Form AEC-483
1/74
10 CFR 31

U.S. ATOMIC ENERGY COMMISSION

REGISTRATION CERTIFICATE—IN VITRO TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

Form Approved
Budget Bureau No.
38—RO 160

Section 31.11 of 10 CFR 31 establishes a general license authorizing physicians, clinical laboratories, and hospitals 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, or hospital has filed Form AEC-483 and received from the Commission a validated copy of Form AEC-483 with registration number.

Wherever the words "Atomic Energy Commission" or "Commission" appear in this registration, they mean the Nucle Regulatory Commission created by Public Law 93-433 and Executive Order No. 11834.

MEDICAL ACUPUNCTURE RESEARCH
Institute of AMERICA
1034 Stelton Rd.
Piscataway, N.J 08854

DR. LUPO T. CARLOTA, President

INSTRUCTIONS

1. Submit this form in triplicate to:
   Director of Licensing
   ATTN: Materials Branch
   Regulation
   U.S. Atomic Energy Commission
   Washington, D.C. 20545
2. Please print or type the name and address (including zip code) of the registrant physician, clinicial laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At AEC, a registration number will be assigned and a validated copy of Form AEC-483 will be returned.)

3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for *(please check one block only)*
   - ☑ a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
   - ☐ b. The above-named clinical laboratory.
   - ☐ c. The above-named hospital.
4. To be completed by the Atomic Energy Commission

Registration number: 3090

FOR THE U. S. NUCLEAR REGULATORY COMMISSION

BY: Clarence A. Hebron 3/24/75

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

5. If place of use is different from address in Item 1, please give complete address:

6. Certification:

I 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 Licensing, within 30 days from the effective date of such change.

d. I have read and understand the provisions of Section 31.11 of AEC 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 Atomic Energy Commission.

Date 2/25/75

By Lupo T. Carlota, M.D.
Signature of person filing form

Printed name and title or position of person filing form
Dr Lupo T. Carlota, M.D.

WARNING—18 U.S.C., Section 1001; Act of June 25, 1948; 62 Stat. 749; 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.