January 13, 1998

Mr. Gary Ross
Clinical Laboratories, Inc.
901 Keystone Industrial Park
Throop, PA 18512

Dear Mr. Ross:

This letter verifies the receipt of the completed NRC Form 483 dated September 24, 1997. This form is a condition of the general license under 10 CFR 31.11 authorizing in-vitro testing with byproduct material under general license.

The form has been assigned registration number 9126. When making changes to any of the information on the form, please reference the registration number and address the correspondence to Director, Office of Nuclear Material Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington DC 20555.

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

Sincerely,

/S/

KimBerly Randall, Registration Specialist
Sealed Source Safety Section
Source Containment and
Devices Branch
Division of Industrial and
Medical Nuclear Safety, NMSS

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