UNITED STATES
NUCLEAR REGULATORY COMMISSION
REGION I
2100 RENAISSANCE BOULEVARD, SUITE 100
KING OF PRUSSIA, PA 19406-2713

July 29, 2015

Docket No. 03030793 License No. 54-28275-02MD
Control No. 588169

Jackie Kavanagh
Senior Manager
Nordion (Canada) Inc.
447 March Road
Ottawa, ON K2K 1X8
Canada

SUBJECT: NORDION (CANADA) INC., LICENSE AMENDMENT, CONTROL NO. 588169

Dear Ms. Kavanagh:

This refers to your license amendment request dated June 1, 2015. Enclosed with this letter is the amended license removing the license condition that required submission of evidence of your registration or licensure with the U.S. Food and Drug Administration as a drug manufacturer.

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; Med, Ind, & Academic Uses;** then **Licensee Toolkits, see our toolkit index page.** You may also obtain these documents by contacting the Government Printing Office (GPO) toll-free at 1-866-512-1800. The GPO is open from 8:00 a.m. to 5:30 p.m. EST, Monday through Friday (except Federal holidays).

Thank you for your cooperation.

Sincerely,

Original signed by Penny Lanzisera

Penny Lanzisera
Senior Health Physicist
Medical Branch
Division of Nuclear Materials Safety

Enclosure:

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SUNSI Review Complete: RElliott

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