

**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.

Licensee

1. Poplar Bluff Regional Medical Center  
d/b/a Three Rivers Health Care, North
2. 2620 N. Westwood Boulevard  
Poplar Bluff, MO 63901

In accordance with two letters dated  
**October 14, 2013,**

3. License number 24-16652-01 is amended in its  
entirety to read as follows:

4. Expiration date March 31, 2022

5. Docket No. 030-11417  
Reference No.

6. Byproduct, source, and/or special  
nuclear material

- A. Any byproduct material  
permitted by 10 CFR 35.100
- B. Any byproduct material  
permitted by 10 CFR 35.200
- C. Any byproduct material  
permitted by 10 CFR 35.300
- D. Cesium-137 permitted by 10  
CFR 35.400
- E. Iodine-125 permitted by 10  
CFR 35.400
- F. Palladium-103 permitted by  
10 CFR 35.400
- G. Any byproduct material  
permitted by 10 CFR 31.11

7. Chemical and/or physical form

- A. Any
- B. Any
- C. Any
- D. Sealed sources (AEA  
Technology Model CDC.T1)
- E. Sealed sources  
(Prostaseed Model I125SL;  
Bard Brachytherapy, Inc.,  
Model STM 1251; and  
Theragenics Corporation  
Model I-Seed AgX100)
- F. Sealed sources (North  
American Scientific Model  
IAPd-103A; and  
Theragenics Corporation  
Model Theraseed 200)
- G. Prepackaged kits

8. Maximum amount that licensee may  
possess at any one time under this  
license

- A. As needed
- B. As needed
- C. 800 millicuries
- D. 1000 millicuries
- E. 500 millicuries
- F. 500 millicuries
- G. 1 millicurie

9. Authorized use:

- A. Any uptake, dilution and excretion study permitted by 10 CFR 35.100.
- B. Any imaging and localization study permitted by 10 CFR 35.200.

**MATERIALS LICENSE  
SUPPLEMENTARY SHEET**

License Number

24-16652-01

Docket or Reference Number

030-11417

Amendment No. 42

- C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300.
- D. through F. Any manual brachytherapy procedure permitted by 10 CFR 35.400.
- G. In vitro studies.

**CONDITIONS**

10. A. Licensed material described in Subitem Nos. 6.A. through 6.G. may be used at the licensee's facilities located at 2620 N. Westwood Boulevard, Poplar Bluff, Missouri.
- B. Licensed material described in Subitem Nos. 6.A. through 6.C. and 6.E. through 6.G. may also be used at the licensee's facilities located at 3100 Oak Grove Road, Poplar Bluff, Missouri.
- C. **Licensed material described in Subitem Nos. 6.A. through 6.B. may also be used at the licensee's facilities located at 3098 Oak Grove Road, Poplar Bluff, Missouri.**
11. Radiation Safety Officer: Jim Smith, M.S.
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 10 CFR 35.13 and 35.14.
- B. The following individuals are authorized users for medical use as indicated:

**Authorized Users**

Subhash B. Gujarati, M.D.

Rubina Mirza, M.D.

Dean K. Rigby, M.D.

Rajinder Moham Gulati, M.D.

Tom Buford Brumitt, D.O.

Emily Militzer, M.D.

Girish Bhatt, M.D.

**Material and Use**

10 CFR 35.300 limited to strontium-89 and samarium-153 and 35.400.

10 CFR 35.100 and 35.200.

10 CFR 35.100 and 35.200.

10 CFR 35.100, 35.200 and 35.300.

10 CFR 35.100, 35.200 and 35.300 (excluding oral administration of sodium iodide I-131 in quantities greater than 33 millicuries).

10 CFR 35.400.

10 CFR 35.200.

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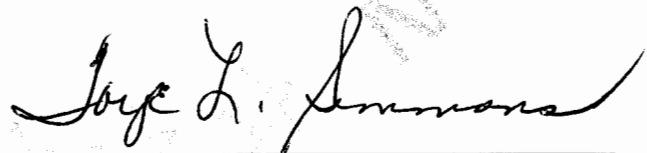
Amendment No. 42

13. The licensee is authorized to transport licensed material in accordance with the provisions of 10 CFR Part 71, "Packaging and Transportation of Radioactive Material."
14. 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 29, 2011; and
- B. Letters dated March 15, 2012, March 16, 2012 (two separate letters), August 27, 2012, October 29, 2012, April 15, 2013, and **October 14, 2013 (two separate letters)**.

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date JAN 21 2014

By

Toye L. Simmons  
Materials Licensing Branch  
Region III