DEC 10 1985

Docket No. : 030-13158
License No.: 50-17686-01
Control No.: 70299

Bartlett Memorial Hospital
3260 Hospital Drive
Juneau, Alaska 99801

Attention: Mr. James R. Burns
Administrator

Gentlemen:

This is in reference to your application dated November 25, 1985 to amend your byproduct material license. In order to complete our review, we need the following additional information:

1. You should submit documentation that Dr. Greening has completed 200 hours of didactic training as described in Item 2.a. of Appendix A to Regulatory Guide 8.20. Although the Supplement A which was submitted with your letter dated November 25, 1985 does indicate that Dr. Greening received training at the Hospital of the University of Pennsylvania, it does not reflect the number of hours of didactic training received in each of the five fields of training indicated on Supplement A.

2. Although the NRC-313M Supplement B lists three hospitals where Dr. Greening received training and experience, only one preceptor signature is given. Separate preceptor statements should be submitted from each institution. These statements should indicate the extent of Dr. Greening's involvement with the nuclear medicine programs. Refer to Item 2.c. of Appendix A to Regulatory Guide 10.8.

   If it is not possible to obtain preceptor statements from physicians who supervised Dr. Greening during his internship, you should ask that each of the three hospitals submit a signed statement to document:

   a. the time interval when Dr. Greening was present at the facility.

   b. his involvement with the nuclear medicine program.

   c. the extent of his involvement (patient selection, dose determination, scan interpretation, patient followup, study of possible complications, etc.)

   d. any authorization granted to Dr. Greening by the hospital's Radiation Safety Committee.

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3. In order to add Dr. Greening as an authorized user for Group III, we must have documentation to verify that he has personally participated in five procedures to elute technetium-99m, including testing of elute, and in five procedures to prepare radiopharmaceuticals from Group III reagent kits. You should submit this documentation. Refer to Item 2b of Appendix A to Regulatory Guide 10.8.

We will continue the review of your amendment request upon receipt of this information. If we do not receive a reply from you within 30 calendar days from the date of this letter, we shall assume that you do not wish to pursue your application. Please reply in duplicate, and refer to Mail Control No. 70299.

Sincerely,

Beth A. Riedlinger
Health Physicist (Licensing)
Nuclear Materials Safety Section

Enclosures:
Appendix A to Regulatory Guide 10.8
Federal Register Notice dated December 2, 1982 (FR 54376)
NRC Form 313M