# E.I. DU PORT DE NEMOURS \& CO. (INC.) MEDICAL PRODUCTS DEPARTMENT 

December 7, 1990

United States Nuclear Regulatory Commission
Attn: Charles E. MacDonald, Chief
Transportation Branch
Division of Safeguards and Transportation, NMSS
Washington, D.C. 20555

Reference: QA Program Approval Number 0384

Dear Mr. MacDonald:
This is a request for an amendment to the the above-referenced Quality Assurance Program Approval for Radioactive Material Packages to reflect a new business name.

On January 1, 1991, a new company will be officially formed by a Joint Venture between the two major corporations, Du Pent and Merck. The name of our company will be changing from Du Font to the Du Font Merck Pharmaceutical Company,

Therefore, effective January 1, 1991, the name and address of our company should be listed on the approval as:

Du Font Merck Pharmaceutical Company<br>33 ! Treble Cove Road<br>North Billerica, Massachusetts 01862

Due to this change in the company name, a revised version of the Quality Assurance Program has been written and is enclosed for your review. Once approved, the scope of the revised program will cover only the operations at the Billerica address. A request for a QA Program Approval for the operations at the Du Pont facility at 549 Albany Street, Boston MA, previously a part of approval number 0384, will be submitted to you under separate cover.

I have enclosed a check in the amozat of $\$ 180.00$ in payment of the amendment processing fee as specified for License Category 10B in the regulations of Title 10 CFR Part 170, Section 170.31.

Please contact me if you require any additional information.

Sincerely,


Francis E. Roy Jr.


Health Physicist
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MEDICAL PRODUCTS DEPARTMENT
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