

DIAGNOSTIC MISADMINISTRATION REPORT

IN1: LICENSEE NAME **Brigham and Women's Hospital** IN2: LICENSE NUMBER **20-17131-13**

IN3: CITY **Boston** IN4: STATE **MA** IN5: EVENT DATE **12/16/88** IN6: REPORT DATE **12/16/88**

IN7: TYPE OF MISADMINISTRATION
 101: WRONG RADIOPHARMACEUTICAL
 102: DOSAGE DIFFERING FROM PRESCRIBED BY 50%
 103: WRONG PATIENT
 104: WRONG ROUTE
 IN8: DID THE MISADMINISTRATION INVOLVE AN ISOTOPE OF IODINE
 1099: YES 1111: NO
 IN9: NUMBER OF PATIENTS WHO RECEIVED A MISADMINISTRATION UNDER THIS REPORT **one(1)**

IN10: INTENDED		IN10A: INTENDED				IN11: GIVEN			
105: NO CLINICAL PROCEDURE	106: NUCLEAR MEDICINE STUDY (Comply with IN10A: INTENDED and IN11: GIVEN)	MILLICURIES	ISOTOPE	CHEMICAL FORM	STUDY	MILLICURIES	ISOTOPE	CHEMICAL FORM	STUDY
<input checked="" type="checkbox"/>	107: X RAY STUDY	25 mCi	Tc-99m	MDP	bone	25 mCi	Tc-99m	TcO ₄ ⁻	N/A
<input type="checkbox"/>	108: ULTRASOUND STUDY								
<input type="checkbox"/>	109: CT STUDY								
<input type="checkbox"/>	110: NMR STUDY								
<input type="checkbox"/>	111: OTHER								

IN12: PRECIPITATOR
 171: REFERRING PHYSICIAN
 172: WARD NURSE
 173: WARD CLERK
 174: NUCLEAR PHARMACY
 175: AUTHORIZED USER
 176: HOT LAB TECHNOLOGIST
 177: IMAGING TECHNOLOGIST
 178: CLINIC RECEPTIONIST
 179: SCHEDULING TECHNOLOGIST
 180: PATIENT
 181: OTHER

IN13: ERROR

HOT LAB	REFERRAL	ADMINISTRATION	OTHER
<input type="checkbox"/> 111: MISLABELED A SYRINGE <input type="checkbox"/> 112: MISLABELED A VIAL OR VIAL SHIELD <input type="checkbox"/> 113: RECONSTITUTED WRONG REAGENT KIT <input type="checkbox"/> 114: PLACED RECONSTITUTED VIAL IN WRONG SHIELD	<input checked="" type="checkbox"/> 115: SELECTED WRONG VIAL WHEN DRAWING DOSAGE <input type="checkbox"/> 116: SET DOSE CALIBRATOR IMPROPERLY <input type="checkbox"/> 117: MISREAD DOSE CALIBRATOR <input type="checkbox"/> 118: MISUNDERSTOOD RADIOPHARMACEUTICAL OR DOSAGE ORDER	<input type="checkbox"/> 120: MISUNDERSTOOD REFERRING PHYSICIAN'S REQUEST <input type="checkbox"/> 121: REQUESTED WRONG STUDY <input type="checkbox"/> 122: REQUESTED STUDY FOR WRONG PATIENT	<input type="checkbox"/> 130: SELECTED WRONG PATIENT <input type="checkbox"/> 131: ANSWERED WAITING ROOM PAGE INTENDED FOR OTHER PATIENT <input type="checkbox"/> 132: BROUGHT WRONG PATIENT TO CLINIC <input type="checkbox"/> 133: SELECTED WRONG SYRINGE FROM DOSAGE CART <input type="checkbox"/> 140: Specify _____

IN14: CONTRIBUTING FACTORS
 180: STUDENT TECHNOLOGIST
 181: NEW EMPLOYEE
 182: FOREIGN LANGUAGE
 183: PATIENT INCOHERENT OR UNCONSCIOUS
 184: ID BRACELET NOT CHECKED
 185: REQUISITION NOT CHECKED
 186: PATIENT CHART NOT CHECKED
 187: NEW PROCEDURE
 188: HEAVY WORKLOAD
 189: OTHER _____
 IN15: ACTION TAKEN TO PREVENT RECURRENCE
 1C0: IMPROVE SUPERVISION OF PERSONNEL
 1C1: NO ACTION
 1C2: IMPROVE HANDLING
 1C3: VERIFICATION OF PATIENT IDENTIFICATION
 1C4: REINSTRUCT PERSONNEL
 1C5: REPRIMAND PERSONNEL
 1C6: OTHER _____

IN16: EFFECT ON PATIENTS NONE APPARENT SEE ABSTRACT

IN17: ABSTRACT (If more space is required, attach additional sheets.)
 A 40 year old woman with back pain was referred for a bone scan to obtain clarification after two sets of spine radiographs were inconclusive. The nuclear medicine technologist withdrew and administered 25 mCi from a multidose vial labeled "TcO₄⁻" instead of "MDP".
 License No. 20-17131-03
 Docket No. 030-15070
 MLEP-8 9-002

RADIATION OFFICER (Printed Name) **David E. Drum, M.D.** SIGNATURE *David E. Drum* TELEPHONE **(617) 732-6056** DATE **12/29/88**

NUCLEAR REGULATORY COMMISSION USE
 IN18: 1999: YES 1111: NO
 IN19: AS
 IN20: REGIONAL LOG NUMBER
 IN21: ACCESSION NUMBER
 IN22: INITIALS
Return Original to Region I

1X30

REGION I
NMSS LICENSEE EVENT REPORT

License No. 20-17131-03

Docket No. 030-15070

MLER-RI-89 - 002

I. ACTION CONTROL DATA

Licensee BRIGHAM AND WOMEN'S HOSPITAL

Event Description diagnostic misadministration

Event Date 12/16/88

Report Date 12/16/88

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 type="checkbox"/> Calculation Adequate |
| <input checked="" type="checkbox"/> Cause of Event | <input type="checkbox"/> Letter to Licensee requesting additional information |

Completed by: [Signature]

Date 3/1/89

Reviewed by: [Signature]

Date 3/9/89

V. SPECIAL INSTRUCTIONS OR COMMENTS

F/11