## Appendix

## NOTICE OF VIOLATION

United Hospital, Inc.

License No. 22-01914-02

As a result of the inspection conducted on December 2, 1981, and in accordance with the Irterim Enforcement Policy, 45 FR 66754 (October 7, 1980), the following violations were identified:

 License Condition 19 states that the licensee shall possess and use licensed material in accordance with statements, representations, and procedures listed in application dated January 31, 1979 and referenced letters.

This application states that quality control checks on your dose calibrator will include daily constancy, quarterly linearity and annual accuracy tests.

Contrary to the above, none of the referenced tests were performed prior to September 1981.

This is a Severity Level V violation (Supplement VII).

 License Condition 19 states that the licensee shall possess and use licensed material in accordance with statements, representations, and procedures listed in application dated January 31, 1979 and referenced letters.

This application states that your survey program will include weekly wipes for removable contamination.

Contrary to the above, no wipe surveys were performed between May 6 and June 5, 1981; June 20 and July 31, 1981; nor during several other periods.

This is a Severity Level V violation (Supplement VII).

 License Condition 19 states that the licensee shall possess and use licensed material in accordance with statements, representations, and procedures listed in application dated January 31, 1979 and referenced letters. This application states that your Medical Isotopes Committee will meet not less than once in each calendar quarter.

Contrary to the above, your Committee did not meet during the second or third quarters of 1981.

This is a Severity Level V violation (Supplement VII).

With respect to item 1, the inspection showed that action had been taken to correct the identified item of noncompliance and to prevent recurrence. Consequently, no reply to this item of noncompliance is required and we have no further questions regarding this matter. With respect to items 2 and 3, pursuant to the provisions of 10 CFR 2.201 you are required to submit to this office within thirty days of the date of this Notice a written statement or explanation in reply, including for each item of noncompliance: (1) corrective action taken and the results achieved; (2) corrective action to be taken to avoid further noncompliance; and (3) the date when full compliance will be achieved. Under the authority of Section 182 of the Atomic Energy Act of 1954, as amended, this response shall be submitted under oath or affirmation. Consideration may be given to extending your response time f r good cause shown.

Dated /2-11-8/

D. G. Wiedeman, Acting Chief Materials Radiation Protection

Section 1