U.S. NUCLEAR REGULATORY COMMISSION

Amendment No. 24

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, 37, 39, 40, 70 and 71, 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 letter dated October 08, 2021,	4. Expiration Date: April 30, 2022
1. Hancock Regional Hospital		
2. P.O. Box 827 801 N. State St. Greenfield, IN 46140	3. License No.: 13-16730-01 is amended in its entirety to read as follows:	5. Docket No.: 030-11551 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	9. Authorized use
A. Any byproduct material permitted by 10 CFR 35.100	A. Any	A. As Needed	A. For use in uptake, dilution and excretion studies permitted by 10 CFR 35.100.
B. Any byproduct material permitted by 10 CFR 35.200	B. Any	B. As Needed	B. For use in imaging and localization studies permitted by 10 CFR 35.200.
C. Any byproduct material permitted by 10 CFR 35.300	C. Any	C. 1 curie total	C. For any use permitted by 10 CFR 35.300.

MATERIALS LICENSE SUPPLEMENTARY SHEET	License No.: 13-16730-01	Docket or Reference No.: 030-11551
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CONDITIONS

10. A. Licensed material shall be used or stored at the licensee's facilities located at 801 N. State St., Greenfield, Indiana 46140.

    B. Licensed material listed in Subitem Nos. 6.A. and 6.B. shall be used at the licensee's facilities located at Hancock Surgical Center, 1 Memorial Sq., Greenfield, Indiana 46140.

11. The Radiation Safety Officer (RSO) for this license is Justin P. Chang, M.D.

12. Licensed material shall only be used by, or under the supervision of:

    A. Individuals permitted to work as authorized users in accordance with 10 CFR 35.13 and 10 CFR 35.14.

    B. The following individuals are authorized users for the material and medical uses as indicated:

Authorized User (M.D.,D.O.,etc.)	Material and Use
Mark Allen, M.D.	10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131)
James Blahunka, M.D.	10 CFR 35.100,10 CFR 35.200,10 CFR 35.300
James R. Bognanno, M.D.	10 CFR 35.100,10 CFR 35.200
Kevin E. Burton, M.D.	10 CFR 35.100,10 CFR 35.200
Justin P. Chang, M.D.	10 CFR 35.100, 10 CFR 35.200, and 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131 in quantities less than or equal to 33 millicuries)
Julia J. Compton, M.D.	10 CFR 35.300 (limited to the parenteral administration of radium Ra-223 dichloride)
Thomas C. Dugan, M.D.	10 CFR 35.300
Stanley S. Givens, M.D.	10 CFR 35.300
Stefan Andrew Hoff, M.D.	10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131)

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Authorized User (M.D.,D.O.,etc.)	Material and Use
Tarun R. Jindal, M.D.	10 CFR 35.100, 10 CFR 35.200, and 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131)
Kenyon K. Kopecky, M.D.	10 CFR 35.100,10 CFR 35.200,10 CFR 35.300
David Kurlander, M.D.	10 CFR 35.100,10 CFR 35.200,10 CFR 35.300
Christopher A. Leagre, M.D.	10 CFR 35.300
Robert Liebross, M.D.	10 CFR 35.300
Michael L. Lutz, M.D.	10 CFR 35.100,10 CFR 35.200,10 CFR 35.300
Gregory A. Merchun, M.D.	10 CFR 35.100,10 CFR 35.200,10 CFR 35.300
Thomas N. Murphy, M.D.	10 CFR 35.100,10 CFR 35.200
Dennis Lee Myers, M.D.	10 CFR 35.100,10 CFR 35.200
Mark J. Paluszny, M.D.	10 CFR 35.100,10 CFR 35.200
Frank W. Peyton, M.D.	10 CFR 35.300 (limited to the parenteral administration of radium Ra-223 dichloride)
Marc J. Pinchouck, M.D.	10 CFR 35.100,10 CFR 35.200
Paul W. Sheets, M.D.	10 CFR 35.100,10 CFR 35.200
Mark Allan Sparrow, M.D.	10 CFR 35.300 (limited to the oral administration of sodium iodide I-131)
Stephan M. Stockberger, Jr., M.D.	10 CFR 35.100, 10 CFR 35.200, and 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131)
Michael L. Swack, M.D.	10 CFR 35.100,10 CFR 35.200
Perry E. Wethington, M.D.	10 CFR 35.100,10 CFR 35.200

13. 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.

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A. Application dated October 24, 2011 (ML113081683)
B. Letter dated August 28, 2013 (ML13254A403)
C. Letter dated October 8, 2021 (ML21309A061)
D. Letter dated January 7, 2022 (ML22011A097)

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date: January 11, 2022

By: Frank P. Tran Digitally signed by Frank P. Tran Date: 2022.01.11 15:04:15 -06'00'

Frank P. D. Tran
Region 3