NRC FORM 374

U.S. NUCLEAR REGULATORY COMMISSION

Amendment No. 33

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. 0C02120 317646

Licensee	
1. Riverview Hospital	In accordance with the letter dated **October 28, 2008,** 3. License number 13-16286-01 is amended in its entirety to read as follows:
2. 395 Westfield Road P.O. Box 220 Noblesville, IN 46060	4. Expiration date October 31, 2010 5. Docket No. 030-10729 Reference No.

6. Byproduct, source, and/or special nuclear material	7. Chemical and/or physical form	8. Maximum amount that licensee may possess at any one time under this license
A. Any byproduct material permitted by 10 CFR 35.100	A. Any	A. As needed
B. Any byproduct material permitted by 10 CFR 35.200	B. Any	B. As needed
C. Any byproduct material permitted by 10 CFR 35.300	C. Any	C. As needed (not to exceed one curie of I-131)
D. Any byproduct material permitted by 10 CFR 31.11	D. Prepackaged Kits	D. As needed

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. In vitro studies.

CONDITIONS

10. Licensed material shall be used only at the licensee's facilities located at 395 Westfield Road, Noblesville, Indiana.

11. The Radiation Safety Officer for this license is Phillip R. Partlan, R.T.

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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
Anthony J. Cossell, M.D.	10 CFR 35.100, 35.200, and 31.11.
William E. McGraw, M.D.	10 CFR 35.100 and 35.200.
John Mark Michael, M.D.	10 CFR 35.100, 35.200 and 35.300.
Richard L. Hallett, M.D.	10 CFR 35.100, 35.200 and 35.300.
Peter D. Arfken, M.D.	10 CFR 35.100, 35.200 and 35.300.
John A. Morton, M.D.	10 CFR 35.100, 35.200, 35.300 and 31.11.
Margaret Brengle, M.D.	10 CFR 35.100 and 35.200.
Jeffrey I. Reider, M.D.	10 CFR 35.100 and 35.200.
Homer F. Beltz, M.D.	10 CFR 35.100 and 35.200.
Michael S. Skulski, M.D.	10 CFR 35.100 and 35.200.
Richard T. Beeler, M.D.	10 CFR 35.100, 35.200 and 35.300.
Steven A. Fritsch, M.D.	10 CFR 35.100 and 35.200.
Frank J. Pistoia, M.D.	10 CFR 35.100 and 35.200.
Michael A. Kuharik, M.D.	10 CFR 35.100 and 35.200
Janalyn P. Ferguson, M.D.	10 CFR 35.100 and 35.200.
Jane S. Mitchell, M.D.	10 CFR 35.100 and 35.200.
Jack J. Moss, M.D.	10 CFR 35.100 and 35.200.
Lori J. Wells, M.D.	10 CFR 35.100 and 35.200.
Eric D. Elliott, M.D.	10 CFR 35.100 and 35.200.
Charles A. Lerner, M.D.	10 CFR 35.100 and 35.200.

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Thomas Hagman, M.D.	10 CFR 35.100, 35.200 and 35.300.
Brian J. Weigel, M.D.	10 CFR 35.100, 35.200 and 35.300.
Jack M. Drew, M.D.	10 CFR 35.100, 35.200 and 35.300.
David R. Gulliver, M.D.	10 CFR 35.100, 35.200 and 35.300.
J. Michael Phelps, Jr., M.D.	10 CFR 35.100, 35.200, and 35.300 (Oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).
Timothy L. Davis, M.D.	10 CFR 35.100, 35.200, and 35.300 (Oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).
Vincent P. Mathews, M.D.	10 CFR 35.100 and 35.200.
Theodore P. Labus, M.D.	10 CFR 35.100, 35.200 and 35.300.
Cristina N. Shinaver, M.D.	10 CFR 35.100, 35.200 and 35.300.
Edward R. Bartley, M.D.	10 CFR 35.100 and 35.200.
Caryn Cockerill Anderson, M.D.	10 CFR 35.100 and 35.200.
Warren Kent Hansen, M.D.	10 CFR 35.100, 35.200, and 35.300 (Oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).
Carlo Robert Lazzaro, M.D.	10 CFR 35.100, 35.200, and 35.300 (Oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).
Charles C. Mulry, M.D.	10 CFR 35.100, 35.200, and 35.300 (Oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).
Eric E. Beltz, M.D.	**10 CFR 35.100, 35.200, and 35.300 (Limited to the oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries).**

13. 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 10CFR 30.35(d) for establishing decommissioning financial assurance.

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14. The licensee is authorized to transport licensed material in accordance with the provisions of 10 CFR Part 71, "Packaging and Transportation of Radioactive Material."

15. 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 CFR 35.26. 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 September 21, 2000 (except Item 12.1); and

    B. Facsimiles dated November 20, 2000, July 26, 2007, July 7, 2008, and July 14, 2008, and;

    C. Letters dated August 2, 2001, August 23, 2001, November 6, 2001, January 14, 2002 (with enclosed close-out survey of old Nuclear Cardiology room), March 26, 2002 (with enclosed closeout survey of old nuclear medicine department), March 28, 2002, June 9, 2006 and June 19, 2007; and

    D. Facsimiles letter dated September 7, 2006, **December 8, 2008**.

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date DEC 16 2008

By James Mullauer

James R. Mullauer, M.H.S.
Materials Licensing Branch
Region III