NRC Form 483
1-76
10 CFR 31

U.S. NUCLEAR REGULATORY COMMISSION

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

RECEIVED JUL 28 1982 U.S. Nuclear Regulatory Commission NMSS Mail Section

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. The use of byproduct material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, or hospital has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

H.P. Moreno-Campos MD
909 Woodward Avenue #101
Pontiac, Michigan 48053

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)*
   - X 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 Nuclear Regulatory Commission.

INSTRUCTIONS

1. Submit this form in triplicate to:
   Office of Nuclear Material Safety and Safeguards
   ATTN: Radioisotopes Licensing Branch
   U.S. Nuclear Regulatory Commission
   Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. (Position the first letter of the address below the right dot and do not extend the address beyond the right dot.) If NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.

Registration number: 6389

FOR THE U. S. NUCLEAR REGULATORY COMMISSION

Shirley A. Crutchfield July 14, 1982

If there is an error in printing, leave that space blank — we will enter it as corrected by NRC. If there is a change of information from a previously registered clinical laboratory, 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 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 the Registration Certificate is filed with the Nuclear Regulatory Commission.

Date _____

By X H Moreno Campos MD

Printed name and title or position of person filing form

WARNING—18 U.S.C., Section 100[illegible] of Title 2[illegible] U.S.C., [illegible] to make a willfully false statement or representation [illegible] to any matter within the jurisdiction [illegible]