



UNITED STATES  
**NUCLEAR REGULATORY COMMISSION**  
REGION I  
475 ALLENDALE ROAD  
KING OF PRUSSIA, PENNSYLVANIA 19406-1415

December 31, 2008

Docket No. 03036417  
Control No. 143093

License No. 29-30840-01

Ralph DeBellonia  
Administrator  
PET Scan of New Jersey  
315 Elmora Avenue, Suite 208  
Elizabeth, NJ 07208

SUBJECT: PET SCAN OF NEW JERSEY, REQUEST FOR ADDITIONAL INFORMATION  
CONCERNING APPLICATION FOR AMENDMENT TO LICENSE, CONTROL  
NO. 143093

Dear Mr. DeBellonia:

This is in reference to your letter dated December 8, 2008 requesting to amend Nuclear Regulatory Commission License No. 29-30840-01. In order to continue our review, we need the following additional information:

1. You requested for the addition of oral administration of sodium iodide, Iodine 131. However, you did not request for its authorized use. Please state your use for iodine 131 (e.g. diagnostic use, therapy for hyperthyroidism, therapy for thyroid ablation, or other uses).
2. You have requested Allen C. Pomerantz, M.D. to be your radiation safety officer. Please submit Dr. Pomerantz's training and qualification to be the radiation safety officer. Enclosed is the recommended NRC Form 313A (RSO) form for documenting and attesting to his qualification. If Dr. Pomerantz was a radiation safety officer previously, please include the NRC license number or a copy of the agreement state license listing him as the radiation safety officer.
3. NUREG-1556, Volume 9, Revision 2, Appendix S, address that a review of administrations requiring a written directive be included in your Quality Management Program. Your QMP did not address this item. Please include a section addressing the review of administrations requiring a written directive in your QMP. It is recommended that you review NUREG-1556, Volume 9, Revision 2, Appendix S for guidance.
4. You did not address the report of medical events to the NRC in your QMP. Please commit to: Notify by telephone the NRC Operations Center no later than the next calendar day after discovery of a medical event and submit a written report to the appropriate NRC Regional Office listed in 10 CFR 30.6 within 15 days after the discovery of the medical event, as required by 10 CFR 35.3045. Also notify the referring physician and the patient as required by 10 CFR 35.3045.

Current NRC regulations and guidance are included on the NRC's website at [www.nrc.gov](http://www.nrc.gov); select **Nuclear Materials; Medical, Academic, and Industrial Uses of Nuclear Material**; then **Regulations, Guidance, and Communications**. 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 7:00 a.m. to 6:30 p.m. EST, Monday through Friday (except Federal holidays).

We will continue our review upon receipt of this information. Please reply to my attention at the Region I Office and refer to Mail Control No. 143093. If you have any technical questions regarding this deficiency letter, please call me at (610) 337-5366.

If we do not receive a reply from you within 30 calendar days from the date of this letter, we will assume that you do not wish to pursue your application.

Sincerely,

***Original signed by Dennis R. Lawyer***

Dennis R. Lawyer  
Health Physicist  
Commercial and R&D Branch  
Division of Nuclear Materials Safety

Enclosure:  
NRC Form 313A (RSO)

cc:  
Natalio Damien, M.D., Radiation Safety Officer

DOCUMENT NAME: G:\WordDocs\Current\Lic Def Letter\L29-30840-01.143093.doc

**SUNSI Review Complete: DLawyer**

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| OFFICE | DNMS/RI       | N | DNMS/RI |  | DNMS/RI |  |  |  |
| NAME   | DLawyer /DC9/ |   |         |  |         |  |  |  |
| DATE   | 12/31/2008    |   |         |  |         |  |  |  |

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