UNITED STATES
NUCLEAR REGULATORY COMMISSION
REGION I
475 ALLENDALE ROAD
KING OF PRUSSIA, PENNSYLVANIA 19406-1415

October 10, 2008

Docket No. 030-31689 License No. 19-07538-05
Control No. 142818

Robert Parker, Ph.D.
Acting Director
Laboratory of Clinical Pharmacology
Office of Testing and Research
Center of Drug Evaluation and Research
U. S. Food and Drug Administration, DHHS
Life Sciences Building #64, Room 2022
10903 New Hampshire Avenue
Silver Spring, MD 20993

SUBJECT: LABORATORY OF CLINICAL PHARMACOLOGY, LICENSE AMENDMENT, CONTROL NO. 142818

Dear Dr. Parker:

This refers to your license amendment request. Enclosed with this letter is the amended license.

Please review the enclosed document carefully and be sure that you understand and fully implement all the conditions incorporated into the amended license. If there are any errors or questions, please notify the U.S. Nuclear Regulatory Commission, Region I Office, Licensing Assistance Team, (610) 337-5239, so that we can provide appropriate corrections and answers.

An environmental assessment for this action is not required, since this action is categorically excluded under 10 CFR 51.22(c)(14).

Current NRC regulations and guidance are included on the NRC's website at www.nrc.gov; select **Nuclear Materials; Medical, Academic, and Industrial Uses of Nuclear Material;** then **Regulations, Guidance, and Communications.** You may also obtain these documents by contacting the Government Printing Office (GPO) toll-free at 1-888-293-6498. The GPO is open from 7:00 a.m. to 8:00 p.m. EST, Monday through Friday (except Federal holidays).

Thank you for your cooperation.

Sincerely,

Original signed by Elizabeth Ullrich

Betsy Ullrich
Senior Health Physicist
Commercial and R&D Branch
Division of Nuclear Materials Safety

Enclosure
Amendment No. 10

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SUNSI Review Complete: EUllrich
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