UNITED STATES
NUCLEAR REGULATORY COMMISSION
REGION I
475 ALLENDALE ROAD
KING OF PRUSSIA, PENNSYLVANIA 19406-1415

April 10, 2007

Docket No. 03007584 License No. 37-06575-03
Control No. 139775

Patrick Alberts
Chief Operating Officer
Monongahela Valley Hospital, Inc.
1163 Country Club Road
Monongahela, PA 15063

SUBJECT: MONONGAHELA VALLEY HOSPITAL, INC., SECOND CORRECTED COPY OF LICENSE, CONTROL NO. 139775

Dear Mr. Alberts:

Enclosed is the second Corrected Copy of Amendment No. 43 for License No. 37-06575-03. In accordance with the telephone call on April 10, 2007, Condition Nos. 6.D, 7.D, 8.D, and 9.D, as well as Condition 12, have been changed to reflect the fact that you do not possess 10 CFR 35.400 material and do not have a physician user authorized for such material.

We apologize for any inconvenience this error may have caused.

Sincerely,

Original signed by Pamela J. Henderson

Pamela J. Henderson, Chief
Medical Branch
Division of Nuclear Materials Safety

Enclosure:
Second Corrected Copy of Amendment No. 43

cc:
Michael J. Semon, Radiation Safety Officer

DOCUMENT NAME: C:\FileNet\ML071010009.wpd

SUNSI Review Complete: RMcKinley
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OFFICE	DNMS/RI	N	DNMS/RI	N	DNMS/RI			
NAME	RMcKinley/RWM		PHenderson/PJH					
DATE	4/10/2007		4/10/2007					

OFFICIAL RECORD COPY

Amendment No. 43

CORRECTED COPY NO. 2

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	In accordance with the letter dated December 6, 2006
1. Monongahela Valley Hospital, Inc.	3. License number 37-06575-03 is amended in its entirety to read as follows:
2. 1163 Country Club Road Monongahela, Pennsylvania 15063	4. Expiration date October 31, 2014
	5. Docket No. 030-07584 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
A. Any byproduct material permitted by 10 CFR 35.100	A. Any	A. As needed
B. Any byproduct material permitted by 10 CFR 35.200	B. Any	B. As needed
C. Any byproduct material permitted by 10 CFR 35.300	C. Any	C. 1000 millicuries
D. Any byproduct material permitted by 10 CFR 31.11	D. Prepackaged Kits	D. 5 millicuries
E. Cesium 137	E. Sealed Sources (Compagnie ORIS Industrie Model CSM-3 series, distributed by CIS-US, Inc.)	E. Not to exceed 150 millicuries per source and 1 curie total

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.
C. Any diagnostic study or therapy procedure permitted by 10 CFR 35.300 for which the patient may be released under the provisions of 10 CFR 35.75
D. In vitro studies.
E. For possession and storage only.

MATERIALS LICENSE
SUPPLEMENTARY SHEET

CORRECTED COPY

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Docket or Reference Number
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Amendment No. 43
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CONDITIONS

10. Licensed material may be used or stored only at the licensee's facilities located at 1163 Country Club Road, Monongahela, Pennsylvania.

11. The Radiation Safety Officer for this license is Michael J. Semon, 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	Material and Use
Abdul S. Chaudry, M.D.	35.100; 35.200; 35.300; In vitro studies
Vinod N. Chablani, M.D.	35.100; 35.200; 35.300; In vitro studies
Douglas Wilson, M.D.	35.100; 35.200; Oral administration of sodium iodide iodine-131 for imaging and localization studies and treatment of hyperthyroidism and cardiac dysfunction; In vitro studies
M. Mohsin Rahman, M.D.	35.100; 35.200; 35.300

    C. The following individuals are authorized users for non-medical uses as indicated.

Authorized Users	Material and Use
Michael J. Semon, M.S.	Cesium 137 (Possession and storage only)

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. Sealed sources or detector cells containing licensed material shall not be opened or sources removed from source holders by the licensee.

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Amendment No. 43
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15. The licensee shall conduct a physical inventory every six months, or at other intervals approved by the U.S. Nuclear Regulatory Commission, to account for all sources and/or devices received and possessed under the license. Records of inventories shall be maintained for 5 years from the date of each inventory and shall include the radionuclides, quantities, manufacturer's name and model numbers, and the date of the inventory.

16. A. Sealed sources shall be tested for leakage and/or contamination at intervals not to exceed the intervals specified in the certificate of registration issued by the U.S. Nuclear Regulatory Commission under 10 CFR 32.210 or under equivalent regulations of an Agreement State.

    B. In the absence of a certificate from a transferor indicating that a leak test has been made within the intervals specified in the certificate of registration issued by the U.S. Nuclear Regulatory Commission under 10 CFR 32.210 or under equivalent regulations of an Agreement State, prior to the transfer, a sealed source received from another person shall not be put into use until tested and the test results received.

    C. Sealed sources need not be tested if they contain only hydrogen-3; or they contain only a radioactive gas; or the half-life of the isotope is 30 days or less; or they contain not more than 100 microcuries of beta- and/or gamma-emitting material or not more than 10 microcuries of alpha-emitting material.

    D. Sealed sources need not be tested if they are in storage and are not being used; however, when they are removed from storage for use or transferred to another person and have not been tested within the required leak test interval, they shall be tested before use or transfer. No sealed source shall be stored for a period of more than 10 years without being tested for leakage and/or contamination.

    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 by 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 5 years.

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."

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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 July 26, 2004 [ML042220301]
B. Letter dated September 10, 2004 [ML042660502]
C. Letter dated March 18, 2005 [ML051160300]
D. Letter dated May 25, 2005 [ML051600025]

For the U.S. Nuclear Regulatory Commission

Date April 10, 2007

By ***Original signed by Richard McKinley***

Richard McKinley
Medical Branch
Division of Nuclear Materials Safety
Region I
King of Prussia, Pennsylvania 19406