

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 1. SSM Audrain Health Care, Inc. d/b/a SSM Health St. Mary's Hospital - Audrain 2. 620 E Monroe Street Mexico, MO 65265		In accordance with letter dated March 05, 2020.	4. Expiration Date: August 31, 2025
			5. Docket No.: 030-08599 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. 500 millicuries total	C. For any use permitted by 10 CFR 35.300.
D. Any byproduct material permitted by 10 CFR 31.11	D. Prepackaged Kits	D. 100 millicuries total	D. For use in in-vitro studies.

**MATERIALS LICENSE
SUPPLEMENTARY SHEET**

License Number

24-15122-01

Docket or Reference Number

030-08599

Amendment No. 30

CONDITIONS

10. Licensed material may be used or stored at the licensee's facilities located at 620 E Monroe Street, Mexico, Missouri, 65265.
11. The Radiation Safety Officer (RSO) for this license is Kenneth Wohlt, M.S.
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

Karen J. Baranski, M.D.

10 CFR 35.100,10 CFR 35.200

George Cyriac, M.D.

10 CFR 31.11,10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Lance Faler, M.D.

10 CFR 35.100,10 CFR 35.200

Robert M. Fischer, M.D.

10 CFR 35.100,10 CFR 35.200

Megan Maine Gau, M.D.

10 CFR 35.100,10 CFR 35.200

Robert Gresick, M.D.

10 CFR 35.100,10 CFR 35.200

Larry Handlin, D.O.

10 CFR 35.100,10 CFR 35.200

Allen Ernest Hillard, M.D.

10 CFR 31.11,10 CFR 35.100,10 CFR 35.200; 10 CFR 35.300 (limited to the oral administration of sodium iodide I-131 in quantities less than or equal to 33 mCi)

Neal Richard Meyer, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Mark Haskell Monroe, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Rebecca J. Mueller, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

Ronald Palmer, M.D.

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300

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

Thomas M. Schroyer, M.D.

John Joseph Stephens, M.D.

Andre Strzembosz, M.D.

Material and Use

10 CFR 35.100,10 CFR 35.200

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

10 CFR 35.100,10 CFR 35.200,10 CFR 35.300



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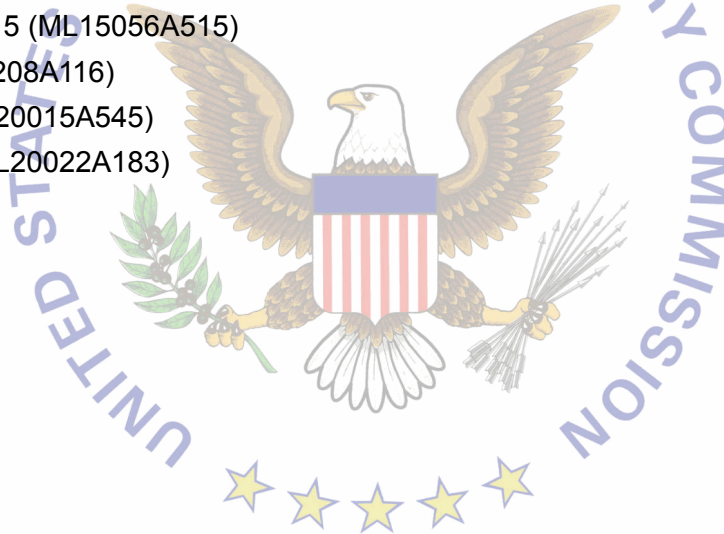
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13. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below. This license 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.

- A. Application dated January 22, 2015 (ML15056A515)
- B. Letter dated July 24, 2015 (ML15208A116)
- C. Letter dated January 8, 2020 (ML20015A545)
- D. Letter dated January 22, 2020 (ML20022A183)



FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date: April 20, 2020

By: _____

Laura B. Cender
Region 3