uranium before they received it (personal communication, 1975). On this basis, we suggest that the following nuclear reactions are responsible for the ^{239}Np :

$$^{238}U + ^{1}n \rightarrow ^{239}U$$

(thermal-neutron cross section = 2.7 barns)

$$^{239}\text{U} \rightarrow ^{239}\text{Np} + \beta$$

$$(T_{1/2} = 23 \text{ min}).$$

Molybdenum-98 has a thermal-neutron cross section of 0.14 barns. If we ignore the difference between the half-lives of ⁹⁹Mo (66.6 hr) and ²³⁹Np (56.5 hr), we calculate that a contamination level of approximately 40 μ Ci of ²³⁹Np per curie of ⁹⁹Mo would be expected from the 2 ppm of uranium contamination in the ⁹⁸Mo. This is about 25 times the maximum amount estimated for the first elution.

CONCLUSIONS

Iodine-131, ruthenium-103, and neptunium-239 were found in the eluates of ⁹⁹Mo-^{99m}Tc generators loaded with ⁹⁹Mo manufactured by thermal-neutron irradiation of ⁹⁸Mo. The level of contamination did not exceed maximum permissible levels (10) at any time, although on several occasions the contamination would have approached the maximum permissible levels less than 12 hr after elution.

Unfortunately, ¹³¹I and ²³⁹Np are not detected with adequate sensitivity in routine ⁹⁹Mo breakthrough tests. Detecting 6 nCi of ¹³¹I per millicurie of ^{99m}Tc is difficult with equipment normally available in a nuclear medicine laboratory, and detecting 10 nCi of ²³⁹Np per millicurie of ^{99m}Tc is completely impossible. Therefore, the control of these impurities must rest with the manufacturer. The manufacturer should ensure that the parent ⁹⁹Mo does not contain enough impurities to yield a product that is unacceptable. On the other hand, "after the fact" testing is possible for any nuclear medicine laboratory and should be carried out as a routine procedure so that the quality of the product is well established on a continuing basis.

REFERENCES

1. ROBINSON GD: Impurities in ^{90m}Tc sodium pertechnetate produced by methyl-ethyl-ketone extraction. J Nucl Med 13: 318-320, 1972

2. AMWAR M, LATHROP K, ROSSKELLY D, et al.: Pertechnetate production from "Mo by liquid-liquid extraction. J Nucl Med 9: 298-299, 1968

3. BILLINGHURST MW, GROOTHEDDE M, PALSER R: Radiochemical purity of ^{60m}Tc pertechnetate. J Nucl Med 15: 266-269, 1974

4. CROSBY EH: Radiochemical purity of short lived technetium-99m from commercial suppliers. *Radiology* 93: 435-439, 1969

5. WOOD DE, BOWEN BM: ⁶⁶Zr and ¹²⁴Sb in ⁶⁶Mo-^{66m}Tc generators. J Nucl Med 12: 307-309, 1971

6. BARRALL RC: Reduction of radioactive impurities in radiopharmaceuticals. J Nucl Med 13: 570, 1972

7. PODOLAK EM: ¹³¹Cs, ³⁶Rb and ⁴⁰Co in ³⁰Mo-⁹⁰mTc generator eluate. J Nucl Med 13: 388-390, 1972

8. SMITH EM: Properties, uses, radiochemical purity and calibration of Tc^{som}. J Nucl Med 5: 871-882, 1964

9. HEATH RK: Table of isotopes. In Handbook of Chemistry and Physics, 4th ed. Cleveland, Ohio, CRC Press, 1973– 1974, pp B250–B544

10. Information Letter No. 417: Health Protection Branch, Department of National Health and Welfare, Canada

11. General Electric Co.: Radiopurity analysis of General Electric lot No. 17

12. Atomic Energy of Canada: "Molybdenum specifications. February 12, 1974

13. Union Carbide: Fission product generator specifications

ERRATUM

In the article "Thallium-201 for Myocardial Imaging: Appearance of the Normal Heart" (*J Nucl Med* 17: 583–589, 1976), it was incorrectly stated that Dr. David J. Cook had been supported by a Clinical Research Fellowship from the Post-Graduate Committee in Medicine of the University of Sydney. Actually, it was Dr. Ian Bailey who had received this Fellowship. Also, the unit of measurement (cm) was omitted from Table 2.