

## 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, 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/

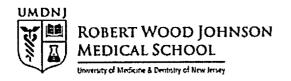
Traci Kime, Licensing Assistant
Materials Safety & Inspection Branch
Division of Industrial
and Medical Nuclear Safety
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OFC	MSIB	
NAME	Traci Kime	
DATE	1/28/2003	

OFFICIAL RECORD COPY



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. of byproduct material that the registrant is entitled to neceive 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 bist @nrc.gov, and to the Dask 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 like Office control without the secondary and a a means used to impute a second of the NRC may not conduct or sponsor, and a person is not required to respond to, the information collection.

was a physicians, clinical laboratories, hospitals, and veterinarians in the		
orizing physicians, clinical laboratories, hospitals, and veterinarians in the of byproduct material for <i>in vitro</i> clinical or laboratory tests not involving or the radiation therefrom to human beings or animals. Possession of e physician, clinical laboratory, hospital, or veterinarian in the practice of the Commission a validated copy of NRC Form 483 with a registration		
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.		
1 —		
Veterinarian in the practice of veterinary medicine.		
4. REGISTRATION		
REGISTRATION NUMBER:		
9205 FOR THE U.S. NUCLEAR REGULATORY COMMISSION		
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.)		
nív. Diagnostic Laboratores - MEB 215, New Brunswi		
RTIFICATION . NJ 00901		
olete.		
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.		
change in the information furnished by a registrant on this registration Safety and Safeguards within 30 days from the effective date of such		
1 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this comply with those provisions as to all byproduct material which he e general license for which this Registration Certificate is filed with the		
comply with those bindiginis as to all papionnel material miner in		

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