaging and localization study permitted by 10 CFR 35.200.</li> <li>C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300.</li> <li>D. Diagnostic medical use of sealed sources permitted by 10 CFR 35.500 in compatible devices registered pursuant to 10 CFR 30.32(g).</li> <li>E. <u>In vitro</u> studies.</li> </ul>						

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MATERIALS LICENSE SUPPLEMENTARY SHEET		Docket or Reference Number 030-01251;06-30218-01			
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	COND	ITIONS			
10.	<ol> <li>Licensed material may be used or stored only at the licensee's facilities located at 64 Robbins Street, Waterbury, Connecticut and 455 Chase Parkway, Waterbury, Connecticut.</li> </ol>				
11.	1. The Radiation Safety Officer for this license is Gerald J. Randall, M.S.				
12.	12. Licensed material is only authorized for use by, or under the supervision of:				
	A. Individuals permitted to work as an authorized user in accordance with 10 CFR 35.13 and 35.14.				
	B. The following individuals are authorized us	ers for medical use as indicated:			
	Authorized Users	Material and Use			
	Bernard Percarpio, M.D.	35.300			
	Erick A. Hyson, M.D.	35.100; 35.200; 35.500; Oral administration of sodium iodide iodine-131 for imaging and localization studies and treatment of hyperthyroidism and cardiac dysfunction; In vitro studies			
	Anthony R. Carter, M.D.	35.100; 35.200; 35.500; Oral administration of sodium iodide iodine-131 for imaging and localization studies and treatment of hyperthyroidism and cardiac dysfunction In vitro studies			
	Jeffrey A. Bitterman, M.D.	35.300 <u>In vitro studies</u>			
	Gerald R. Berg, M.D.	35.100; 35.200; 35.500 35.300, except thyroid carcinoma <u>In vitro studies</u>			
	Stephen Stein, M.D.	35.100; 35.200; 35.500; Oral administration of sodium iodide iodine-131 for imaging and localization studies and treatment of hyperthyroidism and cardiac dysfunction <u>In vitro studies</u>			

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	Authorized Users	Material and Use		
	Kenneth S. Allen, M.D.	35.100; 35.200; 35.500 In vitro studies		
	Anita L. Bourque, M.D.	35.100; 35.200; 35.500 In vitro studies		
	Duncan Belcher, M.D.	35.100; 35.200; 35.300		
	Andrew Lawson, M.D.	35.100; 35.200 35.300, except thyroid carcinoma		
	Marco Verga, M.D.	35.100; 35.200		
	Joseph Ravalese, III, M.D.	35.300		
	Kevin G. Kett, M.D.	35.100; 35.200		
	Mark L. Ruggiero, M.D.	35.200		
	Douglas Housman, M.D.	Oral administration of sodium iodide I-131		
13.	In addition to the possession limits in Item 8, the licensed material to quantities below the minimum decommissioning financial assurance.	licensee shall further restrict the possession of m limit specified in 10 CFR 30.35(d) for establishing		
14.	The licensee is authorized to transport licensed r 10 CFR Part 71, "Packaging and Transportation			

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15. Date	<ul> <li>accordance with the statements, representa including any enclosures, listed below. This that are required to be submitted in accorda condition does not limit the licensee's ability as provided for in 10 CFR 35.26. The U.S. govern unless the statements, representation and correspondence are more restrictive that</li> <li>A. Application dated July 28, 2005 [ML052]</li> <li>C. Letter dated August 25, 2005 [ML052]</li> <li>C. Letter dated September 28, 2005 [ML090]</li> <li>E. Letter dated March 12, 2009 [ML0907]</li> <li>F. Letter dated May 6, 2009 [ML0913208]</li> <li>G. Letter dated May 11, 2011 [ML111450]</li> </ul>	ations, and s license ince with v to make Nuclear I ons, and an the re 5216009 380456] 0529201 0430338 780240] 581] 0662]	condition applies only to those procedures the regulations. Additionally, this license changes to the radiation protection program Regulatory Commission's regulations shall procedures in the licensee's application gulations. 6]