



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

July 20, 2016

Docket No. 03011936
Control No. 590661

License No. 10-12044-03

Michael A. Weber
COL, U.S. Army
Commanding
Department of the Army
Dwight David Eisenhower Army Medical Center
Attn: MCHF-PMS-HP
300 Hospital Road
Fort Gordon, GA 30905-5650

**SUBJECT: DEPARTMENT OF THE ARMY, DWIGHT DAVID EISENHOWER ARMY
MEDICAL CENTER, LICENSE AMENDMENT, CONTROL NO. 590661**

Dear COL Weber:

This refers to your license amendment request dated March 17, 2016. Enclosed with this letter is the amended license changing the radiation safety officer from MAJ Douglas Barrickman to Paul A. Gallagher and adding the use of Iodine-125 low dose rate brachytherapy seeds (Best Medical International, Inc. Model 2301) for the localization of non-palpable lesions. Please note that your Radiation Safety Committee can approve the use, authorized user(s), and your facilities under your broad scope license. Therefore, your procedures were not reviewed, and your letter and its attachments were not tied down under License Condition 22.

Please note that you must meet the requirements for temporary implants and develop, implement, and maintain the appropriate procedures in the following regulations: 10 CFR 35.40(a), (b)(6), (c), and (d), 35.41, 35.67, 35.75, 35.310, 35.404, 35.406, 35.410, and 35.432. You must maintain records for seed localization in accordance with the requirements for temporary implants to include the following: 10 CFR 35.2024, 35.2026, 35.2041, 35.2060, 35.2067, 35.2075, 35.2310, 35.2404, 35.2406, and 35.2432. You are also required to report any medical event, except for those that result from patient intervention, in accordance with 10 CFR 35, Subpart M, to include: 10 CFR 35.3045, 35.3047, and 35.3067.

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).

M. Weber

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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 James P. Dwyer

James Dwyer
Chief, Medical Branch
Division of Nuclear Materials Safety
Region I

Enclosure:
Amendment No. 46

cc:
Paul A. Gallager, Radiation Safety Officer

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SUNSI Review Complete: RGallagher

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NAME	RGallagher/RLG		JDwyer/JDwyer					
DATE	7/20/2016		7/20/2016					

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