NRC FORM 483

U. S. NUCLEAR REGULATORY COMMISSION

EXPIRES: 6-30-99

(9-96)

REGISTRATION CERTIFICATE - in vitro TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE the transfer to the American desirable to

Estimated burden per response to comply with this mandatory information collection request: 7 minutes. The validated registration serves as evidence to suppliers of byproduct material that the registrant is entitled to receive the byproduct meterial. Forward comments regarding burden a formation and Records Management Branch (T-6 F33), U.S. Regulatory Commission, Washington, DC 2055-001, Passacht Reduction Project (3150-0038), Office of Man Reduction Project (3150-0038), Office of Man Reduction Project (3150-0038), Office of Man in its not required to respond to a polic e a currently valid OMB control numb

Section 31:11 of 10 CFR 37 establishes a general license authorizing physicians, clinical laboratories, hospitals, and veterinarians in the practice of veterinary medicine to possess certain small quantities of byproduct material for in vitro clinical or laboratory tests not involving the internal or external administration of the byproduct material or the radiation therefrom to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, hospital, or veteriarian in the practice of veterinary medicine, has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with a registration number.

4 SIEST SUB ADDOCAG OF ADDITION OF
1. NAME AND ADDRESS OF APPLICANT (See Instruction 3.B. below)
B. Rabel Florald MA
B. Rafael Eletalde MD Medical Genetics Institute S.C.
4555 W. Schvoeder Drive, Suite 180
Milwauker, Wi 53223
TELEPHONE NUMBER (Invivid Arm Code)

2. APPLICATION (Check one box only)

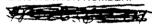
APPROVED BY OMB: NO. 3150-0038

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

- A. 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.

4. REGISTRATION

REGISTRATION NUMBER:



INSTRUCTIONS:

A. Submit this form in duplicate to:

414) 357-6555

Medical, Academic and Commercial Use 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 v. a.

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

(If this an initial registration, leave this seblank – number to be assigned by NRC. If this is a change of marmation from a previously registered general license, include your registration number.)

5. If place of use is different from address listed above, give complete address:

6. CERTIFICATION

I hereby certify that:

- A. All information in this registration certificate is true and complete.
- 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.
- 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 A	ND TITLE OF APPLICANT
E. RAFAEL	ELEJALDE, MD

SIGNATURE OF APPLICANT

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. SECTION 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.