## Lawrence Berkeley National Laboratory

**Recent Work** 

## Title

CARRIER-FREE RADIOISOTOPES FROM CYCLOTRON TARGETS XX. PREPARATION AND ISOLATION OF Ru97,103 FROM MOLYBDENUM

**Permalink** https://escholarship.org/uc/item/2sw7q4jc

## **Authors**

Gile, Jeanne Garrison, Warren Hamilton, Joseph G.

## **Publication Date**

1951-08-22

COPY Z UCRL- 1419 UNCLASSIFIED

# íц **WERKE** CALIFORNIA L O JNIVERSITY

## TWO-WEEK LOAN COPY

This is a Library Circulating Copy which may be borrowed for two weeks. For a personal retention copy, call Tech. Info. Division, Ext. 5545

# RADIATION LABORATORY

## DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.

#### CARRIER-FREE RADIOISOTOPES FROM CYCLOTRON TARGETS

# XX. PREPARATION AND ISOLATION OF Ru97,103 FROM MOLYBDENUM\*

## Jeanne D. Gile, Warren M. Garrison, and Joseph G. Hamilton

Crocker Laboratory, Radiation Laboratory, and Divisions of Medical Physics, Experimental Medicine, and Radiology, University of California, Berkeley and San Francisco, California

### July 12, 1951

Alpha-particle bombardment of molybdenum produces the 2.8-day  $\operatorname{Ru}^{97}$  and the 42-day  $\operatorname{Ru}^{103}$  by the nuclear reactions,  $\operatorname{Mo}(\mathbf{d},\operatorname{xn})\operatorname{Ru}^{97}$ , and  $\operatorname{Mo}^{100}(\mathbf{d},\operatorname{n})\operatorname{Ru}^{103}$ .<sup>(1)</sup> This paper reports a method of isolating this activity in the carrier-free state from the target element and from radioisotopes of Tc (from  $\mathbf{d}$ , pxn reactions) which are produced concurrently by the 40-Mev alpha-particles from the 60-inch cyclotron at the Crocker Laboratory.

The target was a block of C.P. molybdenum metal, silver-soldered to a watercooled copper target plate. It was bombarded with 40-Mev alpha-particles for a total of 70  $\mu$ a-hr. at an average beam intensity of  $8 \mu$ a. The bombarded surface was removed by milling off approximately 0.5 g of Mo. The Mo chips were fused with 5 g of KOH and 0.5 g of KNO<sub>3</sub> in a nickel crucible at 500° C. for 15 minutes. The fused mass was dissolved in a minimum amount of hot water and centrifuged to remove insoluble material. The strongly basic solution was transferred to an allglass distilling flask,<sup>(2)</sup> a stream of Cl<sub>2</sub> was bubbled through the solution, and the active ruthenium was distilled as the RuO<sub>4</sub> which was collected in a trap containing 12 <u>N</u> HCl cooled with ice. Under these conditions the technetium isotopes remain in the distilling flask with the target material. The acid

This document is based on work performed under Contract No. #-7405-eng-48-A for the AEC.

(1) Nuclear Data, National Bureau of Standards, Sept. 1, 1950.

(2) L. A. Scherrer, J. Research Nat. Bur. Stand., 21, 95, (1938).

UCRL-1419

distillate containing the radio-ruthenium was evaporated to a small volume, made basic with NaOH, and redistilled into a trap containing 12 <u>N</u> HCl. The second distillation was done to insure complete separation from the target material. The final HCl solution was evaporated to dryness on 20 mg of NaCl and the activity dissolved quantitatively with the addition of 2 ml of distilled water.

-3-

An aliquot of the preparation was added to a solution containing carrier amounts of Mo and Ru. The activity was quantitatively recovered in the Ru fraction following chemical separation. The radiation characteristics were obtained by aluminum and lead absorption measurements and showed the 0.2-Mev conversion electron and the 0.23-Mev gamma-ray previously reported <sup>(3)</sup> for Ru<sup>97</sup>. The activity showed the 2.8-day period for nearly 15 half-lives at which time it began to lengthen to a period of approximately 40 days, presumably due to Ru<sup>103</sup>.

We wish to thank Professor G. T. Seaborg for helpful suggestions, the staff of the 60-inch cyclotron for bombardments, and Miss Margaret Gee for technical assistance in counting.

(3) D. T. Eggen and M. L. Pool, Phys. Rev. <u>74</u>, 57, (1948).