

MATERIALS LICENSE

Amendment No. 65

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.

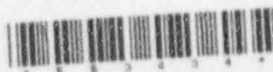
302426

Licensee		In accordance with letter dated March 10, 1997	
1. Wausau Hospital Center		3. License Number 48-01032-01 is amended in its entirety to read as follows:	
2. 333 Pine Ridge Boulevard Wausau, WI 54401		4. Expiration Date December 31, 2000	
		5. Docket or Reference No. 030-03416	
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 identified in 10 CFR 35.100	A. Any radiopharmaceutical identified in 10 CFR 35.100	A. As needed	
B. Any byproduct material identified in 10 CFR 35.200	B. Any radiopharmaceutical identified in 10 CFR 35.200	B. As needed	
C. Any byproduct material identified in 10 CFR 35.300	C. Any radiopharmaceutical identified in 10 CFR 35.300	C. As needed	
D. Any byproduct material identified in 10 CFR 35.400	D. Any brachytherapy source identified in 10 CFR 35.400	D. 1200 millicuries	
E. Any byproduct material identified in 10 CFR 35.500	E. Sealed sources identified in 10 CFR 35.500	E. As needed	
F. Any byproduct material identified in 10 CFR 31.11	F. Prepackaged Kits	F. As needed	

120121

9. Authorized Use:

- A. Medical use described in 10 CFR 35.100.
- B. Medical use described in 10 CFR 35.200.



COPY 80

**MATERIALS LICENSE
SUPPLEMENTARY SHEET**

License Number

48-01032-01

Docket or Reference Number

030-03416

Amendment No. 65

- C. Medical use described in 10 CFR 35.300.
- D. Medical use described in 10 CFR 35.400.
- E. Medical use described in 10 CFR 35.500 in devices which have been evaluated and approved for licensing purposes by the U.S. Nuclear Regulatory Commission or an Agreement State.
- F. In vitro studies.

CONDITIONS

10. Locations of Use:

- A. Licensed material shall be received, stored and used at the licensee's facility located at 333 Pine Ridge Boulevard, Wausau, Wisconsin.
- B. Licensed material identified in 10 CFR 35.100, 35.200 (excluding generators), 35.500 and 31.11 may be used at temporary job sites of medical care facilities anywhere in the United States where the U.S. Nuclear Regulatory Commission maintains jurisdiction for regulating the use of licensed material.

11. Radiation Safety Officer: Raymond F. Wery, M.S.

12. Authorized Users:

- A. Dolorus Buchler, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- B. Jacob H. Martens, M.D., for material in 10 CFR 35.100, 35.200, 35.300, 35.400 and 31.11.
- C. Paul Harari, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- D. G. H. Brister, M.D., for material in 10 CFR 35.100, 35.200, 35.300 and 31.11.
- E. E. B. Koschmann, M.D., for material in 10 CFR 35.100, 35.200, and 31.11.
- F. Henry Kanemoto, M.D., for material in 10 CFR 35.100, 35.200 and 31.11.
- G. William Mahony, M.D., for material in 10 CFR 35.100, 35.200 and 31.11.
- H. A. R. Bourque, M.D., for material in 10 CFR 35.400.
- I. Vina Luthra, M.D., for material in 10 CFR 35.100, 35.200 and 31.11.

COPY

MATERIALS LICENSE
SUPPLEMENTARY SHEETLicense Number
48-01032-01Docket or Reference Number
030-03416

Amendment No. 65

- J. Philip R. Albert, M.D., for material in 10 CFR 35.100, 35.200, 35.300 and 31.11.
- K. Guy W. Holmes, for material in 10 CFR 31.11.
- L. John R. Sorenson, M.D., for material in 10 CFR 35.100, 35.200, 35.300, and 31.11.
- M. Donald M. Nowinski, Jr., M.D., for material in 10 CFR 35.100, 35.200, iodine-131 for treatment of hyperthyroidism and 31.11.
- N. Christopher J. Kelley, M.D., for material in 10 CFR 35.100, 35.200 and 31.11.
- O. Timothy Kinsella, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- P. Minesh Mehta, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- Q. Mark Ritter, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- R. Richard Steeves, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- S. Judith Anne Stitt, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- T. George W. Wirtanen, M.D., for material in 10 CFR 35.300, 35.400 and 35.500.
- U. George R. Edmonson, M.D., for material in 10 CFR 35.100, 35.200 and 31.11.
- V. Darryl R. Barton, M.D., for material in 10 CFR 35.300 and 35.400.
- W. James S. Collison, M.D., for material in 10 CFR 35.100, 35.200 and 35.300.
- X. Bruce S. Salo, M.D., for material in 10 CFR 35.100, 35.200, 35.300, 35.500 and 31.11.
- Y. Joseph A. Heitz, M.D., for material in 10 CFR 35.100 and 35.200.
- Z. Seyed Mohsen Sharifi Takieh, M.D., for material in 10 CFR 35.100 and 35.200 (limited to studies of cardiac imaging and functions).
- AA. Paul A. Luetmer, M.D., for material in 10 CFR 35.100 and 35.200 (limited to studies of cardiac imaging and functions).

COPY

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SUPPLEMENTARY SHEET**

License Number

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Amendment No. 65

13. Pursuant to Title 10, Chapter 1, Code of Federal Regulations, Part 40, "Domestic Licensing of Source Material," the licensee is authorized to possess, use, transfer, and import up to 999 kilograms of depleted uranium contained as shielding material in the molybdenum-99/technetium-99m generators authorized by this license.
14. 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.
15. The licensee may transport licensed material in accordance with the provisions of 10 CFR Part 71, "Packaging and Transportation of Radioactive Material."
16. This license is based on the licensee's statements and representations listed below:
 - A. Application dated May 30, 1990;
 - B. Letters dated October 7, 1990 (with attachments), November 8, 1990 (with attachments), January 8, 1991, June 24, 1991 (with enclosures), July 31, 1991 (with enclosures), October 24, 1991 (signed by Bruce Thomadsen, Ph.D.), August 13, 1991, and February 12, 1992; May 10, 1993 (with attachments, excluding request to add physician user);
 - C. Letters received October 2, 1991 (with enclosures, responding to NRC August 13, 1991 letter) October 18, 1991 and November 13, 1992; and
 - D. Letters dated March 30, 1995 and May 5, 1995 (with attachments).

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date

MAY 29 1997

By

[Signature]
Nuclear Materials Licensing Branch, Region III

COPY

S5

(FOR LFMS USE)
INFORMATION FROM LTS

BETWEEN:

License Fee Management Branch, ARM
and
Regional Licensing Sections

Program Code: 02120
Status Code: 0
Fee Category: 7C 2B
Exp. Date: 20001231
Fee Comments:
Decon Fin Assur Req'd: N

LICENSE FEE TRANSMITTAL

A. REGION

1. APPLICATION ATTACHED

Applicant/Licensee: WAUSAU HOSPITAL CENTER
Received Date: 970317
Docket No: 3003416
Control No.: 302426
License No.: 48-01032-01
Action Type: Amendment

2. FEE ATTACHED

Amount: 440
Check No.: 151660

3. COMMENTS

Signed
Date

D. Hersey
3-17-97

B. LICENSE FEE MANAGEMENT BRANCH (Check when milestone 03 is entered /)

1. Fee Category and Amount: 7C 2B \$440

2. Correct Fee Paid. Application may be processed for:

Amendment ☒
Renewal ☐
License ☐

3. OTHER

Signed
Date

SC 3/20/97

MAR 24 1997

Log	Mar 9 III
Remitter	
Check No.	151660
Amount	\$440
Fee Category	7C
Type of Fee	AMD
Date Check Rec'd	3/19/97
Date Completed	3/20/97
By:	SC

1997 MAR 19 AM 11:25



March 10, 1997

U.S. Nuclear Regulatory Commission
Region III Office
801 Warrenville Road
Lisle, Ill 60532-4351

Wausau Hospital would like to amend its NRC License (# 48-01032-01) to add Dr. Paul A. Luetmer as an authorized user for material in 10 CFR 35.100 and 35.200 (limited to studies of cardiac imaging and function).

Enclosed you will find copies of Dr. Luetmer's training and experience as described in the NRC regulatory guide 10.8, Guide for the Preparation of Applications for Medical Use Programs and a copy of the NRC license for the institution in which Dr. Luetmer received his training. A check for \$440.00 for this amendment is also enclosed.

Should you have questions about this application please contact Ray Wery, Wausau Hospital Radiation Safety Officer, at 715-847-2292.

Sincerely yours;

Gary Bezucha
Vice President, Episodic Care
Wausau Hospital
333 Pine Ridge Boulevard
Wausau, WI 54401

RECEIVED
MAR 17 1997
REGION III

Pm: 3-13-97

302426



February 12, 1997

To Whom It May Concern:

Paul Luetmer, M.D. has completed a one year course conducted by the Medical Physics Section at the Medical Center Hospital of Vermont, Fletcher Allen Health Care. This course consisted of lectures, demonstrations, and reading and problem assignments designed to meet the course requirements specified in 10 CFR Part 35.920.

Further documentation is on file in the Medical Physics office of Fletcher Allen Health Care and is available on request.

Sincerely,

James H. Goodwin, M.S.
Chief Medical Physicist
American Board of Radiology Certified in Radiological Physics
Associate Professor, Department of Radiology, University of
Vermont

Marleen M. Moore, M.S.
Medical Physicist and Radiation Safety Officer
American Board of Radiology Certified in Medical Nuclear Physics
Associate Professor, Department of Radiology, University of
Vermont

**EXHIBIT 2
SUPPLEMENT A**

SUPPLEMENT		U.S. NUCLEAR REGULATORY COMMISSION		
TRAINING AND EXPERIENCE AUTHORIZED USER OR RADIATION SAFETY OFFICER				
1. NAME OF PROPOSED AUTHORIZED USER OR RADIATION SAFETY OFFICER		2. FOR PHYSICIANS, STATE OR TERRITORY WHERE LICENSED		
Paul A. Luetmer		Vermont		
3. CERTIFICATION				
SPECIALTY BOARD A	CATEGORY B	MONTH AND YEAR CERTIFIED C		
4. TRAINING RECEIVED IN BASIC RADIOISOTOPE HANDLING TECHNIQUES				
FIELD OF TRAINING A	LOCATION AND DATE(S) OF TRAINING B	TYPE AND LENGTH OF TRAINING		
		CLOCK HOURS IN LECTURE OR LABORATORY	CLOCK HOURS OF SUPERVISED ON-THE-JOB EXPERIENCE	
a. RADIATION PHYSICS AND INSTRUMENTATION	Medical Center Hosp. of Vermont 7/95 - 6/96	20 hrs lecture 30 hrs assigned	50 hrs	
b. RADIATION PROTECTION	Medical Center Hosp. of Vermont 7/95 - 6/96	16 hrs lecture 20 hrs assigned		
c. MATHEMATICS PERTAINING TO THE USE AND MEASUREMENT OF RADIOACTIVITY	Medical Center Hosp. of Vermont 7/95 - 6/96	5 hrs lecture 10 hrs assigned		
d. RADIATION BIOLOGY	Medical Center Hosp. of Vermont 7/95 - 6/96	15 hrs lecture 20 hrs assigned		
e. RADIOPHARMACEUTICAL CHEMISTRY	Medical Center Hosp. of Vermont 7/95 - 6/96	10 hrs lecture	10 hrs	
5. EXPERIENCE WITH RADIATION. (Actual use of Radioisotopes or Equivalent Experience)				
ISOTOPE	mCi USED AT ONE TIME	LOCATION	CLOCK HOURS	TYPE OF USE
^{99m} Tc	30	Medical Center Hosp. of Vermont	950	Cardiac perfusion, ventriculography
²⁰¹ Tl	2	Medical Center Hosp. of Vermont	10	Cardiac perfusion, myocardial viability

EXHIBIT 3
SUPPLEMENT B

U. S. NUCLEAR REGULATORY COMMISSION

SUPPLEMENT

PRECEPTOR STATEMENT

Supplement B must be completed by the applicant physician's preceptor. If more than one preceptor is necessary to document experience, obtain a separate statement from each.

KEY TO COLUMN C

1. PROPOSED PHYSICIAN USER'S NAME AND ADDRESS

PULL NAME

Paul A. Luchner

STREET ADDRESS

6 Village Vale Drive

CITY

STATE

ZIP CODE

Shelburne

UT

05702

KEY TO COLUMN C

PERSONAL PARTICIPATION SHOULD CONSIST OF:

- 1-Supervised examination of patients to determine the suitability for radiodiagnosis diagnosis and/or treatment and recommendation for prescribed dosage.
- 2-Collaboration in dose calibration and actual administration of dose to the patient including calculation of the radiation dose, related measurements and plotting of data.
- 3-Adequate period of training to enable physician to manage radioactive patients and follow patients through diagnosis and/or course of treatment.

2. CLINICAL TRAINING AND EXPERIENCE OF ABOVE NAMED PHYSICIAN	
1. Name of the Physician	Dr. [Name]
2. Address	[Address]
3. Telephone	[Telephone]
4. Specialties	[Specialties]
5. Training	[Training]
6. Experience	[Experience]
7. Other	[Other]

[illegible]

PROPOSED PHYSICIAN USER

Paul A. Lufner

PRECEPTOR STATEMENT (Continued)

2. CLINICAL TRAINING AND EXPERIENCE OF ABOVE NAMED PHYSICIAN (Continued)

ISOTOPE A	CONDITIONS DIAGNOSED OR TREATED B	NUMBER OF CASES INVOLVING PERSONAL PARTICIPATION C	COMMENTS (Additional information or comments may be submitted in duplicate on separate sheets.) D
P-32 (Sodium)	TREATMENT OF POLYCYTHEMIA VERA, LEUKEMIA, AND BONE METASTASES		
P-32 (Colloid)	INTRACAVITARY TREATMENT		
I-131	TREATMENT OF THYROID CARCINOMA		
	TREATMENT OF HYPERTHYROIDISM		
Au-198	INTRACAVITARY TREATMENT		
Co-60 or Co-137	INTERSTITIAL TREATMENT		
	INTRACAVITARY TREATMENT		
I-125 or Ir-192	INTERSTITIAL TREATMENT		
Co-60 or Co-137	TELE THERAPY TREATMENT		
Sr-90	TREATMENT OF EYE DISEASE		
	RADIOPHARMACEUTICAL PREPARATION		
Mo-99/ Tc-99m	GENERATOR	2	
Sr-90/ Y-90	GENERATOR		
Tc-99m	REAGENT KITS	1	
Other			

3. DATES AND TOTAL NUMBER OF HOURS RECEIVED IN CLINICAL RADIOISOTOPE TRAINING

LOCATION
Medical Center Hosp of Vermont
(Name changed to Fletcher Allen Health Care)
Burlington Vermont

DATES
7/1/93 -
6/30/97

CLOCK HOURS OF EXPERIENCE
1160

4. THE TRAINING AND EXPERIENCE INDICATED ABOVE WAS OBTAINED UNDER THE SUPERVISION OF:

a. NAME OF SUPERVISOR

Kenneth A. Brown M.D.

b. NAME OF INSTITUTION

Fletcher Allen Health Care

c. MAILING ADDRESS

Box 111 Colchester Ave

d. CITY

Burlington VT

e. MATERIALS LICENSE NUMBER(S)

44-10187-03

5. PRECEPTOR'S SIGNATURE

7. PRECEPTOR'S NAME (Please type or print)

Kenneth A. Brown, M.D.

8. DATE

3/3/97

TOTAL P. 02

No. 38711



State of Wisconsin

Department of Regulation and Licensing

Medical Examining Board

Hereby certifies that

PAUL A LUETMER, MD

was granted a license to practice

Medicine and Surgery

in the State of Wisconsin in accordance with Wisconsin law
on the 25th day of April, 1997.

The authority granted herein must be renewed each biennium by the granting authority.

In witness thereof, the State of Wisconsin

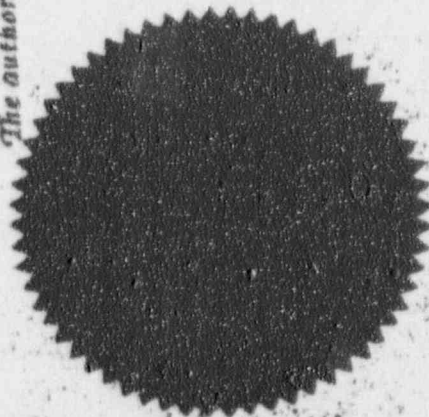
Medical Examining Board

has caused this certificate to be issued under

the seal of the Department of Regulation and Licensing.

This certificate issued this 1st day of May, 1997.

Patricia L. D.
Blair J. J. J.
Secretary





State of Vermont



The Board of Medical Practice
Grants this license as a
Physician

Paul Anthony Lettmer MD
#6 Village Way Drive
Chester VT 05482

2-1-96

LICENSE NUMBER 2430
UNDER THE PROVISIONS OF V.S.A.

EFFECTIVE DATE 1/1/96
TITLE
CHARTER

EXPIRES 11-30-1998

SPECIAL INSTRUCTIONS



State of Vermont



Physician

Paul Anthony Lettmer MD
#6 Village Way Drive
Chester VT 05482
LICENSE NUMBER 2430
EXPIRES 11/30/1998

EFFECTIVE DATE 1/1/1996
CHARTER

11-30-98

MATERIALS LICENSE

Amendment No. 53

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 March 29, 1995,	
1. Fletcher Allen Health Care		3. License Number 44-10187-03 is amended in its entirety to read as follows:	
2. Colchester Avenue Burlington, Vermont 05401		4. Expiration Date December 31, 1997	
		5. Docket or Reference No. 030-03289	
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 identified in 10 CFR 35.100	A. Any radiopharmaceutical identified in 10 CFR 35.100	A. As needed	
B. Any byproduct material identified in 10 CFR 35.200	B. Any radiopharmaceutical identified in 10 CFR 35.200	B. As needed	
C. Any byproduct material identified in 10 CFR 35.300	C. Any radiopharmaceutical identified in 10 CFR 35.300	C. 500 millicuries	
D. Any byproduct material identified in 10 CFR 35.400	D. Any brachytherapy source identified in 10 CFR 35.400	D. 2.5 curies	
E. Any byproduct material identified in 10 CFR 31.11	E. Prepackaged Kits	E. 3 millicurie	
F. Hydrogen 3	F. Any	F. 1 millicurie	
G. Depleted Uranium	G. Cadmium plated metal	G. 91 kilograms	
9. Authorized use			
A. Any uptake, dilution and excretion procedure approved in 10 CFR 35.100.			
B. Any imaging and localization procedure approved in 10 CFR 35.200.			
C. Any radiopharmaceutical therapy procedure approved in 10 CFR 35.300.			
D. Any brachytherapy procedure approved in 10 CFR 35.400.			
E. through F. <u>In vitro</u> studies.			
G. Shielding in a linear accelerator.			

CONDITIONS

10. Licensed material may be used only at the licensee's facilities located at 111 Colchester Avenue, Burlington, Vermont; 101 College Parkway, Colchester, Vermont; and the Degoesbriand Laboratory, One Prospect Street, Burlington, Vermont.
11. The Radiation Safety Officer for this license is Marleen M. Moore, M.S.

**MATERIALS LICENSE
SUPPLEMENTARY SHEET**

License number
44-10187-03

Docket or Reference number
030-03289

Amendment No. 53

12. Licensed material listed in Item 6 above is only authorized for use by, or under the supervision of, the following individuals for the materials and uses indicated:

Authorized Users

Material and Use

Jorge Calles, M.D.	35.100
Robert Naylor, M.D.	35.100; 35.200
Jonathan Fairbank, M.D.	35.100; 35.200; 35.300
Linda Roe, M.D.	35.100; 35.200; 35.300
Norman Sturtevant, M.D.	35.100; 35.200; 35.300
Kenneth Brown, M.D.	35.200 for cardiovascular clinical studies
Thomas Roland, M.D.	35.400
Peter Swift, M.D.	35.400
John McDay, M.D.	35.400
Mary Tang, M.D.	in vitro studies
Bruce McPherson, M.D.	in vitro studies
Marleen M. Moore, M.S.	Depleted Uranium

13. The licensee shall conduct a physical inventory every three months to account for all sealed sources and devices containing licensed material received and possessed pursuant to 10 CFR 35.59 and 35.400 and every six months for all other sealed sources and devices.
14. The licensee is authorized to hold radioactive material with a physical half-life of less than 65 days for decay-in-storage before disposal in ordinary trash, provided:
- Waste to be disposed of in this manner shall be held for decay a minimum of ten half-lives.
 - Before disposal as ordinary trash, the waste shall be surveyed at the container surface with the appropriate survey instrument set on its most sensitive scale and with no interposed shielding to determine that its radioactivity cannot be distinguished from background. All radiation labels shall be removed or obliterated.
 - A record of each such disposal permitted under this License Condition shall be retained for three years. The record must include the date of disposal, the date on which the byproduct material was placed in storage, the radionuclides disposed, the survey instrument used, the background dose rate, the dose rate measured at the surface of each waste container, and the name of the individual who performed the disposal.

**MATERIALS LICENSE
SUPPLEMENTARY SHEET**

License number
44-10187-03

Docket or Reference number
030-03289

Amendment No. 53

15. Notwithstanding the requirements of 10 CFR 35.315(a)(4) the licensee may determine dose rates in contiguous restricted and unrestricted areas as described in the letter dated October 9, 1992.
16. Radioactive waste generated shall be stored in accordance with the statements, representations, and procedures included with the waste storage plan described in the licensee's letter/application dated August 14, 1992.
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, except for minor changes in the medical use radiation safety procedures as provided in 10 CFR 35.31. 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. Letter dated September 25, 1991
 - B. Letter dated August 14, 1992
 - C. Letter dated October 9, 1992
 - D. Letter dated December 19, 1994
 - E. Letter dated March 29, 1995
 - F. Letter dated September 11, 1995

Date SEP 25 1995

For the U.S. Nuclear Regulatory Commission

By

Eric H. Phelps

Nuclear Materials Safety Branch
Region I

King of Prussia, Pennsylvania 19406

MAY 29 1997

Gary Bezucha
Vice President
Wausau Hospital Center
333 Pine Ridge Boulevard
Wausau, WI 54401

Dear Mr. Bezucha:

Enclosed is Amendment No. 65 to your NRC Material License No. 48-01032-01 in accordance with your request.

Please review the enclosed document carefully and be sure that you understand all conditions. If there are any errors or questions, please notify the U.S. Nuclear Regulatory Commission, Region III office at (630) 829-9887 so that we can provide appropriate corrections and answers.

Please be advised that your license expires at the end of the day, in the month, and year stated in the license. Unless your license has been terminated, you must conduct your program involving byproduct materials in accordance with the conditions of your NRC license, representations made in your license application, and NRC regulations. In particular, note that you must:

1. Operate in accordance with NRC regulations 10 CFR Part 19, "Notices, Instructions and Reports to Workers; Inspections," 10 CFR Part 20, "Standards for Protection Against Radiation," and other applicable regulations.
2. Notify NRC, in writing, within 30 days:
 - a. When an authorized user or Radiation Safety Officer permanently discontinues performance of duties under the license or has a name change; or
 - b. When the mailing address listed on the license changes. (No fee is required if the location of byproduct material remains the same.)
3. In accordance with 10 CFR 30.36(b) and/or license condition, notify NRC, promptly, in writing, and request termination of the license when you decide to terminate all activities involving materials authorized under the license.

302424

4. Request and obtain a license amendment before you:
 - a. Receive or use byproduct material for a clinical procedure permitted under Part 35 but not permitted by your license issued pursuant to this Part;
 - b. Permit anyone, except individuals described in 10 CFR 35.13(b), to work as an authorized user under the license;
 - c. Change Radiation Safety Officers;
 - d. Order byproduct material in excess of the amount, or radionuclide, or form different than authorized on the license;
 - e. Add or change the areas of use or address or addresses of use identified in the license application or on the license; or
 - f. Change ownership of your organization.
5. Submit a complete renewal application with proper fee or termination request at least 30 days before the expiration date of your license. You will receive a reminder notice approximately 90 days before the expiration date. Possession of byproduct material after your license expires is a violation of NRC regulations. A license will not normally be renewed, except on a case-by-case basis, in instances where licensed material has never been possessed or used.

In addition, please note that NRC Form 313 requires the applicant, by his/her signature, to verify that the applicant understands that all statements contained in the application are true and correct to the best of the applicant's knowledge. The signatory for the application should be the licensee or certifying official rather than a consultant.

You will be periodically inspected by NRC. Failure to conduct your program in accordance with NRC regulations, license conditions, and representations made in your license application and supplemental correspondence with NRC will result in enforcement action against you. This could include issuance of a notice of violation, or imposition of a civil penalty, or an order suspending, modifying or revoking your license as specified in the General Statement of Policy and Procedure for NRC Enforcement Actions. Since serious consequences to employees and the public can result from failure to comply with NRC requirements, prompt and vigorous enforcement action will be taken when

G. Bezucha

-3-

dealing with licensees who do not achieve the necessary meticulous attention to detail and the high standard of compliance which NRC expects of its licensees.

Sincerely,

Original Signed By
Gidget Watson
Nuclear Materials Licensing Branch

License No.: 48-01032-01
Docket No.: 030-03416

Enclosure: Amendment No. 65

DOCUMENT NAME: M:\03003416.CL7

To receive a copy of this document, indicate in the box: "C" = Copy without attachment/enclosure "E" = Copy with attachment/enclosure "N" = No copy

OFFICE	DNMS/RIII								
NAME	GWATSON:jaw								
DATE	05/28/97 GW								

OFFICIAL RECORD COPY



UNITED STATES
NUCLEAR REGULATORY COMMISSION

REGION III
801 WARRENVILLE ROAD
LISLE, ILLINOIS 60532-4351

March 18, 1997

Raymond F. Wery, M.S.
Radiation Safety Officer
Wausau Hospital Center
333 Pine Ridge Boulevard
Wausau, WI 54401

SUBJECT: ACKNOWLEDGEMENT OF CORRESPONDENCE
(Letter Dated 03/10/97)

Dear Licensee:

In response to your request, we have completed the initial processing, which is an administrative review of your application for a(n):

☐ New License ☒ Amendment ☐ Renewal
☐ Termination ☐ Auth User (Amendment not required)
☐ Other _____

No administrative deficiencies were identified during this initial review. However, it should be noted that a technical review may identify omissions in the submitted information.

It appears that your request is routine (see 1-3 below, as applicable).

1. New and amendment actions are normally processed within 90 days, unless we find major deficiencies, or policy issues requiring central program office assistance.
2. Renewal actions are normally processed within 180 days, however, under timely filing (before expiration), you may continue to operate under your existing license.
3. Termination actions are normally processed within 90 days, unless confirmatory surveys following decontamination/decommissioning activities are involved.

A copy of your correspondence has been forwarded to our Licensing Fee and Debt Collection Branch (301/415-6097) for approval of the fee category and amount, if required.

If you have a compelling safety or business-related reason for requesting expedited review, please contact the Materials Licensing Branch at (630) 829-9887. We will try to complete your request as soon as practicable. Any correspondence about this request should reference the control number.

Nuclear Materials Support Branch

Mail Control No. 302426
License No. 48-01032-01