UNITED STATES
NUCLEAR REGULATORY COMMISSION
WASHINGTON, D.C. 20555-0001

January 28, 2003

Robert Wood Johnson Medical School
ATTN: Eugene G. Martin, Ph.D.
Associate Professor of Pathology
and Laboratory Medicine
Administrative Director, UDL
University Diagnostic Laboratories
One Robert Wood Johnson Place -MEB 212
New Brunswick, NJ 08901

Dear Dr. Martin:

This letter is in reference to your letter dated January 9, 2003, requesting that your NRC License #9205 be terminated as the clinical laboratory has been shut down and is no longer in service effective November 2002.

Your request to repeal registration 9205 for the general license under 10 CFR Part 31.11 has been granted.

I have removed your license from our files.

If you have any questions, please feel free to contact me at (301) 415-8140.

Sincerely,

Traci Kime

Traci Kime, Licensing Assistant
Materials Safety & Inspection Branch
Division of Industrial
and Medical Nuclear Safety
Office of Nuclear Material Safety
and Safeguards

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/RA/

Traci Kime, Licensing Assistant
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DOCUMENT NAME: P.\RobertWoodJohnson483 wpd

To receive a copy of this document, indicate in the box: "C"= Copy w/o att/encl. "E" = Copy w/att/encl."N" = No copy

OFC	MSIB			
NAME	Traci Kime			
DATE	1/28/2003			

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UMDNJ
ROBERT WOOD JOHNSON
MEDICAL SCHOOL
University of Medicine & Dentistry of New Jersey

Department of Pathology and Laboratory Medicine
MEB 212

January 9, 2003

Materials Safety Branch (T-8 F5)
Division of Industrial and Medical Nuclear Safety
Office of Nuclear Material Safety and Safeguards
U.S Nuclear Regulatory Commission
Washington D.C. 20555-0001

Re: Termination of Registration Certificate Number 9205

Dear Sir or Madam:

We were granted the enclosed Registration Certificate Number 9205 for - *in vitro* Testing with Byproduct Material under General License in September 2001.

We are requesting that the Registration Certificate 9205 be terminated as the clinical laboratory has been shut down and is no longer in service effective November 2002.

All radioactive materials were disposed of through the Rutgers University, Rutgers Environmental Health and Safety (REHS) Department, which is contracted by UMDNJ to perform Health Physics services for their Piscataway and New Brunswick campuses REHS also performed closeout surveys for fixed and removable contamination and found all results were at or below background levels

Thank you for your assistance, if you have any questions please feel free to contact me.

Sincerely,

Eugene G. Martin, Ph.D.
Associate Professor of Pathology & Laboratory Medicine
Administrative Director, UDL

Cc: Mary Aldrich
Evan Cadoff, M.D.

:C FORM 483 (999)

U.S. NUCLEAR REGULATORY COMMISSION

REGISTRATION CERTIFICATE -- *In vitro* TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

APPROVED BY OMB: NO. 3150-0038 EXPIRES: 07/31/2002

Estimated burden per response to comply with this mandatory collection request: 7 minutes. The validated registration serves as evidence to suppliers of byproduct material that the registrant is entitled to receive the byproduct material. Send comments regarding burden estimate to the Records Management Branch (T-6 F33), U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001, or by Internet e-mail to bjs1@nrc.gov, and to the Desk Officer, Office of Information and Regulatory Affairs, NEOB-10202, (3150-0038), Office of Management and Budget, Washington, DC 20503. If a means used to impose an information collection does not display a currently valid OMB control number, the NRC may not conduct or sponsor, and a person is not required to respond to, the information collection.

:ction 31.11 of 10 CFR 31 establishes a general license authorizing physicians, clinical laboratories, hospitals, and veterinarians in the actice of veterinary medicine to possess certain small quantities of byproduct material for *in vitro* clinical or laboratory tests not involving e internal or external administration of the byproduct material or the radiation therefrom to human beings or animals. Possession of product material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, hospital, or veterinarian in the practice of terinary medicine, has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with a registration mber.

NAME AND ADDRESS OF APPLICANT *(See Instruction 3 B. below)*

Eugene G. Martin, Ph.D.
JMDNJ - Robert Wood Johnson Medical School
University Diagnostic Laboratories
One Robert Wood Johnson Place - MEB 212
New Brunswick, NJ 08901

TELEPHONE NUMBER (Include Area Code)

(732) 235-8110

2. APPLICATION *(Check one box only)*

I hereby apply for a registration number pursuant to 10 CFR 31, Section 31.11, for use of byproduct materials for:

- ☐ Myself, a duly licensed physician authorized to disperse drugs in the practice of medicine.
- ☒ The above-named clinical laboratory.
- ☐ The above named hospital.
- ☐ Veterinarian in the practice of veterinary medicine.

INSTRUCTIONS

Submit this form in duplicate to:

Materials Safety Branch (T-8 F5)
Division of Industrial and Medical Nuclear Safety
Office of Nuclear Material Safety and Safeguards
U.S. Nuclear Regulatory Commission
Washington, DC 20555-0001

(At NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.)

In the box above, print or type the name, address (including ZIP Code), and telephone number of the registrant physician, clinical laboratory, hospital, or veterinarian in the practice of veterinary medicine for whom or for which this registration form is filed.

4. REGISTRATION

REGISTRATION NUMBER:

9205

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Traci Kime

Traci Kime October 5, 2001

(If this an initial registration, leave this space blank -- number to be assigned by NRC. If this is a change of information from a previously registered general license, include your registration number.)

place of use is different from address listed above, give complete address

JMDNJ-Robert Wood Johnson Medical School-Univ. Diagnostic Laboratores - MEB 215, New Brunswick NJ 08901

6. CERTIFICATION

hereby certify that:

A. All information in this registration certificate is true and complete.

B. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.

C. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.

D I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form), and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the U S Nuclear Regulatory Commission

PRINTED OR TYPED NAME AND TITLE OF APPLICANT	SIGNATURE	DATE
Eugene G. Martin, Ph.D.		9/21/01

WARNING: FALSE STATEMENTS IN THIS CERTIFICATE MAY BE SUBJECT TO CIVIL AND/OR CRIMINAL PENALTIES. NRC REGULATIONS REQUIRE THAT SUBMISSIONS TO THE NRC BE COMPLETE AND ACCURATE IN ALL MATERIAL RESPECTS. 18 U.S.C. 1001 MAKES IT A CRIMINAL OFFENSE TO MAKE A WILLFULLY FALSE STATEMENT OR REPRESENTATION TO ANY DEPARTMENT OR AGENCY OF THE UNITED STATES AS TO ANY MATTER WITHIN ITS JURISDICTION.