

UNITED STATES NUCLEAR REGULATORY COMMISSION WASHINGTON, D.C. 20555-0001

April 18, 2014

Steven Trombly, MD 5195 15 Mile Road Sterling Heights, MI 48310

Dear Dr. Trombly:

This letter verifies receipt of the completed NRC Form 483 dated April 2, 2014. This form is a condition of the general license under 10 CFR 31.11 authorizing in-vitro testing with byproduct material under a general license.

The form has been assigned registration number **9360.** When making changes to any of the information on the form, please reference the registration number and address the correspondence to the Director, Office of Federal and State Materials and Environmental Management Programs, Washington, DC 20555.

If you have any questions or need further assistance, please contact me at (301) 415-8140.

Sincerely,

Licensing Assistant

Licensing Branch

Division of Materials Safety and State Agreements

Office of Federal and State Materials and Environmental Management Programs

Enclosure: NRC Form 483 - 9360

NRC FORM 483

(2-2012)

U.S. NUCLEAR REGULATORY COMMISSION

REGISTRATION CERTIFICATE -- in vitro TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

APPROVED BY OMB: NO. 3150-0038

EXPIRES: 01/31/201

Estimated burden par response to comply with this mandatory collection request. § minutes. The validated registration serves as evidence to suppliers of byproduci material that the registrant is entitled to receive the byproduct material. Send comments regarding burden estimate to the Information Services Branch (T-5 F53), U.S. Nuclear Regulatory Commission, Weshington, DC 2055-0001, or by internet e-mail to infocollects. Resource@mic.gov, and to the Desk Officer, Office of Information and Regulatory Affairs, NEO6-10202, (3150-0038), Office of Management and Budget Washington, DC 20533. If a means used to timpose an information collection does not display a currently valid CMB control number, the NRC may not conduct or spensor and a person is not required to respond to, the information collection.

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.

with a registration number.		
TELEPHONE NUMBER (Include Area Code):	2. APPLICATION (Check one box only) I hereby apply for a registration number pursuant to 10 CFR 31, Section 31.11, for use of by product materials for: Myself, a duty icensed 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 NUMBER: 9.360 FOR THE U.S. NUMBER: 9.360 (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.)	
INSTRUCTIONS A. Submit this form to: Source Safety and Security Branch (T-8 E24) Division of Materials Safety & State Agreements and Environmental Management Programs 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 filled. 5. If place of use is different from address listed above, give complete address.		
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I hereby certify that:

6. CERTIFICATION

- 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 this Registration Certificate is filed with the U.S. Nuclear Regulatory Commission.

PRINTED OR TYPED NAME AND TITLE OF APPLICANT	SIGNATURE	DATE,
Steven D. Trombly M.D.	1. Krom by	14.2.14

WARNING: FALSE STATEMENTS IN THIS CERTIFICATE MAY BE SUBJECT TO CIVIL ÁNDIOR 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.

NRC FORM 483 (2-2012)

Steve Trombly, M.D., P.C. 5195 Fifteen Mile Road Sterling Heights, MI 48310

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