LETTERS TO THE EDITOR



DISSOLUTION OF METALS BY AC ELECTROLYSIS

Dear Sir:

Reference is made to the subject article by W. D. Box in the August 1966 issue of *Nuclear Applications*. It is of interest to note that the two Russian references published in 1959 and 1963 have an antecedent in the Wohlwill gold-refining process of 1878.

For the electrolysis of high-silver anodes Wohlwill superimposed a low frequency ac on the dc. This counteracted passivity by causing the adherent AgCl to flake off and permitted the use of a higher current density.

Process details are covered in *Electrochemical* Engineering by C. L. Mantell, McGraw Hill, 1960.

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FLUORINE-18 FROM RECOIL DEUTERONS

Dear Sir:

Radiochemical and gamma-ray spectrometry studies have shown the presence of ¹⁸F in the heavy-water moderator-coolant of the Georgia Tech Research Reactor (GTRR). At 1-MW power level the saturation concentration is $3 \times 10^{-3} \ \mu \text{Ci/ml}$.

Several possible mechanisms are available to explain the production of ¹⁸F activity:

(a)	$^{10}O(t,n)$ ^{10}F
(b)	$^{17}O(d,n)$ ^{18}F
(c)	18 O (<i>d</i> ,2 <i>n</i>) 18 F

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We have not been able to produce a measurable quantity of ${}^{18}{\rm F}$ by the irradiation of the same heavy

water in a well-thermalized region of the GTRR, indicating that mechanism (a) involving tritons from the deuterium (n,γ) tritium or from ⁶Li (n,α) tritium is not the explanation. Furthermore, thermal/fast neutron flux ratio measurements using the ⁵⁸Ni(n,p) ⁵⁸Co reaction (threshold 2.9 MeV) have shown the ratio to be greater than 500 for the position used.

This leaves mechanisms (b) and (c) which are caused by high-energy recoil deuterons from the fast fission neutron moderation process. Deuterons of the required energy could be found near the fuel-plate heavy-water interface.

Work reported by Tagami et al.¹ has demonstrated the formation of ¹⁸F in the light-water-moderated Hitachi Training Reactor and ascribed the production to the ¹⁸O (p,n) ¹⁸F reaction. The heavy water within the GTRR does contain approximately 0.5% light water. However, a calculation based on the quantity of ¹⁸F found by Tagami, correcting for the different reactor power and irradiation time still leaves the ¹⁸F production in the GTRR an order of magnitude higher than that reported for the recoil proton mechanism. Differences in neutron spectra in the two reactors would certainly have an effect as would the method of water irradiation. These factors are being evaluated.

Work to reach a more definite conclusion regarding the mechanism of ¹⁸F production in the GTRR is continuing. Currently, experiments involving the irradiation of heavy water with 14-MeV neutrons from the ³H $(d,n)^4$ He reaction indicate the formation of ¹⁸F. If confirmed, this will support the recoil deuteron hypothesis.

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Corrigendum

The following credit should have accompanied "Neutron Activation Analysis of Blood and Blood Serum for Copper and Zinc," by C. C. Thomas, Jr., G. P. Tercho, and J. A. Sondel of Western New York Nuclear Research Center at Buffalo, N. Y. in *Nucl. Appl.*, **3**, 53 (1967):

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^{1.} Takashi Tagami, Masaaki Yamamoto, and Yasuo Osawa, "Formation of ¹³N and ¹⁸F by Recoil Proton Flux in Water Moderated Reactor," NSJ-Tr-39, Japan Atomic Energy Research Institute (1965).