



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

December 21, 2017

Dianon Pathology
ATTN: Jeremy Ginn, MT
Clinical Lab Manager
840 Research Parkway
Oklahoma City, OK 73104

SUBJECT: TERMINATION OF REGISTRATION CERTIFICATE – *IN VITRO* TESTING
WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE NO. 8929

Mr. Ginn:

This letter is in response to your letter dated November 9, 2017, requesting an update to your Registration Certificate No. 8929. The purpose of this letter is to inform you of the termination of Dianon Pathology General License No. 8929, authorizing the use of by product material for *in vitro* testing in accordance with Title 10 of the *Code of Federal Regulations* (10 CFR) 31.11, "General license for use of byproduct material for certain *in vitro* clinical or laboratory testing." The U.S. Nuclear Regulatory Commission (NRC) is terminating your general license because the address of use of the licensed material is not under NRC jurisdiction. You should contact the State of Oklahoma and comply with their regulations.

In accordance with 10 CFR 2.390 a copy of this letter will be available electronically for public inspection in NRC's Public Document Room or from the Publicly Available Records component of NRC's document system (ADAMS). ADAMS is accessible from the NRC Web site at <http://www.nrc.gov/NRC/ADAMS/index.html> (the Public Electronic Reading Room).

If you have any questions, please contact Katie Wagner, the General License Tracking System Program Manager, at (301) 415-6202 or by electronic mail at Katie.Wagner@nrc.gov.

Sincerely,

/RA/

Celimar Valentin-Rodriguez, Ph.D.
Materials Safety Licensing Branch
Division of Material Safety, State, Tribal
and Rulemaking Programs
Office of Nuclear Material Safety
and Safeguards

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DATE	12/21/2017	12/21/2017	12/21/2017

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