

Medical-Surgical Division/3M

3M Center
St. Paul, Minnesota 55144-1000
612/733-1110

3M

October 1, 1987

Bruce S. Mallet, Ph.D.
Chief, Materials Licensing
U.S. Nuclear Regulatory Commission
799 Roosevelt Road
Glen Ellyn, Illinois 60137

Dear Dr. Mallet:

This letter is a follow-up to my letters of March 16, 1987 and June 11, 1987, regarding revisions to labeling materials for I-125 Seeds.

Enclosed with this letter are three package inserts for I-125 Seeds, one each for models 6702, 6711, and 6720. In accordance with our previous agreements, these inserts have been revised to provide additional precautionary information to customers, specifically pertaining to reuse of I-125 Seeds. These revised inserts are included with each Seed shipment beginning October 1, 1987.

These inserts represent the final aspect in our process of labeling revision for I-125 Seeds, and we trust that we have satisfied the agency's concerns with regard to providing adequate instructions to customers who reuse the product. If you have any questions regarding this matter, please feel free to call (612/733-6421).

Sincerely yours,

Jacquelyn D. Bush
Jacquelyn D. Bush
Sr. Regulatory Affairs Specialist
3M Medical-Surgical Division
3M Center, Building 270-4A-05
St. Paul, MN 55144-1000

clrk

CONTROL NO 84259

9105220210 B80108
REG3 LIC30
22-00057-59MD PDR

*refee due
See 1/88
memo to
J. Jackson*

RECEIVED

OCT 05 1987

REGION III

OCT 5 1987