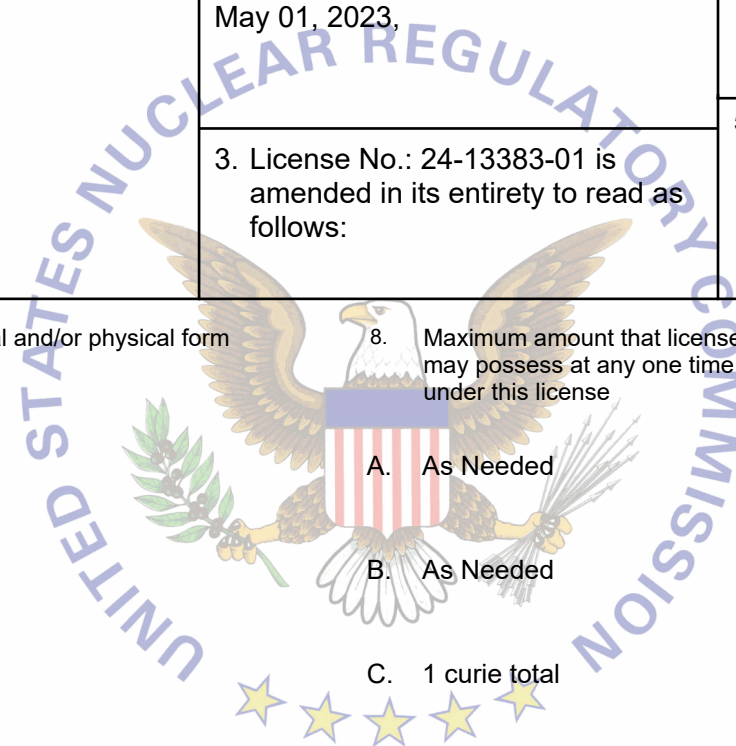


**MATERIALS LICENSE**

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.

<p style="text-align: center;">Licensee</p> <p>1. Christian Hospital Dept. of Nuclear Medicine</p> <p>2. 11133 Dunn Rd. St. Louis, MO 63136</p>		<p>In accordance with letter dated May 01, 2023,</p>	<p>4. Expiration Date: January 31, 2037</p>
		<p>3. License No.: 24-13383-01 is amended in its entirety to read as follows:</p>	<p>5. Docket No.: 030-02382 Reference No.:</p>
<p>6. Byproduct, source, and/or special nuclear material</p> <p>A. Any byproduct material permitted by 10 CFR 35.100</p> <p>B. Any byproduct material permitted by 10 CFR 35.200</p> <p>C. Any byproduct material permitted by 10 CFR 35.300</p> <p>D. Any byproduct material permitted by 10 CFR 31.11</p>	<p>7. Chemical and/or physical form</p> <p>A. Any</p> <p>B. Any</p> <p>C. Any</p> <p>D. Prepackaged Kits</p>	<p>8. Maximum amount that licensee may possess at any one time under this license</p> <p>A. As Needed</p> <p>B. As Needed</p> <p>C. 1 curie total</p> <p>D. 5 millicuries total</p>	<p>9. Authorized use</p> <p>A. For use in uptake, dilution and excretion studies permitted by 10 CFR 35.100.</p> <p>B. For use in imaging and localization studies permitted by 10 CFR 35.200.</p> <p>C. For any use permitted by 10 CFR 35.300.</p> <p>D. For use in in-vitro studies.</p>



**MATERIALS LICENSE  
SUPPLEMENTARY SHEET**

License No.: 24-13383-01

Docket or Reference No.:  
030-02382

Amendment No. 79

CONDITIONS

10. Licensed material may be used or stored at the licensee's facilities located at 11133 Dunn Rd., St. Louis, Missouri 63136.
11. The Radiation Safety Officer (RSO) for this license is Scott J. Surovi.
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 medical uses as indicated:

Authorized User (M.D.,D.O.,etc.)

Material and Use

Shahed N. Badiyan, M.D.

10 CFR 35.300

Farrokh Dehdashti, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Tyler J. Fraum, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Saul N. Friedman, M.D., Ph.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Robert J. Gropler, M.D.

10 CFR 35.100,10 CFR 35.200

Paul W. Hargan, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Malak Itani, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

David A. Leitman, D.O.

10 CFR 35.100,10 CFR 35.200; 10 CFR 35.300 (limited to oral administration of sodium iodide I-131)

Joyce C. Mhlanga, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Jeff M. Michalski, M.D.

10 CFR 35.300

Smita S. Parikh, M.D.

10 CFR 35.100,10 CFR 35.200

Maria Rosana Ponisio, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

**MATERIALS LICENSE  
SUPPLEMENTARY SHEET**

License No.: 24-13383-01

Docket or Reference No.:  
030-02382

Amendment No. 79

Authorized User (M.D., D.O., etc.)

Henry D. Royal, M.D.

Thomas H. Schindler, M.D.

Barry A. Siegel, M.D.

James Galen Stewart, M.D.

Amir Iravani Tabrizipour, M.D.

Wade L. Thorstad, M.D.

Wenzel Vas, M.D.

Richard L. Wahl, M.D.

Jerold W. Wallis, M.D.

Material and Use

10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300

10 CFR 35.100, 10 CFR 35.200

10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300

10 CFR 35.100, 10 CFR 35.200

10 CFR 35.100, 10 CFR 35.200; 10 CFR 35.300 (limited to oral administration of sodium iodide I-131)

10 CFR 35.300

10 CFR 31.11, 10 CFR 35.100, 10 CFR 35.200

10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300

10 CFR 35.100, 10 CFR 35.200, 10 CFR 35.300



**MATERIALS LICENSE  
SUPPLEMENTARY SHEET**

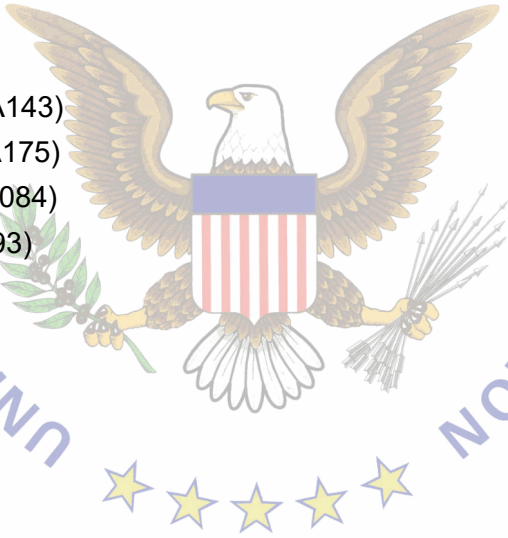
License No.: 24-13383-01

Docket or Reference No.:  
030-02382

Amendment No. 79

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 statements, representations, and 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 impose on the licensee requirements that are more restrictive than or in addition to the regulations.

- A. Application dated July 21, 2021 (ML21214A143)
- B. Letter dated February 26, 2021 (ML21063A175)
- C. Letter dated February 11, 2022 (ML22074A084)
- D. Letter dated August 18, 2022 (ML22256A193)
- E. Letter dated May 1, 2023 (ML23145A143)



FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date: June 26, 2023By: \_\_\_\_\_  
Magdalena R. Gryglak  
Region 3