NRC FORM 374

U.S. NUCLEAR REGULATORY COMMISSION

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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.

deliver or transfer such material to persons at shall be deemed to contain the conditions spapplicable rules, regulations and orders of the below.	ithorized pecified in	to receive it in a n Section 183 o	occordance with the re of the Atomic Energy	gulation Act of 1	ns of the applicable Part(s). This license 954, as amended, and is subject to al		
Licensee			In accordance with letter dated				
			October 17, 200	08,			
1, St. Joseph Health System LLC			3. License number 13-00418-02 is amended in its				
d/b/a St. Joseph Hospital	entirety to read as follows:						
2. 700 Broadway			4. Expiration date May 31, 2011				
Fort Wayne, IN 46802			5. Docket No. 030-01581				
			Reference No.				
Byproduct, source, and/or special nuclear material	7. Che	mical and/or ph	ysical form	ро	eximum amount that licensee may ssess at any one time under this ense		
Any byproduct material permitted by 10 CFR 35.100	A.	Any		Α.	As needed		
B. Any byproduct material permitted by 10 CFR 35.200	B.	Any		В.	As needed		
C. Any byproduct material permitted by 10 CFR 35.300	C.	Any		C.	As needed		
D. Any byproduct material permitted by 10 CFR 35.500	D.	Sealed soul	rce (Gammatron, OS-213A)	D.	One source not to exceed 300 millicuries		

9. Authorized Use:

- A. Any uptake, dilution and excretion study permitted by 10CFR 35.100.
- B. Any imaging and localization study permitted by 10CFR 35.200.
- C. Any diagnostic study or therapy procedure permitted by 10CFR 35.300.
- D. For storage only incident to disposal.

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CONDITIONS

- 10. Licensed material may be used or stored only at the licensees facilities located at 700 Broadway, Fort Wayne, Indiana.
- 11. The Radiation Safety Officer for this license is Dakshesh Patel, M.D.
- 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 10CFR 35.13 and 35.14.
 - B. The following individuals are authorized users for medical use as indicated:

Authorized Users	Material and Use
Brett Hagedorn M.D.	10 CFR 35.100, 35.200.
John Pasalich, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.
Stephen R. Phillip, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.
Marc Thomas, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.
David Sorg, M.D.	Oral administration of sodium iodide-131 in quantities equal to or less than 33 millicuries.
Diane D. Daly, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.
Frederick N. Vandeman, M.D.	10 CFR 35.100, 35.200, 35.300 (Oral administration of sodium iodide-131 in quantities equal to or less than 33 millicuries) and 35.500.
Christopher Kowalski, M.D	10 CFR 35.100, 35.200.
Joseph R. Decamp, M.D.	10 CFR 35.100, 35.200, 35.500 and oral administration of sodium iodide-131 in quantities equal to or less than 33 millicuries.
John L. Bormann, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.
Michael E. Parker, M.D.	10 CFR 35.100, 35.200, 35.300 and 35.500.

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Authorized	<u>Users</u>	Material and U	<u>se</u>					
Andre B. Stoval	I, M.D.	10 CFR 35.100, 35	5.200.					
Pamela Lee Stra	ange, M.D.	10 CFR 35.100, 35	5.200, 35.300 and 35.	500.				
John Rock, M.D).	10 CFR 35.100 and	d 35.200.					
Richard P. Step	hens, M .D.	10 CFR 35.100, 35.200, 35.300 and 35.500.						
James C. Wehre	enberg, M.D.	10 CFR 35.100 an	d 35.200.					
James A. Arata,	M.D.	10 CFR 35.100, 35.200, 35.300, and 35.500.						
David B. Janizel	k, M.D.	10 CFR 35.100, 35.200, 35.300, and 35.500.						
Christine Anne	Tremper, M.D.	10 CFR 35.100 an	d 35.200.					
Randall J. Phillip	os, M.D.	10 CFR 35.100, 35	5.200, 35.300 and 35.	500.				
John R. Kim, M.	D.		5.200, 35.300 (oral adr tities equal to or less t					
Richard Sibley, I	M.D.	10 CFR 35.100 and	d 35.200.					
Dakshesh Patel	M.D.	10 CFR 35.100 and	d 35.200.					
Shilpa Kashyap,	M.D.	10 CFR 35.100 and	d 35.200.					
H. Steven Beyer	r, M.D.	10 CFR 35.300 ().						
Eric V. Heatwole	e, M.D.	10 CFR 35.100 and	d 35.200.					
Shawn Johnson	, M.D.	10 CFR 35.100, 35	5.200, 35.300.					
Brian Kim, M.D.		10 CFR 35.100 and	d 35.200					
Sandeep S. Ahlu	uwalia, M.D.	10 CFR 35.100, 35 administration of so	.200 and 35.300 (limit odium iodide-131)	ed to th	e ora	al		
John C. Lacunza	a, M.D.	10 CFR 35.100 and	d 35.200					
Linda Gould Hi	ppenhammer, M.D.	10 CFR 35.100, 35	.200 and 35.300.					
Steven Hossler	, M .D.	10 CFR 35.100 and	d 35.200					

- 13. For iodine-131 therapeutic procedures that do not meet the criteria of 10 CFR 35.75, the licensee shall implement the procedures in Appendix P, "Model Procedure for Radiation SafetyDuring Iodine Therapy", found in Regulatory Guide 10.8, Revision 2, dated August 1987.
- 14. In addition to the possession limits in Item 8, the licensee shall further restrict the possession of licensed material to quantities below the minimum limit specified in 10 CFR 30.35(d) for establishing decommissioning financial assurance.
- 15. The licensee is authorized to transport licensed material in accordance with the provisions of 10 CFR Part 71, "Packaging and Transportation of Radioactive Material."
- 16. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below. This license condition applies only to those procedures that are required to be submitted in accordance with the regulations. Additionally, this license condition does not limit the licensee's ability to make changes to the radiation protection program as provided for in 10 CFR35.26. The U.S. Nuclear Regulatory Commission's regulations shall govern unless the statements, representations, and procedures in the licensee's application and correspondence are more restrictive than the regulations.
 - A. Application dated December 26, 2000.
 - B. Letter dated October 17, 2008.

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

DEC 2 9 2008

Date

James R. Mullauer, M.H.S.

Materials Licensing Branch

Region III