



- Vascular Ultrasound • Bone Densitometry • Ultrasound
- X-Ray • CAT Scan • Mammography • Nuclear Medicine

October 6, 2008

United States Nuclear Regulatory Commission
Region III
2443 Warrenville Road, Suite 210
Lisle, IL. 60532-4352

RE: Amendment to NRC License No. 21-25921-01
MDO Imaging Associates, PLC

Attn: To Whom It May Concern:

We would like to amend our current NRC license to reflect the following changes.

1. We request approval for the use of Byproduct Material 35.300 for Written Directives. We have developed and will implement and maintain written Procedures in accordance with the requirements in 10 CFR 35.40, 35.41, 35.2040 and 35.2041.
2. We are updating Dr. John Frederick, D.O. and Dr. Robert Yochim, M.D., as authorized users for 10 CFR 35.300. They are listed as 35.300 authorized users on license #21-26740-01, amendment #10 (please find enclosed copy).
3. Please add Dr. Michael A. Gedwill, D.O. as an authorized user for 10 CFR 35.100 and 35.200. He is currently listed for these uses on license #21-26740-01, amendment #10 (please find enclosed copy).
4. We would like to remove Dr. David T. Salvati, D.O. from our NRC license.

We appreciate your assistance in this amendment.

If you have any questions or require additional information, please contact our consultant, Laura Luna at (734) 662-3197.

Sincerely,


Michael J. Sinelli, COO
MDO Imaging Associates, PLC


John J. Frederick, D.O., RSO
MDO Imaging Associates, PLC

RECEIVED OCT 15 2008

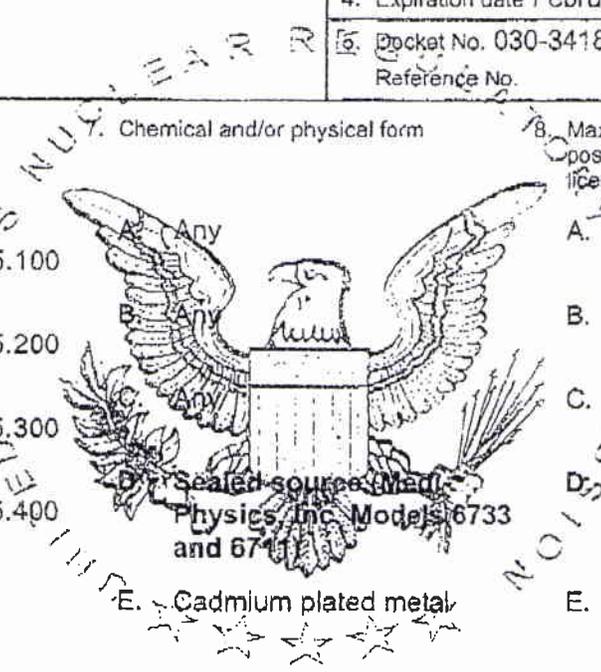
MATERIALS LICENSE

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.

<p style="text-align: center;">Licensee</p> <p>1. Genesys Regional Medical Center Nuclear Medicine Department</p> <p>2. One Genesys Parkway Grand Blanc, MI 48439</p>	<p>In accordance with letter dated July 30, 2007,</p> <p>3. License number 21-26740-01 is amended in its entirety to read as follows:</p> <p>4. Expiration date February 29, 2012</p> <p>5. Pocket No. 030-34188 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. Any byproduct material permitted by 10 CFR 35.400</p> <p>E. Uranium depleted in Uranium-235</p> <p>F. Cesium-137</p> <p>G. Gadolinium-153</p>	<p>7. Chemical and/or physical form</p> <p>A. Any</p> <p>B. Any</p> <p>C. Any</p> <p>D. Sealed source (Medi Physics, Inc. Models 6733 and 6741)</p> <p>E. Cadmium plated metal</p> <p>F. Sealed source (Amersham/Tech Ops Model 77032)</p> <p>G. Sealed sources (Isotope Products Laboratories Model A-3410 or Dupont Merck Pharmaceutical Co. Model NES-8426)</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. As needed (not to exceed 1 curie of I-131)</p> <p>D. One Curie</p> <p>E. As needed</p> <p>F. 165 millicuries</p> <p>G. 28 sources, not to exceed 240 millicuries total. Four replacement sources not to exceed 20 millicuries each</p>
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<p>9. Authorized Use:</p> <p>A. Any uptake, dilution and excretion study permitted by 10 CFR 35.100.</p> <p>B. Any imaging and localization study permitted by 10 CFR 35.200.</p> <p>C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300.</p>



**MATERIALS LICENSE
SUPPLEMENTARY SHEET**

License Number
21-26740-01

Docket or Reference Number
030-34188

Amendment No. 10

- D. Any manual brachytherapy procedure permitted by 10 CFR 35.400.
- E. Shielding in a linear accelerator.
- F. To be used in Amersham/Tech Ops Model 773 calibration device for survey instrument calibration.
- G. Twenty-eight sources of varying activities to be used in a Siemens Medical Systems Profile Attenuation Correction System transmission line source housing device for medical radiography in humans. Four sources in their shipping containers for replacement sources.

CONDITIONS

- 10. A. Licensed material may be used at the licensee's facilities located at One Genesys Parkway, Grand Blanc, Michigan.
- B. Licensed materials in Items 6.C and 6.E may be used at the licensee's facilities located at the Radiation Oncology Center, 302 Kensington Avenue, Flint, Michigan.
- 11. A. Radiation Safety Officer: John J. Frederick, D.O.
- B. Assistant Radiation Safety Officer (Brachytherapy): Ibrahim S. Abdulhay, Ph.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

Material and Use

Byung Ho Chang, M.D.	10 CFR 35.100, 35.200 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
John Dobson, M.D.	10 CFR 35.100, 35.200 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Anthony M. Parillo, M.D.	10 CFR 35.100, 35.200 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Roger J. Rohr, D.O.	10 CFR 35.100, 35.200 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
John J. Frederick, D.O.	10 CFR 35.100, 35.200, 35.300 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.

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Ronald E. Weigel, D.O.	10 CFR 35.100, 35.200, 35.300 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Khalid Latif, M.D.	10 CFR 35.100, 35.200, 35.300 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Robert J. Yochim, M.D.	10 CFR 35.100, 35.200, 35.300 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Haesook S. Kim, M.D.	10 CFR 35.300 and 35.400
Dong-Whan Oh, M.D.	10 CFR 35.300, limited to Sr-89 for treatment of metastatic bone disease and 35.400
Ibrahim S. Abdulhay, Ph.D.	Cesium-137 and/or 10 CFR 35.400 for survey instrument calibration only
Ahmed M. Akl, M.D.	10 CFR 35.300 and 35.400
Meketa Schlega, M.D.	10 CFR 35.100, 35.200, 35.300 (excluding iodine-131 for thyroid carcinoma)
Edward F. Martin, D.O.	10 CFR 35.100, 35.200, 35.300 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
John S. Morrison, D.O.	10 CFR 35.100, 35.200 and gadolinium-153 in Profile Attenuation Correction System devices for medical radiography.
Michael A. Gedwill, D.O.	10 CFR 35.100 and 35.200.

13. 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.
14. The licensee is authorized to receive, possess, and use sealed sources of gadolinium-153 where the radioactivity exceeds the maximum amount of radioactivity specified in this license provided:
- Such possession does not exceed the quantity per source specified in Item 8 by more than 20 percent for gadolinium-153; and
 - Records of the licensee show that no more than the maximum amount of radioactivity per source specified in this license was ordered from the supplier or transferor of the byproduct material; and
 - The levels of radiation for the Siemens Medical Systems E-cam Attenuation Correction Profile device do not exceed those specified in the Sealed Source and Device Registry Sheet.

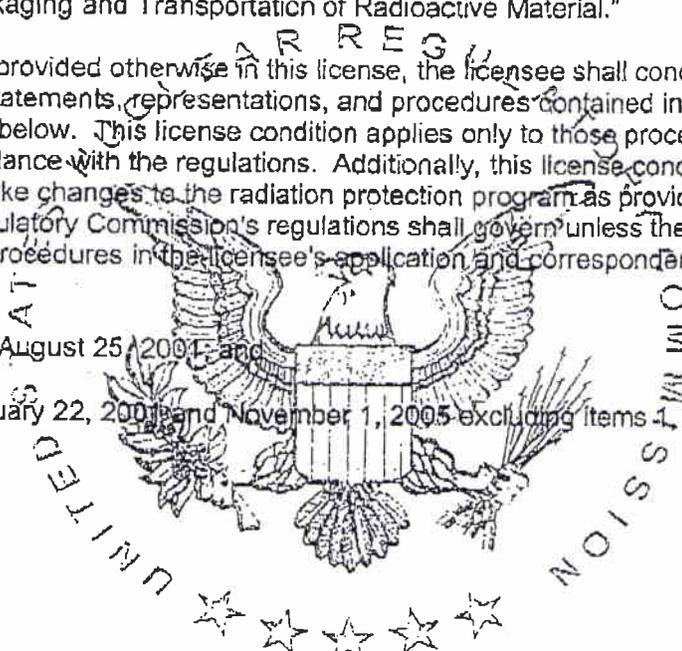
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- 15. Sealed sources containing licensed material shall not be opened or sources removed from source holders by the licensee.
- 16. The licensee shall follow the "Registry of Radioactive Sealed Sources and Devices Safety Evaluation of Device" No. IL-605-D-105-S for the Siemens E-cam Attenuation Correction Profile device as it pertains to description (installation), conditions of normal use, limitations and other considerations of use.
- 17. The licensee is authorized to transport licensed material in accordance with the provisions of 10 CFR Part 71 "Packaging and Transportation of Radioactive Material."
- 18. 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 August 25, 2001, and
 - B. Letters dated January 22, 2001, and November 1, 2005 excluding items 1, 2 and 3.



FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date NOV 01 2007

By *Toye L. Simmons*
 Toye L. Simmons
 Materials Licensing Branch
 Region III

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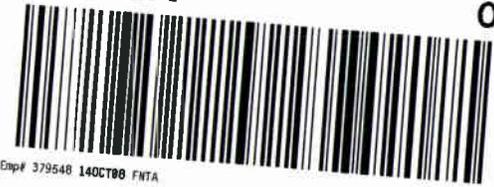
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FedEx Express Saver
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Third business day. Saturday Delivery NOT available.

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