ASSOCIATED ENDOCRINOLOGISTS, P.C.

Karen K. Berris, M.D.
Howard S. Blank, M.D.
Gary W. Edelson, M.D.
Michael Garcia, M.D.
Michael M. Kaplan, M.D.
Lowell R. Schmeltz, M.D.
Charles I. Taylor, M.D.
Carla Ferrise, MSN, FNP-BC

Laboratory Director
Walter E. Szpunar, Ph.D.

Date: August 6, 2015

To: Licensing Branch (T8 E24)
Division of Materials Safety and State Agreements
U.S. Nuclear Regulatory Commission
Washington, DC 20555–0001

License No. 21-24444-01

Subject: License renewal with change of license scope.

Enclosed is NRC Form 483 for Associated Endocrinologists, P.C. The clinical laboratory of our office no longer uses liquid iodine–125 for *in vitro* work. We now use only prepackaged kits containing iodine–125 in units not exceeding 10 microcuries for each use. The kits are for *in vitro* human blood tests not involving internal or external administration of byproduct material, or the radiation therefrom, to human beings or animals. We do not and will not possess more than 200 microcuries of iodine–125 at any one time.

It is our understanding that this use of byproduct material qualifies our clinical laboratory for a general license under section 31.11 of 10 CFR 31.

If there are any problems or questions regarding this application, please contact me at telephone # 248-855-5620 or by email kaplan@endocrinemds.com.

Michael M. Kaplan, MD
President and Radiation Safety Officer
Associated Endocrinologists, P.C.

NRC FORM 483
(03-2015)

U.S. NUCLEAR REGULATORY COMMISSION

APPROVED BY OMB: NO. 3150-0038 EXPIRES: 02/28/2018

Estimated burden per response to comply with this mandatory collection request: 8 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 FOIA, Privacy and Information Collections Branch (T-5 F53), U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001, or by internet e-mail to Infocollects.Resource@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.

REGISTRATION CERTIFICATE -- *IN VITRO TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE*

Section 31.11 of 10 CFR 31 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 veterinarian 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.

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

Associated Endocrinologists, P.C.
6900 Orchard Lake Road, Suite 204
West Bloomfield, MI 48322-3425

TELEPHONE NUMBER (Include Area Code):
248-855-5620

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

A. Submit this form to:

Licensing Branch (T-8 E24)
Division of Materials Safety & State Agreements
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.)

B. 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:

(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.)

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

6900 Orchard Lake Road, Suite LL06
West Bloomfield, MI 48322-3425

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 Federal and State Materials and Environmental Management Programs 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 (on page 2 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
Michael M. Kaplan, MD, President and Radiation Safety Officer Associated Endocrinologists, P.C.	Michael M Kaplan MD	8/24/2015

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.