September 19, 2001

Stanley L. Mills, Ph.D., R.Ph.
President and CEO
Mills Biopharmaceuticals, Inc.
120 NE 26th St.
Oklahoma City, OK 73105

SUBJECT: COMPLETION OF REGISTRATION CERTIFICATE FOR MODELS I-125 AND Pd-103 BRACHYTHERAPY SOURCES

Dear Dr. Mills:

Based on the information and test data submitted in your application dated April 30, 2001, and subsequent correspondence with enclosures thereto, we continue to conclude that the Models I-125 and Pd-103 brachytherapy sources are acceptable for licensing purposes in accordance with the conditions of the enclosed registration certificate (NR-1081-S-101-S).

Please be advised that you must distribute the product in accordance with the statements and representations contained in your application, with enclosures thereto, and the information set out in your registration certificate. As a general rule, you must request and obtain an amendment to the certificate before you make changes or modifications to the information submitted to obtain the certificate.

Please read over the enclosed registration certificate in its entirety and notify us immediately of any errors or omissions.

You are obligated to notify us promptly in writing should you decide to no longer manufacture or offer service support for the product.

Please be aware that, as a holder of an NRC registration, you may be subject to the NRC's licensing and inspection fees in accordance with 10 CFR Part 170, and annual fees in accordance with 10 CFR Part 171. If you have any questions concerning the fee requirements, please contact the License Fee and Accounts Receivable Branch at (301) 415-7544.

If you have any questions, please contact me at (301) 415-7904 or Dr. Seung J. Lee at (301) 415-5787.

Sincerely,
/RA/
John P. Jankovich, Ph.D., Sr. Engineer
Materials Safety and Inspection Branch
Division of Industrial and Medical Nuclear Safety
Office of Nuclear Material Safety and Safeguards

Enclosure: As stated
cc w/encl: S. Crutchfield, LFARB
Enclosure: As stated

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