

From: Michelle Simmons
To: Dmhppphysics@aol.com
Date: Wed, May 4, 2005 1:09 PM
Subject: City Hospital: Request for additional information
Place: LAT

License No. 47-15501-01
Docket No. 03009218
Control No. 136135

Dear Sir:

This is in reference to your application dated December 15, 2004 requesting to renew Nuclear Regulatory Commission License No. 47-15501-01. In order to continue our review, we need the following additional information.

1. Confirm zip code and mailing address (i.e. Dry run or Dry Run & Tavern)
2. Provide licensed limit for 31.11 material (i.e. 5m Ci)
3. Describe areas above/below/behind hot lab and indicate dose rates in these areas.
4. Procedures not reviewed in detail, however cursory review indicates that procedures may not be consistent with currently accepted standards (i.e. ANSI- N323A-1997 survey instrumentation calibration). However, detailed procedures are no longer required to be submitted. Instead you may provide the following statements:
 - A. A statement that: "Radiation monitoring instruments will be calibrated by a person qualified to perform survey meter calibrations."
 - B. A statement that: "Equipment used to measure dosages will be calibrated in accordance with nationally recognized standards or the manufacturer's instructions."
 - C. A statement that: "Either we will perform a prospective evaluation demonstrating that unmonitored individuals are not likely to receive, in one year, a radiation dose in excess of 10% of the allowable limits in 10 CFR Part 20 or we will provide dosimetry that meets the requirements listed under "Criteria" in NUREG 1556, Vol. 9, "Consolidated Guidance About Materials Licenses Program-Specific Guidance About Medical Use Licensees," dated October 2002."
 - D. A statement that: "We have developed and will implement and maintain written procedures for area surveys in accordance with 10 CFR 20.1101 that meet the requirements of 10 CFR 20.1501 and 10 CFR 35.70."
 - E. A statement that: "We have developed and will implement and maintain procedures for safe use of unsealed byproduct material that meet the requirements of 10 CFR 20.1101 and 10 CFR 20.1301. "
 - F. A statement that: "We have developed and will implement and maintain written procedures for safe response to spills of licensed material in accordance with 10 CFR 20.1101."

G. A statement that: "We have developed and will implement and maintain written waste disposal procedures for licensed material in accordance with 10 CFR 20.1101, that also meet the requirements of the applicable section of Subpart K to 10 CFR Part 20 and 10 CFR 35.92."

In order to continue our review, we request that you submit a response as soon as possible.

Sincerely,
Michelle Simmons
Health Physicist
Medical Branch
Division of Nuclear Materials Safety

CC: LAT

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(427901E0.AF5 : 23 : 10032)

Subject: City Hospital: Request for additional information
Creation Date: Wed, May 4, 2005 1:09 PM
From: Michelle Simmons

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MESSAGE	6831	Wednesday, May 4, 2005 1:09 PM

Options

Expiration Date: None
Priority: Standard
Reply Requested: No
Return Notification: None

Concealed Subject: No
Security: Standard