



Technologist reminded the technologists that they must be always vigilant in seeing that the correct syringe is used for the correct patient.

This misadministration was reported by phone to me on 11/18/87. While filing a copy of the report of it in her records on 11/18/87 the Chief Technologist discovered the original and copy of a 8/10/87 letter to me in her files. She immediately forwarded the original to me with a note asking if this meant that the misadministration was not reported to me. The misadministration was not reported to me. I am now reporting it to you. I am clear that this was an oversight as she has reported misadministrations to me prior and after this one promptly and willingly. We have discussed the matter and she will make sure that in future I am informed by phone as well as by mail. Details of this misadministration and action taken are in the attached letter to me dated 8/10/87. The referring physician was Dr R. Robinson.

Names of individuals involved in both events are on file in my office.

Sincerely,

M. Rosemary Kennedy  
Radiation Safety Officer

c.c. Gerald M. Kolodny M.D., Chairman, Radiation Safety  
Committee  
Malcolm Weiner, Associate Vice President

REGION I  
NMSS LICENSEE EVENT REPORT

License No. 20-00747-1E

Docket No. 030-09062

MLER-RI-87- 198

I. ACTION CONTROL DATA

Licensee BETH ISRAEL HOSPITAL

Event Description two (2) diagnostic misadministration

Event Date 11/17/87

Report Date 11/20/87

II. REPORTING REQUIREMENT

- |   |   |
|---|---|
| <input type="checkbox"/> 10 CFR 20.402 - theft or loss            | <input type="checkbox"/> 10 CFR 35.33 Therapeutic Misadministration           |
| <input type="checkbox"/> 10 CFR 20.403(a)(b) overexposure/release | <input checked="" type="checkbox"/> 10 CFR 35.33 Diagnostic Misadministration |
| <input type="checkbox"/> 10 CFR 20.405 - 30 day report            | <input type="checkbox"/> License Condition                                    |
| <input type="checkbox"/> Other _____                              |   |

III. REGION I RESPONSE

- |  |                            |
|--|----------------------------|
| <input type="checkbox"/> Immediate Site Inspection | Inspector _____ Date _____ |
| <input type="checkbox"/> Special Inspection        | Inspector _____ Date _____ |
| <input type="checkbox"/> Telephone Inquiry         | Inspector _____ Date _____ |

Licensee Representative and Title \_\_\_\_\_

- PM       Daily Report
- Information entered - Region I log and Outstanding Items List
- Review at next routine inspection

IV. REPORT EVALUATION

- |  |   |
|--|---|
| <input checked="" type="checkbox"/> Description of Event   | <input checked="" type="checkbox"/> Corrective Actions                        |
| <input checked="" type="checkbox"/> Levels of R/M involved | <input checked="" type="checkbox"/> Calculation Adequate                      |
| <input checked="" type="checkbox"/> Cause of Event         | <input type="checkbox"/> Letter to Licensee requesting additional information |

Completed by: E. Ullrich

Date 12/7/87

Reviewed by: D. Kimmey

Date 12/14/87

V. SPECIAL INSTRUCTIONS OR COMMENTS

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