

July 28th, 2011

NRC License # 48-32697-01

Material Licensing Branch U.S. Nuclear Regulatory Commission, Region III 2443 Warrenville Road, Suite 210 Lisle, IL 60532-4352

Northern Shared Medical Services would like to add additional authorized users to our radioactive materials license. The authorized users are as follows:

Authorized Users:

John Baird, M.D.	NRC License# 24-01565-01
Barbara Tellerman, M.D.	NRC License# 24- 01565-01
Charles M. Swaney, M.D.	NRC License# 24-01565-01
Terry J. Elwing, M.D.	NRC License# 24-01565-01
Laura J. Sievert, M.D.	NRC License# 24-01565-01
Maxwell Lazinger, M.D.	NRC License# 24-01565-01
David Perry Brummett, M.D.	NRC License# 24-01565-01
Chad Michael Ruble, M.D.	NRC License# 24-01565-01

Thank you for your kind attention. If any questions should arise, please feel free to contact me 608-839-9956.

Sincerely

Paul Flaten, Director of Health & Safety, ARSO Northern Shared Medical Services

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NRC	FORM	374
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U.S. NUCLEAR REGULATORY COMMISSION

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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 r shall be deemed to contain the conditions specified in S	o use such material for the purpose(s) and at the place(s) designated below; to eccive it in accordance with the regulations of the applicable Part(s). This license ection 183 of the Atomic Energy Act of 1954, as amended, and is subject to all egulatory Commission now or hereafter in effect and to any conditions specified
Licensee	In accordance with letters dated
BEICH 1966	September 23, 2009, and November 17, 2009,
Boone Hospital Center	3. License number 24-01565-01 is amended in its
	entirety to read as follows:
2. 1600 East Broadway	4. Expiration date April 30, 2015
Columbia, MO 65201	5. Docket No. 030-02304
	Reference No.
A. Any byproduct material A. Any byproduct material B. Any byproduct material Permitted by 10 CFR 35 200 Cf. Any byproduct material Permitted by 10 CFR 35 300 D. Any byproduct material Permitted by 10 CFR 35 300 D. Any byproduct material Permitted by 10 CFR 35 400 E. Cesium-137 permitted by 10 CFR 35.400 F. Cesium-137 F. S M G. Any byproduct material A. A	8. Maximum amount that licensee may possess at any one time under this license A. As needed B. As needed
permitted by 10 CFR 31.11	Activities to the second secon
H. Depleted uranium I. M	letal I. 999 kilograms
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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 GFR 35 200.
- C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300.
- D. Any manual brachytherapy procedure permitted by 10 CFR 35.400.
- E. and F. For storage only incident to disposal.
- G. In vitro studies.
- H. For use as shielding material.

CONDITIONS

- 10. Licensed material may be used at the licensee's facilities located at Boone Hospital Center, 1600 East Broadway, Columbia, Missouri
- 11. Radiation Safety Officer for this license is Liesje Myers, CNMT.
- 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	Material and Use
V		10.CPR:85,100 35.200 iodine;131, diagnostic procedures
^	John Baird, M.D.	permitted by 35,300 and 31,11
X	Barbara Tellerman, M.D.	10 OFR85 100, 35.200, iodine 131 diagnostic procedures
		permitted by 35 300 and 31.11.
X	Charles M. Swaney, M.D.	10 CFR 35.100, 35.200, 31.11, and iodine-131 for diagnosti

procedures and the treatment of hyperthyroidism permitted by 35,300.

Mark Bryer, M.D. 10 CFR 35.300 and 35.400.

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Ste	ven Westgate, M.D.	10 CFR 35.300 an	d 35.400.
Jos	eph M. Bean, M.D.	10 CFR 35.300 an	d 35.400.
X Ten	ry J. Elwing, M.D.	10 CFR 35 100, 30 permitted by 35.30	200, Iodine-131 diagnostic procedures 0 and 31.11.
X Lau	ra J. Sieven, M.D.		5.200, iodine-131 diagnostic procedures and for thyroidism permitted by 35.300 and 31.11.
Jam	nes Allen, M.D.	10 CFR 35.300 an	d 35.400.
X Max	well Lazinger, M.D.	10 CFR 35.100, 35 permitted by 35.30	5.200, iodine-131 diagnostic procedures 0 and 31.11.
X pDav	id Perry Brummett, M.D.		5.200, iodine-131 for diagnostic procedures and perthyroidism permitted by 35.300 and 31.11.
Ž-Willi	am E. Decker M.D.	10 CFR 35.300 an	d 35.400
X, Cha	d Michael Ruble, M.D.	40 CFR-35,100, 35 permitted by 35.3	5.200, iodine 131 diagnostic procedures 00 and 31.11.
14. For	sealed sources not associated	with 10 OFR Part 3	5 use the following conditions apply:
ĹÃ.	- 4 2 4. 運動を取り返り 2000年 (2000年) 1 11 がん 200	ficate of registration	r contamination at intervals not to exceed the issued by the U.S. Nuclear Regulatory ement State.
В.	the intervals specified in the	certificate of registra	ndicating that a leak test has been made within ition issued by the A.S. Nuclear Regulatory entire to the transfer, a sealed
	source received from anothe received.	gperson shall not be	publific use until tested and the test results
C.	gamma emitting material or n	ot more than 10 mic	n not more than 100 microcuries of beta and/or crocuries of alpha emitting material.
D.	when they are removed from been tested within the require	storage for use or to edileak test interval, for a period of mor	storage and are not being used. However, ransferred to another person, and have not they shall be tested before use or transfer. No eithan 10 years without being tested for
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- E. The leak test shall be capable of detecting the presence of 0.005 microcurie (185 becquerels) of radioactive material on the test sample. If the test reveals the presence of 0.005 microcurie (185 becquerels) or more of removable contamination, a report shall be filed with the U.S. Nuclear Regulatory Commission in accordance with 10 CFR 30.50(c)(2), and the source shall be removed immediately from service and decontaminated, repaired, or disposed of in accordance with Commission regulations.
- F. Tests for leakage and/or contamination, including leak test sample collection and analysis, shall be performed by the licensee or other persons specifically licensed by the U.S. Nuclear Regulatory Commission or an Agreement State to perform such services.
- G. Records of leak test results shall be kept in units of microcuries and shall be maintained for 3 years.

 //o. CFR 35.67
- 15. The licensee shall conduct a physical inventory every 3 months to account for all sources and/or devices received and possessed pursuant to 10 CFR 35.59, 10 CFR 35.400 and every 6 months for all other sources and/or devices. Records of inventories shall be maintained for 5 years from the date of each inventory, and shall include the information required in 10 CFR 35.59(g).
- 16. Sealed sources containing licensed material shall not be opened or sources removed from source holders by the licensee.
- 17. In addition to the possession limits in Item 8, the licensee shall further restrict the possession of licensed material to quantities below the minimum limit specified in 10 CFR 30.35(d) for establishing decommissioning financial assurance.
- 18. The licensee is authorized to transport licensed material in accordance with the provisions of 10 CER Part 71. Packaging and Transportation of Radioactive Material."

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- 19. 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 March 8, 2005;
 - B. Letters dated September 10, 1990, October 18, 2006, November 14, 2006, January 15, 2007, August 2007, October 3, 2007, December 12, 2008;
 - C. Facsimile dated April 20, 2005, transmitting letter dated October 29, 2004; and
 - D_r Facsimile letter dated April 2, 2007, October 29, 2007 and November 6, 2007.

FOR THE U.S. NUCCEAR REGULATORY COMMISSION

Date NOV 3 0 2009

Toye L. Simmons

Materials Licensing Branch

Region III

Domestic Shipments

· To qualify for the Letter rate, UPS Express Envelopes may only contain correspondence, urgent documents, and/or electronic media, and must weigh 8 oz. or less. UPS Express Envelopes containing items other than those listed or weighing more than 8 oz. will be billed by weight.

International Shipments

- . The UPS Express Envelope may be used only for documents of no commercial value, Certain countries consider electronic media as documents. Visit ups.com/importexport to verify if your shipment is classified as a document.
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LTR

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