

**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. Crittenton Hospital Medical Center</p> <p>2. 1101 W. University Drive Rochester, MI 48307</p>	<p>In accordance with letter dated <b>May 20, 2016</b>,</p> <p>3. License number 21-13562-01 is amended in its entirety to read as follows:</p> <hr/> <p>4. Expiration date August 31, 2021</p> <hr/> <p>5. Docket No. 030-02157 Reference No.</p>
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<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. Iridium-192 permitted by 10 CFR 35.400</p> <p>E. Iodine-125 permitted by 10 CFR 35.400</p> <p>F. Palladium-103 permitted by 10 CFR 35.400</p> <p>G. Cesium-131 permitted by 10 CFR 35.400</p> <p>H. Gadolinium-153 permitted by 10 CFR 35.500</p>	<p>7. Chemical and/or physical form</p> <p>A. Any</p> <p>B. Any</p> <p>C. Any</p> <p>D. Sealed sources (Best Medical International, Inc. Model 81-01 Series)</p> <p>E. Sealed sources (Medi-Physics, Inc. Model 6711 OncoSeed and Model 9011; Bard Brachytherapy, Inc. Model STM 1251, North American Scientific, Inc., Model MED3633, and IsoAid Advantage I-125 (IAI-125A))</p> <p>F. Sealed sources (North American Scientific, Inc., Model MED3633)</p> <p>G. Sealed sources (IsoRay Model CS-1)</p> <p>H. Sealed sources (Isotope Products Labs Models NES-8426 and AEA Technology Model GD.LIN2)</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</p> <p>D. 300 millicuries</p> <p>E. 400 millicuries</p> <p>F. 300 millicuries</p> <p>G. 400 millicuries</p> <p>H. 32 sources, not to exceed 14 sources per tray, not to exceed 120 millicuries per source tray, and 320 millicuries total</p>
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9. Authorized use:
- A. Any uptake, dilution and excretion study permitted by 10 CFR 35.100.

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- B. Any imaging and localization study permitted by 10 CFR 35.200.
- C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300.
- D. through G. Any manual brachytherapy procedure permitted by 10 CFR 35.400.
- H. Diagnostic medical use of sealed sources permitted by 10 CFR 35.500 in compatible devices registered pursuant to 10 CFR 30.32(g).

CONDITIONS

10. Licensed material may be used or stored only at the licensee's facilities located at 1101 W. University Drive, Rochester, Michigan.
11. The Radiation Safety Officer (RSO) for this license is Annie Kalapparambath, M.D.
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

Kanak Varde, M.D.

Kenneth Levin, M.D.

Judith M. Bender, M.D.

Carla Cook, M.D.

Annie Kalapparambath, M.D.

Amf Aref, M.D.

Anna Norris Rabbani, M.D.

Paul Chuba, M.D.

Material and Use

10 CFR 35.100, 35.200, 35.300 (limited to the oral administration of sodium iodide-131 in quantities less than or equal to 33 millicuries), and 35.500.

10 CFR 35.300 and 35.400.

10 CFR 35.100, 35.200, 35.300, and 35.500.

10 CFR 35.400.

10 CFR 35.100, 35.200, 35.300 (limited to the oral administration of sodium iodide 131), and 35.500.

10 CFR 35.400.

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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 December 29, 2010 (excluding request to permit a medical physicist under the supervision of an Authorized Medical Physicist (AMP) to be physically present during HDR treatments instead of the AMP) (ML110030161)
  - B. Letter dated November 30, 2011 (ML113400517)
  - C. Letter dated April 3, 2012 (ML12095A154)
  - D. Letter dated April 23, 2014 (ML14115A303)
  - E. Letter dated April 24, 2015 (ML15117A583)
  - F. Letter dated July 9, 2015 (ML15205A257)
  - G. Letter dated July 16, 2015 (ML15205A259)
  - H. Letter dated, and August 27, 2015 (ML15240A392)

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date JUL 07 2016By Sara A. Forster  
Sara A. Forster, M.S.  
Materials Licensing Branch  
Region III