

SAFETY INSPECTION

97-001

Page 1 of _____

1. LICENSEE

Department of the Army
 Walter Reed Army Medical Center
 Washington, D.C. 20307-5001

2. REGIONAL OFFICE

REGION I
 U S NUCLEAR REGULATORY COMMISSION
 475 ALLENDALE ROAD
 KING OF PRUSSIA PA 19406-1415

3. DOCKET NUMBER(S)

030-01317

4. LICENSE NUMBER(S)

08-01738-02

5. DATE OF INSPECTION

February 11-14, 1997

LICENSEE:

The inspection was an examination of the activities conducted under your license as they relate to radiation safety and to compliance with the Nuclear Regulatory Commission (NRC) rules and regulations and the conditions of your license. The inspection consisted of selective examinations of procedures and representative records, interviews with personnel, and observations by the inspector. The findings as a result of this inspection are as follows:

☐ 1. Within the scope of this inspection, no violations were observed.

☒ 2. The inspector also verified the steps you have taken to correct the violations identified during the last inspection. We have no further questions on those actions at this time.

☒ 3. During this inspection certain of your activities, as described below or attached, were in violation of NRC requirements. This form is a **NOTICE OF VIOLATION**, which is required to be posted in accordance with 10 CFR 19.11.

☐ A. 9705050433 970214
PDR ADOCK 03001317
C PDR was not properly posted to
 indicate the presence of a _____ 10 CFR 20.203(b),(c),(d),(e) or 34.42.

☐ B. _____ of sealed sources were not
 performed at the proper frequencies. 10 CFR _____ or License Condition Number _____

☐ C. Records of _____ were not properly maintained.
 10 CFR _____ or License Condition Number _____

☐ D. Documents were not properly posted or otherwise made available: 10 CFR 19.11

☒ E. Bioassays Certified by: M. A. Linton **DESIGNATED ORIGINAL** RETURN ORIGINAL TO **IE 07**
Reports or notification of the Nuclear Pharmacist were not made in accordance with
 10 CFR 35.315 or License Condition Number _____

☒ F. 19.12 Failure to provide 19.12 instruction to the nuclear
pharmacist in the regulatory requirement to measure
his thyroid burden after preparation of iodine-123 therapy
does in accordance with 10 CFR 35.315(a)(8).

050009

I hereby state that, within 30 days, the actions described by me to the Inspector will be taken to correct the violations identified in the items checked above. This statement of corrective actions is made in accordance with the requirements of 10 CFR 2.201. No further response will be submitted unless required by the NRC.

SIGNATURE - LICENSEE

DATE

SIGNATURE - NRC INSPECTOR

DATE

Michael J. Linton

14 Feb 97

Merissa Hall Darda

2/14/97

