

MASTER

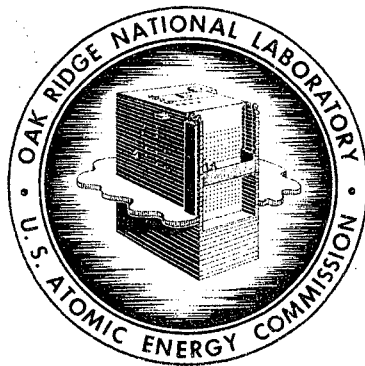
325  
7-27-61

ORNL-3068  
UC-10 - Chemical Separations Processes  
for Plutonium and Uranium

AQUEOUS PROCESSES FOR DISSOLUTION OF  
URANIUM-MOLYBDENUM ALLOY REACTOR

FUEL ELEMENTS

L. M. Ferris



**OAK RIDGE NATIONAL LABORATORY**

operated by

UNION CARBIDE CORPORATION

for the

U.S. ATOMIC ENERGY COMMISSION

Printed in USA. Price \$1.00. Available from the  
Office of Technical Services  
Department of Commerce  
Washington 25, D. C.

LEGAL NOTICE

This report was prepared as an account of Government sponsored work. Neither the United States, nor the Commission, nor any person acting on behalf of the Commission:

- A. Makes any warranty or representation, expressed or implied, with respect to the accuracy, completeness, or usefulness of the information contained in this report, or that the use of any information, apparatus, method, or process disclosed in this report may not infringe privately owned rights; or
- B. Assumes any liabilities with respect to the use of, or for damages resulting from the use of any information, apparatus, method, or process disclosed in this report.

As used in the above, "person acting on behalf of the Commission" includes any employee or contractor of the Commission, or employee of such contractor, to the extent that such employee or contractor of the Commission, or employee of such contractor prepares, disseminates, or provides access to, any information pursuant to his employment or contract with the Commission, or his employment with such contractor.

Contract No. W-7405-eng-26  
CHEMICAL TECHNOLOGY DIVISION  
Chemical Development Section B

AQUEOUS PROCESSES FOR DISSOLUTION OF URANIUM-  
MOLYBDENUM ALLOY REACTOR FUEL ELEMENTS

L. M. Ferris

Technician

J. F. Land

DATE ISSUED

JUL 14 1961

---

Oak Ridge National Laboratory  
Oak Ridge, Tennessee  
Operated by  
Union Carbide Corporation  
for the  
U. S. Atomic Energy Commission

ABSTRACT

Methods for dissolving unirradiated uranium-molybdenum alloy reactor fuels in nitric acid, nitric acid--ferric nitrate, and nitric acid--phosphoric acid solutions were studied on a laboratory scale. Flowsheets based on the results propose dissolution of alloys containing 3% molybdenum in boiling 6 M  $\text{HNO}_3$  to yield stable solutions that are 0.6 M in uranium and 3 to 4 M in nitric acid. The uranium can then be easily decontaminated and recovered in a conventional Purex-type tributyl phosphate solvent extraction process. Alloys containing 10% molybdenum would be dissolved in boiling 11 M  $\text{HNO}_3$ , allowing molybdic oxide to precipitate. The molybdic oxide, which carries 5-10% of the uranium, is removed by centrifugation and the acidity of the supernatant solution adjusted to allow recovery of the uranium by Purex-type solvent extraction procedures. The uranium carried by the molybdic oxide is recovered after the  $\text{MoO}_3$  is dissolved in warm 5 M  $\text{NaOH}$ . Less than 0.1% of the uranium is solubilized during the caustic dissolution.

Alternative methods investigated involve dissolution in nitric acid containing 0.5 to 1 M ferric nitrate to complex the molybdenum. These techniques lead to undesirably large volumes of high-level solvent extraction waste solutions. Phosphate ion is also effective in complexing molybdenum; however, its use in the dissolvent would be purposeless since it must be complexed with iron during solvent extraction.

Rates of reaction of the various alloys and the solubility of molybdic oxide were determined in nitric acid, nitric acid--ferric nitrate, and nitric acid--phosphoric acid solutions.

CONTENTS

	Page
1.0 Introduction	4
2.0 Flowsheets	5
2.1 Alloys Containing 3% Molybdenum	5
2.2 Alloys Containing 10% Molybdenum	6
3.0 Experimental	10
3.1 Decladding	10
3.2 Core Dissolution	11
3.3 Solubility Data	23
3.4 Corrosion Studies	32
4.0 Miscellaneous Related Studies	34
4.1 Dissolution in Dilute Aqua Regia	34
4.2 Other Potential Complexing Agents for Molybdenum	34
4.3 Reduction of Solvent Extraction Waste Volumes	34
4.4 Decladding of Zirconium-clad Fuels	35
5.0 References	35

## 1.0 INTRODUCTION

The objective of the laboratory studies reported here was the development of methods for dissolving uranium-molybdenum alloy reactor fuel elements in aqueous nitrate solutions from which the uranium and plutonium could be decontaminated and recovered by Purex-type (1) solvent extraction processes. The methods evolved are applicable to the processing of the Consumers Public Power (U-10% Mo) and the Detroit Edison blanket (U-3% Mo) fuel elements. Both these elements are clad in stainless steel and bonded with sodium (2,3). Because of the hazard involved in reacting sodium with mineral acids, it was assumed that the cladding and bond would be removed mechanically and that the core alloy would probably be recanned in aluminum before aqueous processing (4a). Laboratory studies consisted of an extensive investigation of dissolution of unirradiated alloys in nitric acid, nitric acid—ferric nitrate, and nitric acid—phosphoric acid solutions since chemical removal of the aluminum can presents no problems.

Preliminary work on the dissolution of uranium-molybdenum alloys in nitric acid, done at ORNL in 1955, has been reported (5). In a process used in the USSR for U-Mo-Mg alloy fuels (6), either ferric nitrate or phosphoric acid was used to complex molybdenum in 5 M  $\text{HNO}_3$  solutions. At Hanford Atomic Products Operation (HAPO) considerable effort has been expended in the development of processes involving dissolution in nitric acid (7) or nitric acid—ferric nitrate (8) solutions. While the recent work at ORNL paralleled that at HAPO somewhat, the objective was different. At HAPO, Redox-type solvent extraction was desired, while Purex was the preferred solvent extraction method at ORNL. As a consequence, dissolution studies at ORNL were aimed at the preparation of solutions containing 2-4 M  $\text{HNO}_3$  instead of the slightly acidic or acid-deficient aluminum nitrate solutions required for the Redox process. Preliminary results of recent ORNL studies were reported previously (4b). Savannah River Laboratory has proposed total dissolution of Detroit Edison fuel (zirconium-clad U-10% Mo alloy). The total dissolution process proposed involves dissolution of the zirconium in dilute hydrofluoric acid followed by addition of 3 M  $\text{HNO}_3$  to dissolve the alloy (9). A dilute solution is maintained so that molybdenum does not

precipitate during dissolution.

In addition to the development of aqueous methods for uranium-molybdenum alloy fuels, a nonaqueous method involving reaction of the alloys with air containing a chlorinating agent has been developed (10). By this method, molybdenum is separated from the uranium as a volatile oxychloride at 300-500°C. The ash is  $U_3O_8$  containing only 5-10% of the original molybdenum and can be dissolved easily in dilute nitric acid to produce solutions suitable as feeds for Purex-type solvent extraction processes.

The preliminary solvent extraction experiments showing the feasibility of using a Purex-type process with solutions containing molybdenum were kindly performed by J. R. Flanary and J. H. Goode, ORNL Chemical Technology Division. Chemical analyses were made by the groups of G. R. Wilson and W. R. Laing of the ORNL Analytical Chemistry Division. The author is particularly indebted to R. L. Sherman for his aid in obtaining x-ray data for the various solid molybdenum compounds encountered in the course of the work. Special credit is also due H. Kubota for developing a technique for determining the acidity of nitric acid-phosphoric acid solutions containing uranium and molybdenum.

## 2.0 FLOWSHEETS

Conditions selected as optimum, primarily from solvent extraction waste volume considerations, result from dissolution of 3% molybdenum alloys in 6 M  $HNO_3$  and the dissolution of 10% molybdenum alloys in 11 M  $HNO_3$ . A stable solution was obtained from the 3% alloys, but most of the molybdenum from a 10% alloy was precipitated. Alternative processes based on the complexing of molybdenum with ferric ion are also presented.

### 2.1 Alloys Containing 3% Molybdenum

The process for elements containing uranium-3% molybdenum alloy (e.g., the Detroit Edison Blanket) involves dissolution of the aluminum can (which may be present as a result of the prior recanning step) in boiling  $NaOH-NaNO_3$  solution (11) followed by dissolution of the alloy in boiling

6 M  $\text{HNO}_3$  (Fig. 1a). Less than 0.1% of the uranium is lost to the decladding solution. Dissolution of the 0.415-in.-dia alloy requires about 6 hr. The product of the core dissolution contains 0.6 M uranium, about 0.04 M molybdenum, and about 3.4 M hydrogen ion. No precipitation occurs at any time during core dissolution. Uranium and plutonium are decontaminated and recovered by Purex-type solvent extraction methods. Preliminary studies (4b) showed that the traces (about 0.013 M) of molybdenum in the feed had no adverse effect on the extraction of uranium with 30% TBP.

Large volumes of solvent extraction waste would result from this process unless the waste solutions were given special treatment. Two alternatives seem feasible: (1) neutralization followed by evaporation, and (2) a boil-down to precipitate the molybdenum and recover acid (Sect. 4.3). The molybdic oxide precipitated during boil-down would be dissolved in sodium hydroxide solution for storage.

In the alternative process for uranium—3% molybdenum fuels, dissolution is achieved in boiling 8 M  $\text{HNO}_3$ —0.5 M  $\text{Fe}(\text{NO}_3)_3$ . Iron was selected as the complexing agent instead of phosphate because it was necessary to complex phosphate with iron prior to solvent extraction. By complexing molybdenum with 0.5 M ferric ion, a solvent extraction feed containing 1 M uranium and about 3 M hydrogen ion can be produced (Fig. 1b). Preliminary solvent extraction studies on solutions spiked with plutonium showed that molybdenum had no adverse effect on the extraction with 30% TBP in 7 stages and that the stage requirements were no greater than those for normal Purex operation (4b).

The chief disadvantage in using iron is that the waste volumes from solvent extraction are very large. Even if the molybdenum could be concentrated to 25 g/liter, about 1100 gal of high-activity waste would be generated for each ton of uranium processed.

## 2.2 Alloys Containing 10% Molybdenum

Fuels containing 10% molybdenum (e.g., the CPPD-1) are dissolved in boiling 11 M  $\text{HNO}_3$  (5) after the aluminum can is removed by the standard caustic decladding method (11) (Fig. 2). Soluble uranium losses to the



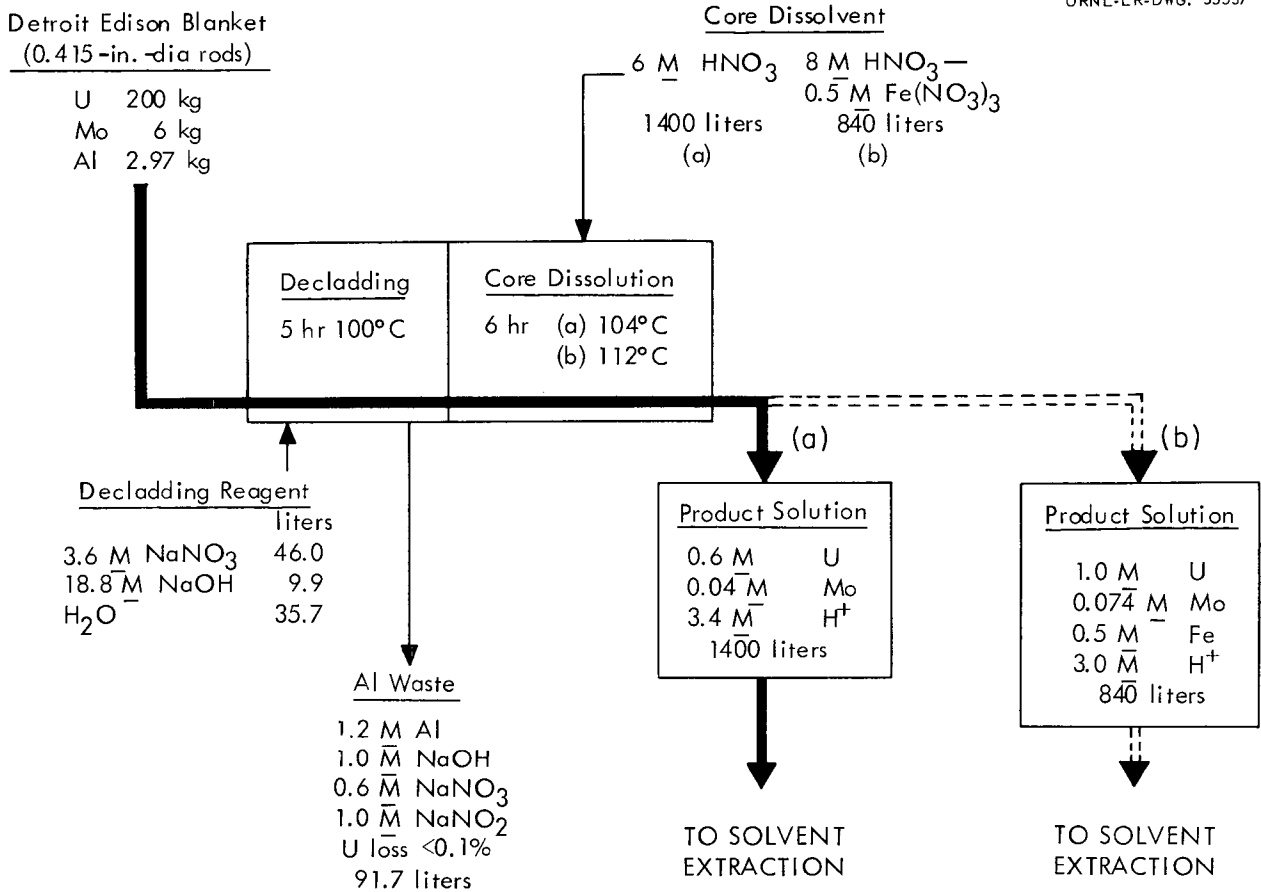


Fig. 1. Processes for dissolution of uranium--3% molybdenum alloy fuels in (a) nitric acid solution and (b) nitric acid--ferric nitrate solution. No molybdenum precipitates during dissolution in either case.

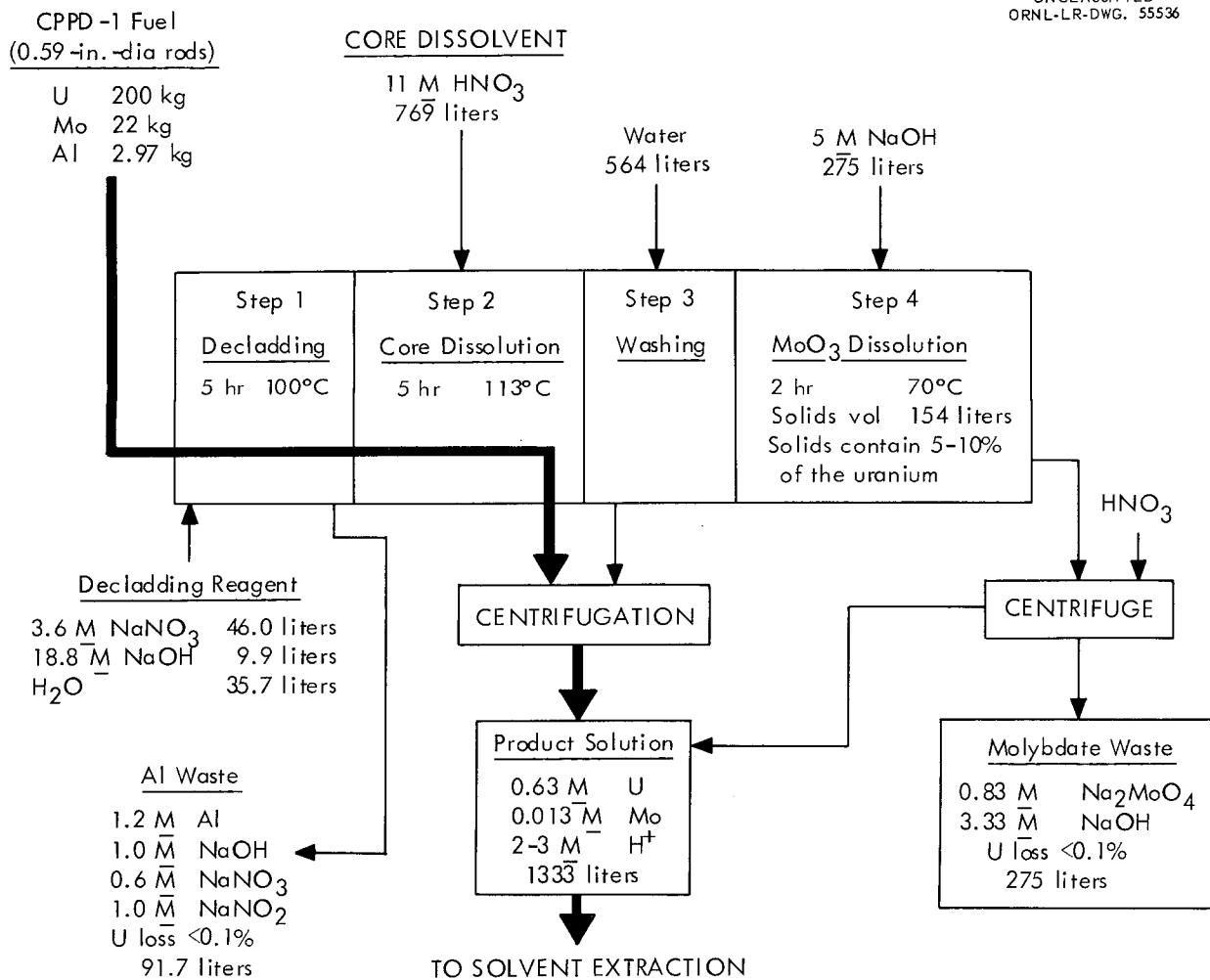


Fig. 2. Process for dissolution of uranium-10% molybdenum alloy fuels in concentrated nitric acid. Molybdenum precipitates during dissolution. Uranium sorbed by MoO<sub>3</sub> is dissolved in nitric acid after the MoO<sub>3</sub> is dissolved in caustic and removed.

decladding solution are less than 0.1%. The 0.59-in.-dia core is dissolved in 5 hr in a volume of boiling acid sufficient to produce a solution containing 260 g of uranium per liter; the acid concentration after dissolution is about 6 M. The uranium concentration is then adjusted to about 0.6 M for solvent extraction.

Most of the molybdenum precipitates, as  $\text{MoO}_3$ , during core dissolution. This solid, after centrifugation, occupies a volume which is 20-30% of the original solution volume. After about three washes, the centrifuged solids contain 5-10% of the uranium (Table 1) and about 2% of the plutonium (7). Recovery of the sorbed uranium is greatly improved when more efficient washing and solid-liquid separation techniques are used. For example, the washed solids contained only about 0.1% of the uranium when the solids were separated by vacuum filtration before washing (Table 1). Residual uranium, and probably plutonium compounds, can be recovered by dissolution in nitric acid after the  $\text{MoO}_3$  is dissolved in 200% stoichiometric excess of hot 5 M NaOH. No significant fraction of the uranium (and probably plutonium) is solubilized when the  $\text{MoO}_3$  is dissolved (Table 1).

The principal advantage of this process is the separation of most of the molybdenum from the uranium prior to solvent extraction. This separation should result in low-volume Purex-type solvent extraction wastes. The chief disadvantage is the necessity of recovering relatively large amounts of solid molybdic oxide containing significant amounts of uranium and plutonium by centrifugation or filtration. The process could also be used for elements containing 3% molybdenum; centrifugation problems with this alloy would undoubtedly be less severe than with the 10% alloy.

Alloys containing 10% molybdenum can be dissolved in nitric acid--ferric nitrate solutions without precipitation occurring. A solvent extraction feed solution containing 0.5 M uranium, 0.14 M molybdenum, and 3.0 M  $\text{HNO}_3$  is obtained by dissolution in boiling 5.5 M  $\text{HNO}_3$ -1 M  $\text{Fe}(\text{NO}_3)_3$ .

Table 1. Dissolution of Uranium-10% Molybdenum Alloy in Boiling 11 M HNO<sub>3</sub>

Solution composition prior to solid-liquid separation:  
about 6 M HNO<sub>3</sub> containing 260 g of uranium per liter

Mo in Alloy, %	Dissolu- tion Time, hr	Separation Method	Wash Solution <sup>a</sup>	Mo in Filtrate or Centrifugate, %	U Content, %	
					Washed Solids	NaOH Solution <sup>b</sup>
8.4	5	Filtration	1 M HNO <sub>3</sub>	6.0	0.1	0.08
8.4	3	Centrifugation	Water	8.4	4.1	0.11
8.4	6	Filtration	Water	5.6	0.1	0.22
10	4	Centrifugation	Water	20.7	7.2	0.005

<sup>a</sup>Wash solutions generally added in three portions.

<sup>b</sup>MoO<sub>3</sub> solids dissolved in 200% excess of hot 5 M NaOH.

### 3.0 EXPERIMENTAL

#### 3.1 Decladding

Both the CPPD-1 core and the Detroit Edison blanket fuels are bonded to stainless steel jackets with sodium. While there appears to be no danger of a vapor-phase explosion if the stainless steel and sodium were dissolved in sulfuric acid (12), sufficient uncertainty exists regarding the containment of the sodium-water reaction that mechanical removal of the clad and bond appears to be the safest approach. After mechanical de-jacketing, the core alloys would probably be recanned in aluminum to facilitate subsequent handling (4a). Chemically, aluminum offers no special problems. Uranium losses were acceptably low when the aluminum was dissolved in sodium hydroxide—sodium nitrate solutions (Table 2). When sulfuric acid is used for dissolution of the original stainless steel cladding, no serious uranium loss results.

Table 2. Losses from Various Uranium-Molybdenum Alloys to Typical Decladding Solutions

Boiling Decladding Solution	Uranium Losses in 24 hr, %		
	From 3% Mo Alloys	From 8.4% Mo Alloys	From 10% Mo Alloys
2 M NaOH—1.78 M NaNO <sub>3</sub>	0.00002	0.00002	0.00005
4 M H <sub>2</sub> SO <sub>4</sub>	0.57	0.022 0.028	--
6 M H <sub>2</sub> SO <sub>4</sub>	0.34 1.6	0.055 0.045	-- --
3.8 M H <sub>2</sub> SO <sub>4</sub> —1.8 M stainless steel	0.31	0.0027 <sup>a</sup>	--

<sup>a</sup>Exposure time 5 hr in this experiment.

### 3.2 Core Dissolution

Dissolution in Nitric Acid. Uranium-molybdenum alloys containing up to 10% molybdenum dissolved rapidly in nitric acid. Because of its low solubility, molybdic oxide usually precipitated during the dissolution. The dissolution rate increased with increasing acid concentration and with the fuel molybdenum content. The three alloys studied dissolved more rapidly than uranium metal, maximum rates at each acid concentration being for the 10% molybdenum alloy (Fig. 3). For each alloy the rate data obeyed approximately the equation

$$\text{Rate (mg min}^{-1}\text{cm}^{-2}) = k (\text{HNO}_3, \text{ M})^n \quad (1)$$

The log-log plots in Fig. 3 show that generally the data could be represented by the above equation when the rate was less than about 100 mg min<sup>-1</sup>cm<sup>-2</sup>. The slopes of these lines, n, indicated a second- or third-power dependence of the rate on the nitric acid concentration (Table 3). Irrespective of the molybdenum content of the alloy, the rate was always greater than 10 mg min<sup>-1</sup>cm<sup>-2</sup> in solutions in which the acid concentration was greater than 5 M. Recent studies have shown that the dissolution rate may be higher after irradiation (13). Rates expressed in mg per minute per square centimeter may be converted to rates in mils per hour by use of the equation

$$\text{Rate (mils/hr)} = 1.3 \text{ Rate (mg min}^{-1}\text{cm}^{-2})$$

UNCLASSIFIED  
ORNL-LR-DWG. 55538

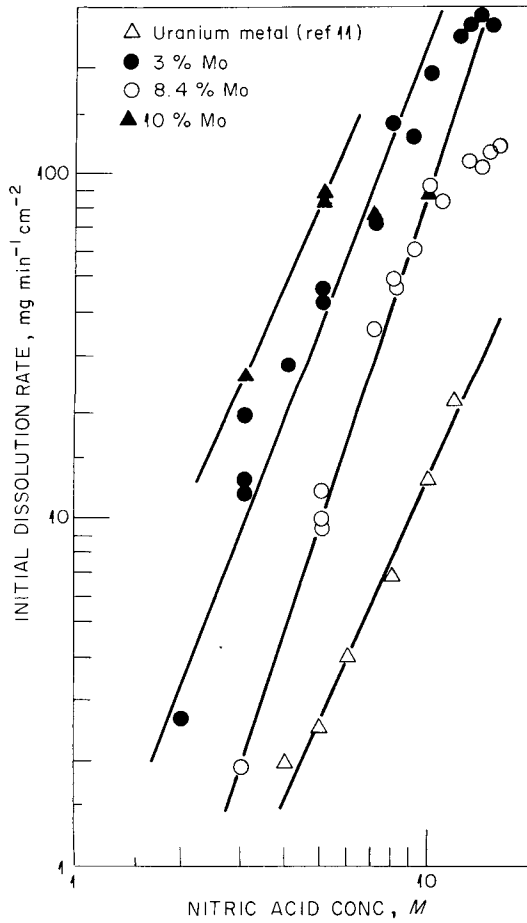


Fig. 3. Initial rates of dissolution of uranium-molybdenum alloys in boiling nitric acid.

UNCLASSIFIED  
ORNL-LR-DWG. 55539

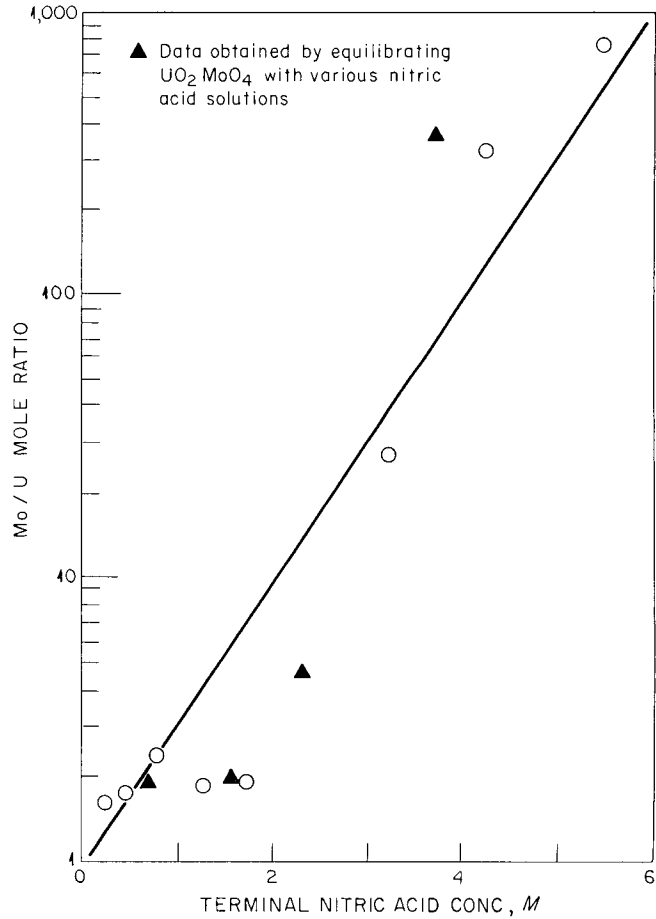


Fig. 4. Molybdenum/uranium mole ratio in solids produced during dissolution of uranium-molybdenum alloys in boiling nitric acid. Final uranium concentration was 1-1.5 M in each case.

Table 3. Dissolution of Uranium-Molybdenum Alloys in Nitric Acid: Constants Obtained When Data Were Expressed by the Equation  

$$\text{Rate (mg min}^{-1}\text{cm}^{-2}) = k (\text{HNO}_3, \text{M})^n$$

Mo Content of Alloy, %	Slope (n)	Intercept (k)
0 <sup>a</sup>	2.23	0.0795
3	2.46	0.763
8.4	3.25	0.0517
10	2.33	2.02

<sup>a</sup>Data for uranium metal taken from reference 11.

Acid Consumption. Dissolution in boiling 3 to 11 M  $\text{HNO}_3$  of alloys containing up to 10% molybdenum resulted in the apparent consumption of 4-7 moles of hydrogen ion for each mole of alloy dissolved. Since the extent to which the nitrogen oxides are refluxed is dependent on the type of condenser used, these data indicate only the approximate stoichiometry of the reaction. Two- to 20-g samples of the alloys were dissolved in 500-ml flasks fitted with two 21-in.-long condensers. With the alloy containing 8.4% molybdenum, between 3.4 and 4.6 moles of hydrogen ion was consumed for each mole of alloy dissolved regardless of the acid concentration (Table 4). The uranium concentration in the final solution varied from 0.4 to 2 M. With both the 3 and 10% alloys the amount of hydrogen ion consumed appeared to increase as the acid concentration increased. For first-approximation calculations it is convenient to assume that 4 moles of nitric acid is required to dissolve 1 mole of alloy.

Formation of Uranyl Molybdates. When the terminal acid concentration was less than about 4 M, the precipitate contained significant amounts of uranium (Table 5). The molybdenum-uranium mole ratio in the solid varied from about 1 (which would correspond to the normal molybdate,  $\text{UO}_2\text{MoO}_4$ ) to about 1000 as

Table 4. Consumption of Acid in Dissolution of Uranium-Molybdenum Alloys in Boiling 3-11 M HNO<sub>3</sub>

Mo in Alloy, %	Initial HNO <sub>3</sub> Conc., M	Alloy Dissolved, moles	H <sup>+</sup> Consumed, moles	Moles H <sup>+</sup> Consumed / Mole Alloy Dissolved
3	3.0	0.0598	0.250	4.2
3	5.0	0.0339	0.122	3.6
3	6.0	0.0883	0.380	4.3
3	8.0	0.0410	0.177	4.3
3	8.0	0.0848	0.398	4.7
3	11.0	0.0832	0.580	7.0
3	11.0	0.0470	0.191	4.1
8.4	3.0	0.0242	0.081	3.4
8.4	8.0	0.0366	0.130	3.6
8.4	11.0	0.0615	0.280	4.6
8.4	11.0	0.0330	0.127	3.9
10	4.0	0.0539	0.192	3.6
10	5.0	0.0686	0.239	3.5
10	5.0	0.0095	0.028	2.9
10	6.0	0.0645	0.238	3.7
10	8.0	0.0706	0.289	4.1
10	8.0	0.0080	0.031	3.9
10	10.0	0.0674	0.340	5.0
10	11.0	0.0727	0.386	5.3
10	11.0	0.0442	0.227	5.1

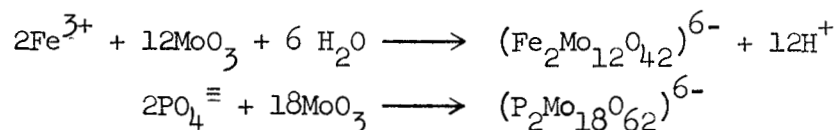
the terminal acid concentration increased from 0 to about 6 M (Fig. 4 and Table 5). Apparently, a series of uranyl molybdates, which are similar to the sodium molybdates (14), is formed in solutions of low acidity. X-ray analyses indicated that the solid was MoO<sub>3</sub> when the terminal acidity was greater than 4 M; therefore uranium carried by the solid during dissolution in concentrated acid is probably only sorbed and can be recovered by washing with water or acid. Water is preferred for the washing to minimize solubilization of the molybdenum (see data on the solubilities of MoO<sub>3</sub> and UO<sub>2</sub>MoO<sub>4</sub> in nitric acid, Sect. 3.3).



Table 5. Solids Produced During Dissolution of Uranium-Molybdenum Alloys in Nitric Acid

Mo in Alloy, %	H <sup>+</sup> Conc, M		U Conc, M	Solids		Mo/U Mole Ratio in Solids
	Initial	Final		U, %	Mo, %	
3	5.0	0.43	1.22	38.9	28.2	1.80
3	8.0	1.25	1.64	40.0	31.1	1.93
3	11.0	5.55	1.58	-	60.1	-
10	5.0	0.80	0.94	40.0	38.5	2.39
10	8.0	3.20	1.05	5.05	55.6	27.3
10	11.0	5.50	1.11	0.21	61.5	727
10	4.0	0.23	0.85	39.6	27.9	1.75
10	6.0	1.68	0.96	40.3	31.4	1.93
10	10.0	4.45	1.12	0.46	60.8	328

Dissolution in Nitric Acid—Ferric Nitrate Solutions. Ferric and phosphate ions strongly complex molybdenum in dilute nitric acid solution. Heteropolymolybdate ions are formed by reactions similar to the following (15, 16):



At each acid concentration, uranium-molybdenum alloys dissolved in nitric acid—ferric nitrate solutions more rapidly than in nitric acid alone. For example, the initial (5-min) rate of dissolution of U-3% Mo alloy in boiling 8 M HNO<sub>3</sub> increased from 142 to about 200 mg min<sup>-1</sup> cm<sup>-2</sup> as the ferric nitrate concentration increased from 0 to 1 M (Table 6). A complete study of the off-gases has not yet been made. However, work at HAPO (8) indicated that the gas mixture evolved during dissolution was about 70% NO and 25% NO<sub>2</sub>. Less than 0.2% hydrogen was present in the gas.

In most cases, the dissolution rate showed a first-order dependence on the ferric nitrate concentration regardless of the nitric acid concentration. The data were found to fit approximately the

Table 6. Initial Rates of Dissolution of Uranium-Molybdenum Alloys in Boiling Nitric Acid—Ferric Nitrate Solutions

Mo in Alloy, %	HNO <sub>3</sub> Conc, M	Fe(NO <sub>3</sub> ) <sub>3</sub> Conc, M	Rate, mg min <sup>-1</sup> cm <sup>-2</sup>
3	3.0	0.0	20
		0.25	30
		0.50	42
		0.75	65
		1.0	93
3	8.0	0.0	142
		0.25	195, 177
		0.50	204, 224
		0.75	201, 199
		1.0	200, 197
3	10.0	0.0	199
		0.25	311, 279
		0.50	241
		0.75	223
		1.0	205
8.4	5.0	0.0	10
		0.25	34
		0.50	60
		0.75	89
		1.0	94
		1.0	103
10	3.0	0.0	26
		0.25	50
		0.50	57
		0.75	76
		1.0	90
10	8.0	0.0	77
		0.25	91, 90
		0.50	114
		0.75	125
		1.0	132
10	10.0	0.0	90
		0.5	126
		0.75	146
		1.0	148

following type of equation:

$$R \text{ (mg min}^{-1}\text{cm}^{-2}\text{)} = k \text{ (HNO}_3, \underline{M})^n + j \text{ (Fe, } \underline{M})^p \quad (2)$$

With an alloy of known composition, the quantity  $k \text{ (HNO}_3)^n$  may be expressed as a constant,  $K$ , at each nitric acid concentration. The  $K$  may be evaluated from the values of  $k$  and  $n$  in Table 3 and the nitric acid concentration.

Equation 2 then becomes

$$R = K + j \text{ (Fe)}^p \quad (3)$$

or, by rearranging and taking logarithms,

$$\log (R - K) = \log j + p \log (\text{Fe}) \quad (4)$$

The quantity  $(R - K)$  is the increase in rate due to the presence of ferric nitrate. Plots of  $\log (R - K)$  vs  $\log (\text{Fe})$  were linear with slopes equal to  $p$ , the order of the reaction with respect to the ferric nitrate concentration (Fig. 5). Values for  $p$  and  $j$  (Table 7) show that the reaction for

Table 7. Constants Obtained When Dissolution Rates in Nitric Acid—  
Ferric Nitrate Solutions were Correlated by the Equation

$$\text{Rate} = k \text{ (HNO}_3)^n + j \text{ (Fe)}^p$$

Mo in Alloy, %	HNO <sub>3</sub> Conc, M	p	j
3	3	1.14	75.9
3	3	1.42	66.7
3	8	0.22	82.8
3	8	0.32	67.0
8.4	5	0.90	91.2
10	3	0.75	61.9
10	8	1.05	64.9
10	10	1.05	64.9

UNCLASSIFIED  
ORNL-LR-DWG. 55535

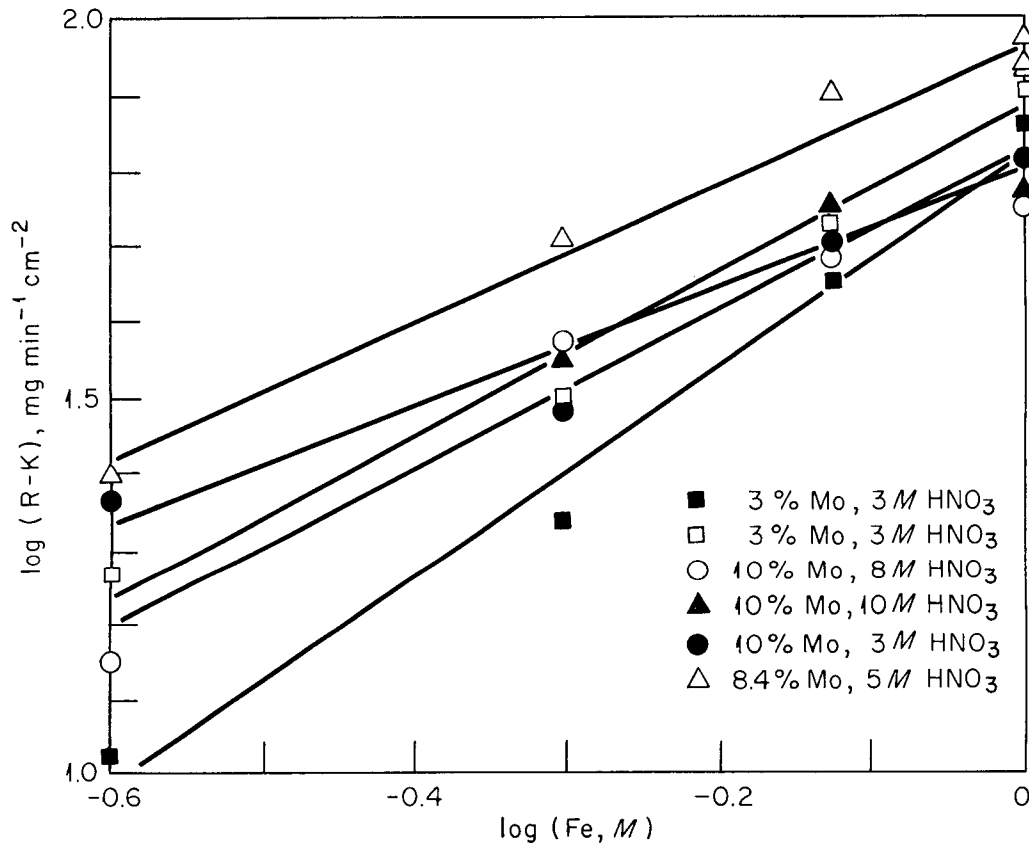


Fig. 5. Correlation of dissolution rates of uranium-molybdenum alloys in boiling nitric acid—ferric nitrate solutions by use of eq. 4. Slopes of the lines indicate a first-order dissolution rate with respect to the ferric nitrate concentration.

the rate increase due to ferric nitrate is approximately first order with respect to the ferric nitrate concentration, with the rate constant being about 65. The fact that the rate data can be correlated by means of an equation of the form of eq. 2 indicates that uranium-molybdenum alloys dissolve by means of separate, competing reactions involving nitric acid and ferric nitrate.

The presence of ferric nitrate apparently had little effect on the amount of acid consumed, which generally varied between 3 and 5 moles per mole of alloy dissolved (Table 8). For convenience it is assumed that 4 moles of nitric acid is consumed in the dissolution of 1 mole of alloy.

Table 8. Consumption of Nitric Acid in Dissolution of Uranium-Molybdenum Alloys in Boiling Nitric Acid--Ferric Nitrate Solutions

Mo in Alloy, %	Initial HNO <sub>3</sub> Conc, M	Fe(NO <sub>3</sub> ) <sub>3</sub> Conc, M	Alloy Dissolved, moles	Acid Consumed, moles	Moles Acid Consumed Mole Alloy Dissolved
0	5.0	1.0	0.0286	0.207	7.2
0	5.0	1.0	0.0321	0.179	5.6
3	5.0	0.3	0.0840	0.332	4.0
3	5.0	0.7	0.0846	0.354	4.2
3	5.0	1.0	0.0872	0.376	4.3
3	8.0	0.5	1.01 <sup>a</sup>	5.36 <sup>a</sup>	5.3
3	8.0	0.5	0.0488	0.266	5.4
3	8.0	0.5	0.0523	0.297	5.7
3	8.0	0.5	0.0414	0.174	4.2
8.4	5.0	0.25	0.732 <sup>a</sup>	2.01 <sup>a</sup>	2.7
8.4	5.0	0.50	0.661 <sup>a</sup>	1.78 <sup>a</sup>	2.7
8.4	5.0	0.75	0.793 <sup>a</sup>	2.50 <sup>a</sup>	3.2
8.4	5.0	1.0	0.931 <sup>a</sup>	3.06 <sup>a</sup>	3.3
8.4	5.0	1.0	0.814 <sup>a</sup>	2.50 <sup>a</sup>	3.1
8.4	5.0	1.0	1.061 <sup>a</sup>	3.54 <sup>a</sup>	3.3
8.4	5.0	1.0	0.924 <sup>a</sup>	3.25 <sup>a</sup>	3.5
8.4	5.0	1.0	0.0652	0.250	3.8
8.4	5.0	1.0	0.0691	0.240	3.5
8.4	5.0	1.0	0.0686	0.253	3.7
8.4	5.0	1.0	0.0702	0.301	4.3
8.4	5.0	1.0	0.0715	0.321	4.5
10	2.0	1.0	0.0301	0.0650	2.2
10	5.0	1.0	0.0604	0.201	3.3
10	8.0	1.0	0.0922	0.359	3.9

<sup>a</sup>Quantities given are per liter of solution.

Densities at 26°C of nitric acid—ferric nitrate solutions containing less than 0.1 M molybdenum are given in Fig. 6 and can be expressed by the equation

$$d^2 \text{ (g/ml)} = 0.0719(\text{HNO}_3, \text{ M}) + 0.425(\text{Fe}, \text{ M}) + 0.994$$

The boiling points of several nitric acid—ferric nitrate and nitric acid—phosphoric acid solutions are given in Table 9.

Table 9. Boiling Points of Several Nitric Acid—Ferric Nitrate and Nitric Acid—Phosphoric Acid Solutions

HNO <sub>3</sub> Conc, M	Fe(NO <sub>3</sub> ) <sub>3</sub> Conc, M	H <sub>3</sub> PO <sub>4</sub> Conc, M	Boiling Point, °C
3	0	0	101
3	0.25	0	102
3	0.5	0	103
3	1.0	0	-
8	0	0	108
8	0.25	0	110
8	0.5	0	112
8	1.0	0	114
10	0	0	112
10	0.25	0	113
10	0.5	0	114
10	1.0	0	118
3	0	0.25	101
3	0	0.5	102
3	0	1.0	103
8	0	0.25	108
8	0	0.5	109
8	0	1.0	110
10	0	0.25	111
10	0	0.5	112
10	0	1.0	113

Dissolution in Nitric Acid—Phosphoric Acid Solutions. Uranium-molybdenum alloys dissolve at high rates in nitric acid—phosphoric acid solutions, with maximum rates at about 0.25 M H<sub>3</sub>PO<sub>4</sub> (Fig. 7). For example, the rate of dissolution of uranium-3% molybdenum alloy in boiling 10 M HNO<sub>3</sub>

UNCLASSIFIED  
ORNL-LR-DWG. 55545

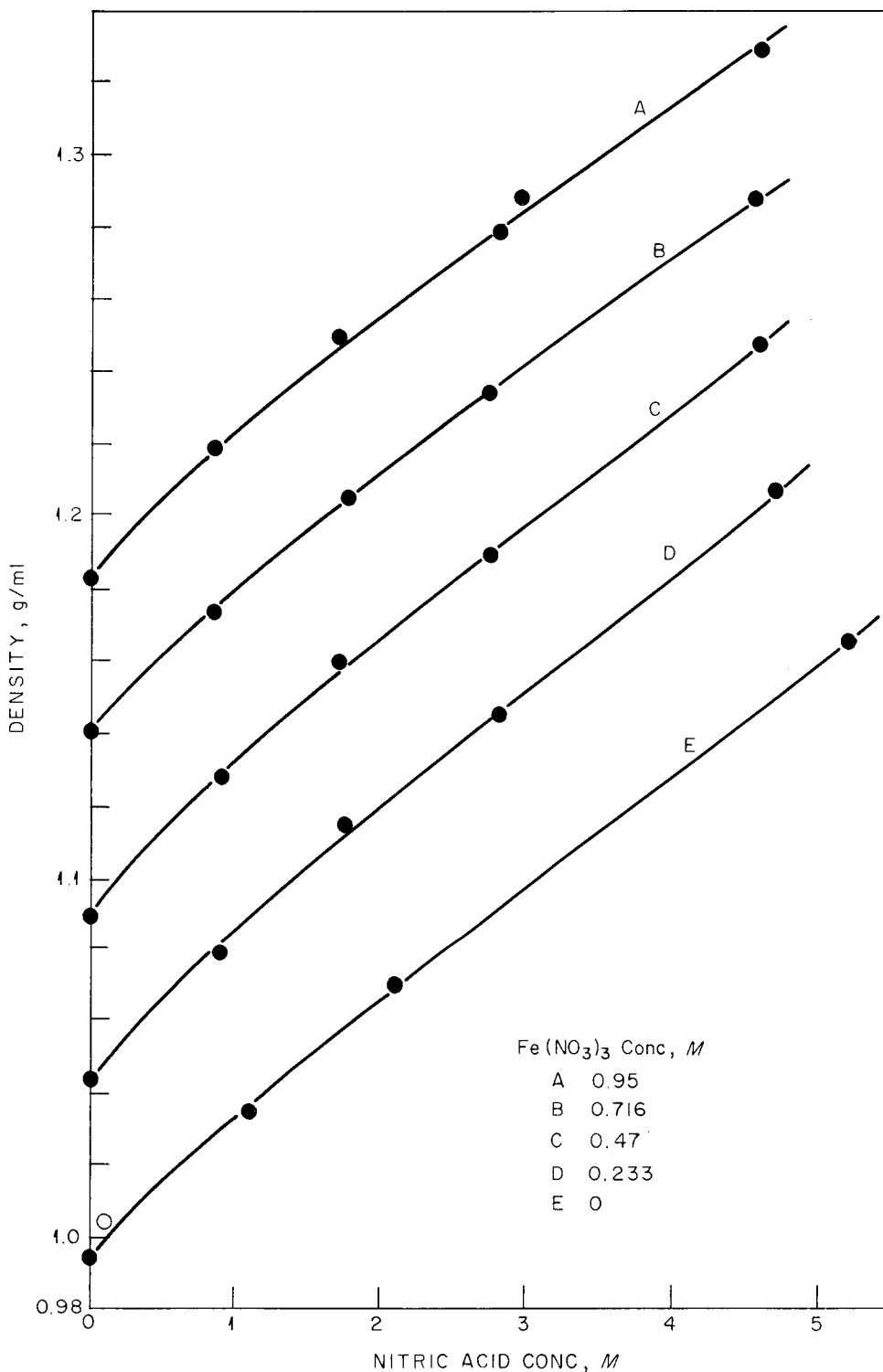


Fig. 6. Densities of nitric acid—ferric nitrate solutions at 26°C (molybdenum concentration less than 0.1 M).

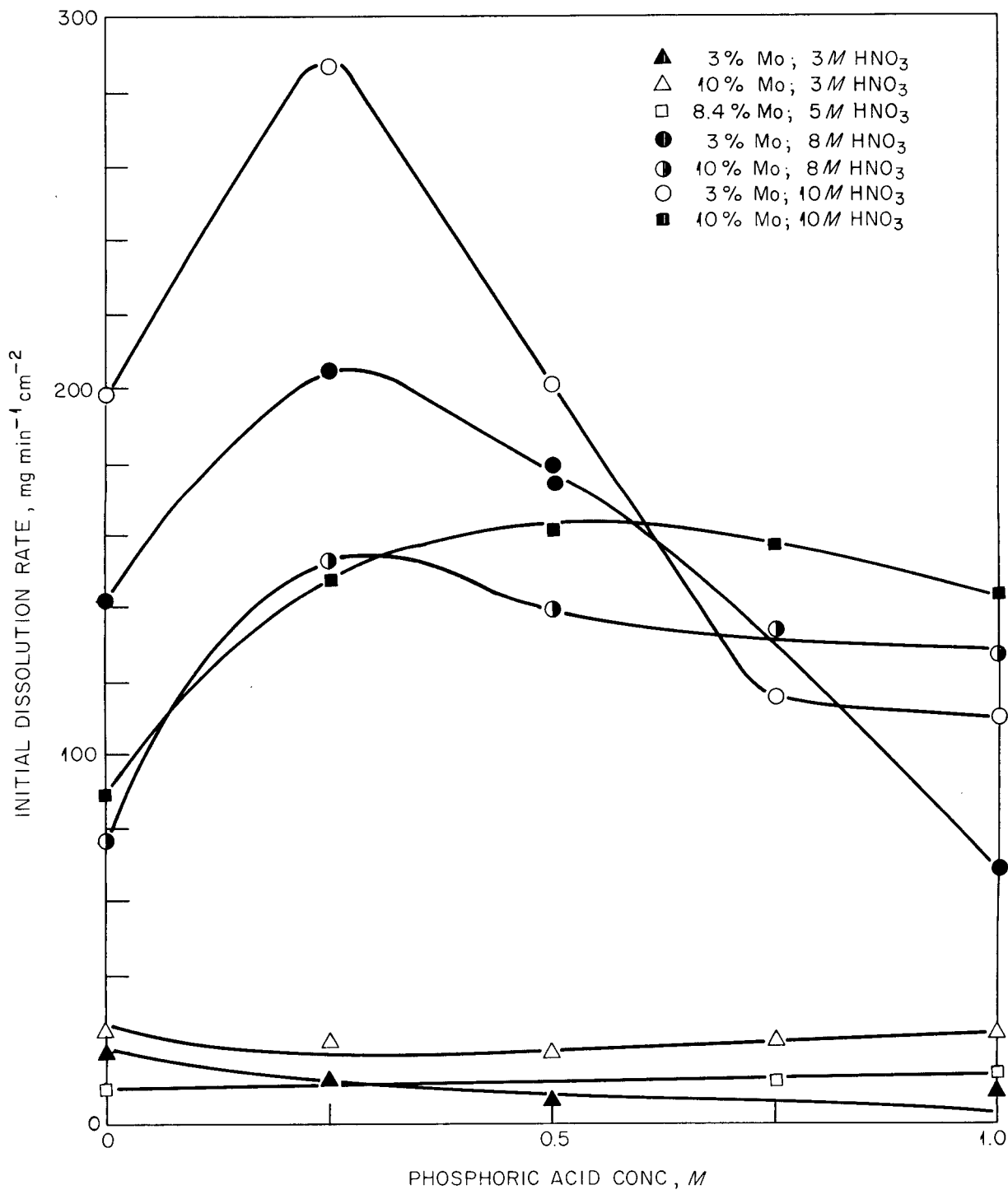


Fig. 7. Effect of nitric acid concentration, phosphoric acid concentration, and alloy composition on the rate of dissolution of uranium-molybdenum alloys in boiling nitric acid—phosphoric acid solutions.



increases from 200 to about 290  $\text{mg min}^{-1}\text{cm}^{-2}$  when the phosphoric acid concentration is increased from 0 to 0.25 M; a further increase in phosphoric acid concentration to 1 M causes a decrease in rate to about 110  $\text{mg min}^{-1}\text{cm}^{-2}$ . As expected, in solutions of constant phosphoric acid concentration the rate increased with increasing nitric acid concentration. With alloys containing 3 and 10% molybdenum, the rate of reaction in boiling 8 to 10 M  $\text{HNO}_3$  containing 0 to 1 M phosphoric acid was always greater than 60  $\text{mg min}^{-1}\text{cm}^{-2}$ .

In boiling 5 to 12 M  $\text{HNO}_3$  containing 0 to 1 M  $\text{H}_3\text{PO}_4$ , between 3 and 5 moles of hydrogen ion was consumed in the dissolution of 1 mole of 10% Mo alloy irrespective of the initial acid or phosphate concentration (Table 10). However, the nitrate ion consumption increased from about 1.5 to about 3 moles per mole of alloy dissolved when the acid concentration was increased from 5 to 12 M. The phosphate concentration did not change significantly during dissolution.

### 3.3 Solubility Data

The solubilities of  $\text{MoO}_3$ ,  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$ , and  $\text{UO}_2\text{MoO}_4$  were measured in various nitric acid, nitric acid—ferric nitrate, nitric acid—phosphoric acid and sodium hydroxide solutions to aid in the determination of optimum process conditions.

The various molybdenum compounds used in the solubility studies were prepared as follows:  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  was freshly precipitated before use by acidifying a solution of c.p. grade sodium molybdate according to the procedure given by Freedman (17). Hydrous  $\text{MoO}_3$  was obtained by digesting about 120 g of c.p. sodium molybdate with 8 M  $\text{HNO}_3$  for about 15 hr, filtering off the supernatant liquid, and digesting the solids again with 5 M  $\text{HNO}_3$  for 5 hr. The final product was  $\text{MoO}_3$  containing about 3% water.  $\text{UO}_2\text{MoO}_4$  was obtained by mixing 25 ml of 1 M  $\text{Na}_2\text{MoO}_4$  solution with 25 ml of 1.08 M  $\text{UO}_2(\text{NO}_3)_2$  solution at room temperature. After water washing, the yellow solid contained 22.2% molybdenum.  $\text{UO}_2\text{MoO}_4$  contains 22.3% molybdenum.

All samples were equilibrated at least 3 weeks before analysis.

Table 10. Consumption of Hydrogen Ion and Nitrate Ion in the Dissolution of Uranium-10% Molybdenum Alloy in Boiling  $\text{HNO}_3$ - $\text{H}_3\text{PO}_4$  Solutions

Initial $\text{HNO}_3$ Conc, M	$\text{H}_3\text{PO}_4$ Conc, M		Moles Consumed per Mole of Alloy	
	Initial	Final	Hydrogen Ion	Nitrate
5.0	0	0	2.9	0.6
5.0	0.05	0.05	3.8	1.4
5.0	0.1	0.11	3.4	1.3
5.0	0.25	0.26	3.4	1.5
5.0	0.5	0.48	2.9	1.6
5.0	1.0	0.98	2.8	1.6
8.0	0	0	3.9	-
8.0	0.05	0.05	3.3	1.8
8.0	0.1	0.11	3.2	1.8
8.0	0.25	0.25	4.9	2.2
8.0	0.5	0.51	3.6	1.8
8.0	1.0	1.1	3.1	1.7
12.0	0.05	0.05	6.3	4.2
12.0	0.1	0.11	4.8	3.0
12.0	0.25	0.27	4.5	3.1
12.0	0.5	0.56	4.5	3.0
12.0	1.0	-	4.0	2.8

Solubility of Molybdenum Oxide in Nitric Acid. At  $26^\circ\text{C}$ , the solubility of  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  attains its maximum value of about 0.13 M when the nitric acid concentration is about 3.5 M (Fig. 8a) (18). The solubility of  $\text{MoO}_3$  in nitric acid is also maximum at about 3 M acid, but is only about 0.05 M (Fig. 8a). The solubilities of  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  and  $\text{MoO}_3$  (19) decrease with increasing temperature in solutions where the nitric acid concentration is greater than about 2 M (Fig. 8a). Data obtained by Cannon (20) with  $\text{MoO}_3 \cdot \text{H}_2\text{O}$  indicate that the solubility is nil in solutions where the acid concentration is greater than about 6 M. For maximum separation of the molybdenum, then, dissolution of 10% alloys in nitric acid must be carried out under conditions which ensure that the acidity of the solution produced is greater than 6 M.

If the uranium had a negligible effect on the solubility, alloys containing 3 and 10% molybdenum could be dissolved in nitric acid to produce

solutions containing about 3 M  $\text{HNO}_3$  and 0.18 and 0.65 M uranium, respectively. Actually, the highest uranium concentration achieved with 3% alloys was about 0.6 M.

Solubility of  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  in Nitric Acid—Uranyl Nitrate Solutions. In solutions where the nitric acid concentration was less than about 2 M, the solubility of  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  was lowered by the presence of uranyl nitrate (Fig. 8b) (18). For example, in about 3.5 M  $\text{HNO}_3$ , the solubility was reduced from 0.13 to 0.07 M when the uranyl nitrate concentration was increased from 0 to 0.95 M. Corresponding data for the solubility of  $\text{MoO}_3$  were not obtained, but it is assumed that uranyl nitrate would have a similar effect, and that the solubility values would be slightly lower than those obtained with the dihydrate. The fact that the solubility of the dihydrate is higher in solutions containing uranium at low acidity is an indication that some complex ion formation occurs between uranium and molybdenum in these solutions.

Solubility of Uranyl Molybdate ( $\text{UO}_2\text{MoO}_4$ ) in Nitric Acid. Since the solids formed in the dissolution of uranium-molybdenum alloys in nitric acid appeared to be a series of uranyl molybdates (Sect. 3.2), it was of interest to determine the solubility behavior of such compounds in nitric acid. When the nitric acid concentration after dissolution was greater than about 4 M, the solids formed were  $\text{MoO}_3$ . The solubility of this compound in nitric acid was discussed above. However, when the dissolution was carried out under such conditions that solids precipitated from solutions with an acid concentration less than 4 M, the solids contained both uranium and molybdenum (Table 5).

The normal molybdate,  $\text{UO}_2\text{MoO}_4$ , was selected for study. In solutions containing less than about 3 M  $\text{HNO}_3$ , uranium and molybdenum from the solids were solubilized to about the same extent (Fig. 9). However, at higher acidities the molybdenum solubility decreased from its maximum value of about 0.18 M; the uranium concentration increased with increasing acid concentration in all cases. These data further illustrate that solids, if formed during dissolution with nitric acid, are best washed either with water or with nitric acid more concentrated than 4 M if significant solubilization of the molybdenum is to be avoided.

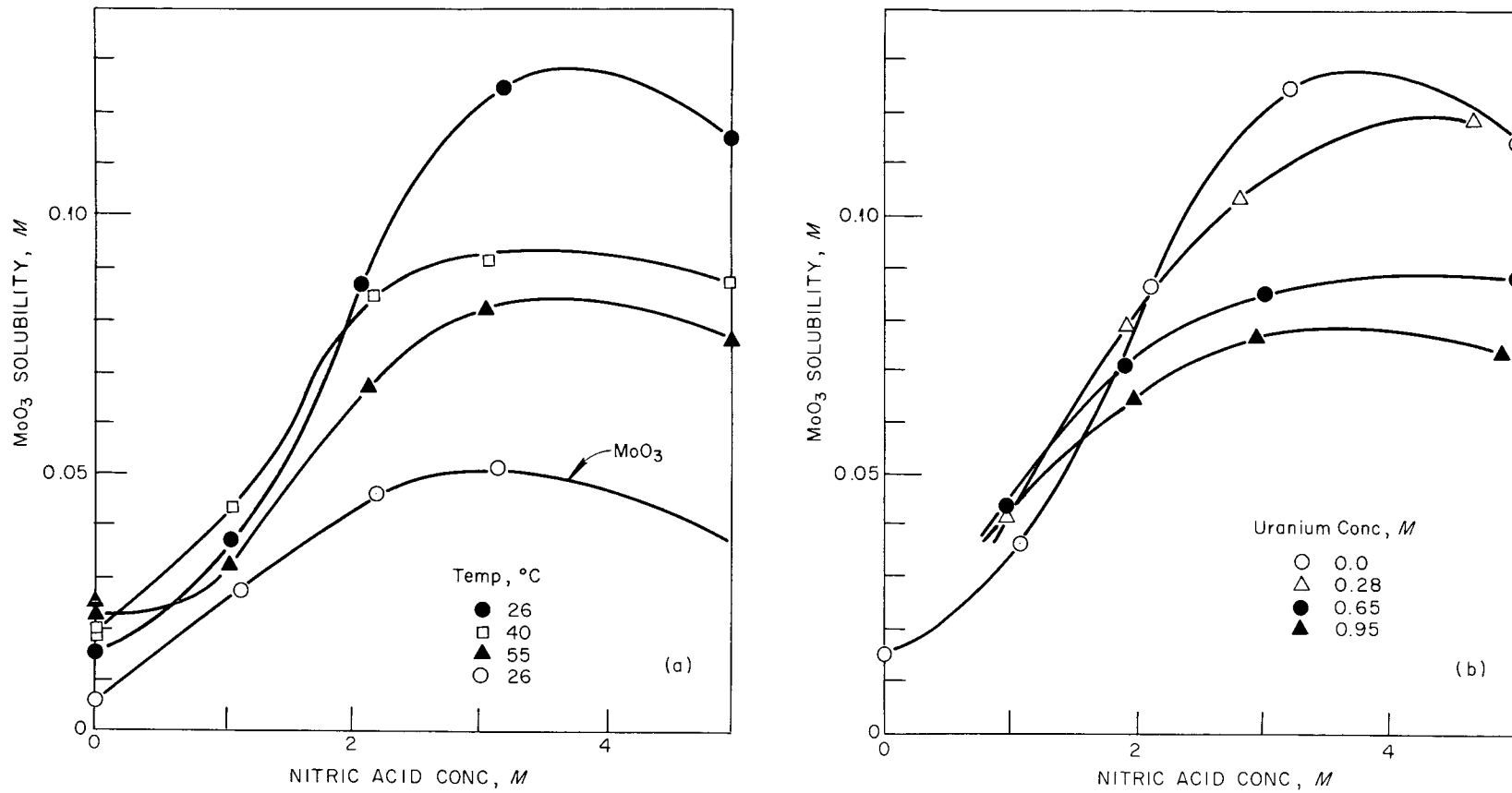


Fig. 8. Solubility (a) of MoO<sub>3</sub>·2H<sub>2</sub>O in nitric acid at various temperatures and of MoO<sub>3</sub> at 26°C; and (b) of MoO<sub>3</sub>·2H<sub>2</sub>O in nitric acid—uranyl nitrate solutions at 26°C.

The uranium content of the solid phase after equilibration decreased with increasing acid concentration (Table 11), as expected from the results given in Fig. 9. Within the limits of error, the variation in composition of the solid phase with acid concentration was the same as that in solids obtained from dissolution of the alloys (see Fig. 4).

Table 11. Solid Phases Resulting from Equilibration of  $UO_2MoO_4$  with Nitric Acid at 26°C

Nitric Acid Conc, M	Composition of Solid Phase, %		Mo/U Mole Ratio in Solids
	Uranium	Molybdenum	
0.73	34.5	26.9	1.94
1.54	36.0	29.0	2.00
2.30	20.6	38.1	4.59
3.56	0.36	43.8	302

Solubility of Molybdenum Oxide in Nitric Acid—Ferric Nitrate Solutions.

Stabilities of solutions resulting from the dissolution of uranium-3% and -10% molybdenum alloys in nitric acid—ferric nitrate solutions were reported by HAPO (8). These data, which showed the maximum alloy concentration attainable in dilute nitric acid solutions containing up to 1 M ferric nitrate, are useful in determining conditions for preparing Redox process solvent extraction feed solutions. The solubility studies reported here were undertaken to obtain data in more concentrated acid solutions and in solutions containing no uranium. The latter data are useful in selecting conditions for storing aqueous Purex-type solvent extraction wastes.

The solubilities of both  $MoO_3$  and  $MoO_3 \cdot 2H_2O$  were determined in nitric acid—ferric nitrate solutions. The solubilities at 26°C were affected by the ferric nitrate and nitric acid concentrations in a similar manner, i.e., the solubility of the dihydrate (18) was about twice that of the oxide (Table 12), other conditions being the same (Figs. 10 and 11). When the

UNCLASSIFIED  
ORNL-LR-DWG. 55541

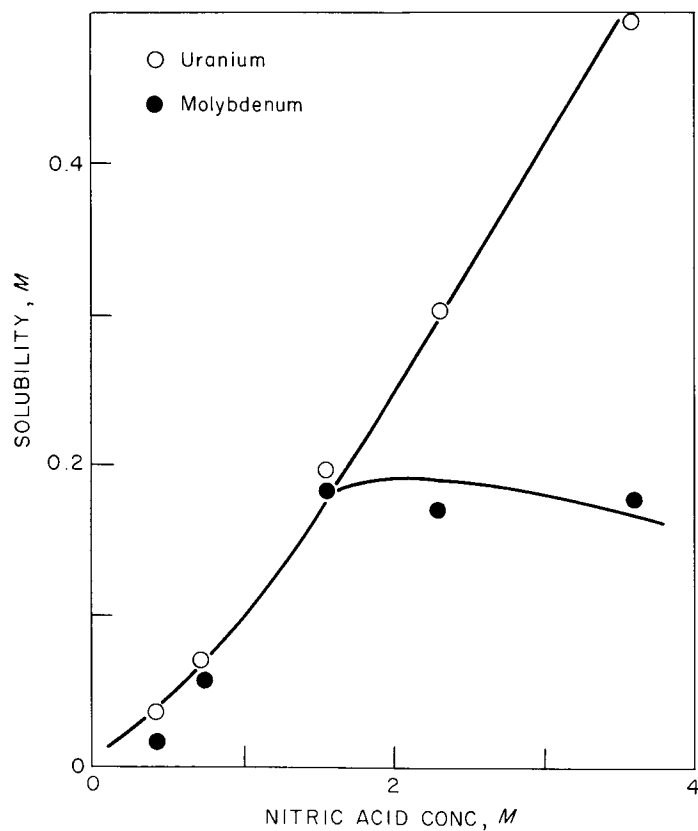


Fig. 9. Solubility of  $UO_2MoO_4$  in dilute nitric acid solutions at 26°C.

UNCLASSIFIED  
ORNL-LR-DWG. 55542

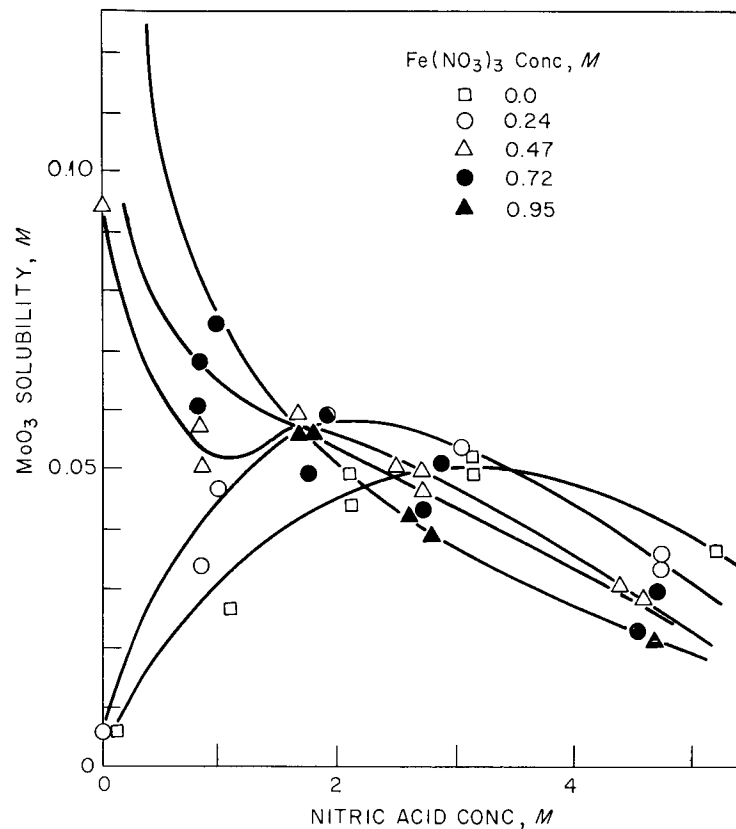


Fig. 10. Solubility of  $MoO_3$  in aqueous nitric acid—ferric nitrate solutions at 26°C.

Table 12. Solubility of MoO<sub>3</sub> in Nitric Acid and Nitric Acid-Ferric Nitrate Solutions at 26°C

Fe Conc, M	H <sup>+</sup> Ion Conc, M	Mo Conc, M	Fe Conc, M	H <sup>+</sup> Ion Conc, M	Mo Conc, M
0	0.11	0.0066	0.716	0.84	0.0608, 0.0677
0	1.08	0.0269	0.717	0.97	0.0743
0	2.12	0.0443, 0.0490	0.713	1.75	0.0493
0	3.14	0.0493, 0.0521	0.752	1.93	0.0597
0	5.20	0.0364	0.713	2.73	0.0434
			0.752	2.88	0.0511
0.233	0.0	0.0056	0.705	4.55	0.0227
0.233	0.86	0.0329, 0.0337	0.717	4.70	0.0295
0.242	1.00	0.0468			
0.242	1.95	0.0598	0.950	0.0	0.262
0.242	3.05	0.0534	0.951	1.67	0.0508, 0.0554
0.231	4.73	0.0333, 0.0332	0.950	1.80	0.0563
0.233	4.75	0.0359	0.950	2.60	0.0426
			0.951	2.78	0.0388
0.476	0.0	0.0941	0.942	4.60	0.0225, 0.0239
0.476	0.81	0.0575	0.950	4.70	0.0217
0.467	0.88	0.0505			
0.476	1.67	0.0595			
0.471	1.73	0.0551			
0.476	2.50	0.0501			
0.473	2.73	0.0461, 0.0499			
0.456	4.40	0.0301			
0.460	4.58	0.0283			

nitric acid concentration was constant and less than 2 M, the solubilities increased with increasing ferric nitrate concentration. On the other hand, the solubilities were always lowered by the addition of ferric nitrate to solutions having nitric acid concentrations greater than about 3 M.

The solubility data point out the advantage of using iron to complex molybdenum in dilute nitric acid solutions. Experience has shown that the data for the dihydrate best describe the solubility limitations to be expected in the dissolution of alloys.

Although the effect of uranium on the solubility of molybdenum oxide in nitric acid—ferric nitrate solutions was not investigated directly, data from HAPO (8) showed the solubility in dilute nitric acid of alloys containing 3 and 10% molybdenum to increase with increasing nitric acid concentration when the ferric nitrate concentration was constant. Solubilities were not determined in solutions more concentrated in nitric acid than 3 to 4 M. Increasing the ferric nitrate concentration increased the solubility in solutions of constant acidity.

Although ferric nitrate is useful in complexing molybdenum during alloy dissolution, its presence in solution complicates subsequent waste handling. For example, dissolution of a fuel containing 3% molybdenum according to the flowsheet shown in Fig. 1b would result in about 1100 gal of high-activity solvent extraction waste solution for each ton of fuel processed even if the waste should be concentrated to 0.25 M molybdenum. The solubility data show that this molybdenum concentration is attainable only in solutions of very low acidity. Extreme care would be required in the partial neutralization and evaporation of the waste to avoid precipitation of ferric hydroxide.

Solubility of MoO<sub>3</sub> in Nitric Acid—Phosphoric Acid Solutions. The solubility of MoO<sub>3</sub> was determined at 26°C in solutions which were initially 0 to 5 M nitric acid containing 0 to 0.5 M phosphoric acid (Table 13). The solubility was highest in solutions containing no nitric acid and passed through a maximum of about 2.2 M when the phosphoric acid concentration was about 0.3 M (Fig. 12). In solutions of constant phosphoric acid concentration the solubility decreased with increasing nitric acid concentration; e.g., from about 2.2 to 0.12 M in solutions containing 0.3 M phosphoric acid when the nitric acid concentration was increased from 0 to 5 M. The solubilities in solutions containing no nitric acid were much higher than those observed by Cannon (20) for the monohydrate, MoO<sub>3</sub>·H<sub>2</sub>O.

As illustrated by the lower curves in Fig. 12, little benefit is derived from the use of phosphoric acid as a complexing agent for molybdenum in solutions where the nitric acid concentration is greater than about 2 M. This behavior, in addition to the fact that ferric ion is required to complex the phosphate in the subsequent solvent extraction step (21), makes the use of phosphate impractical.



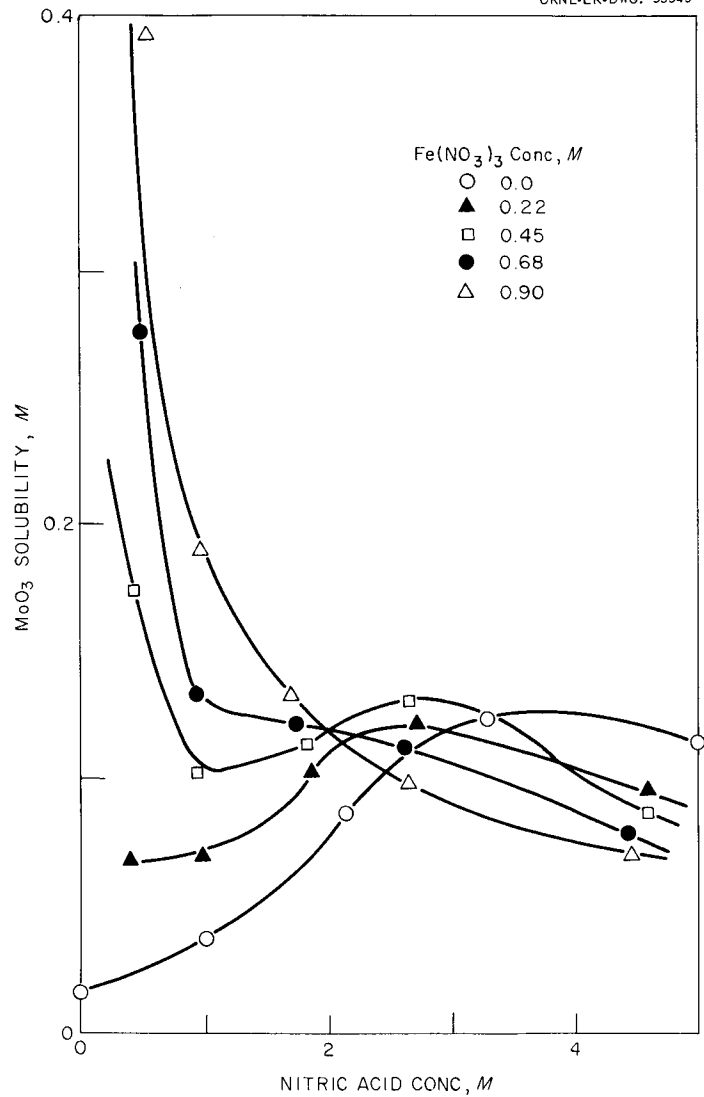


Fig. 11. Solubility of  $\text{MoO}_3 \cdot 2\text{H}_2\text{O}$  in nitric acid—ferric nitrate solutions at  $26^\circ\text{C}$ .

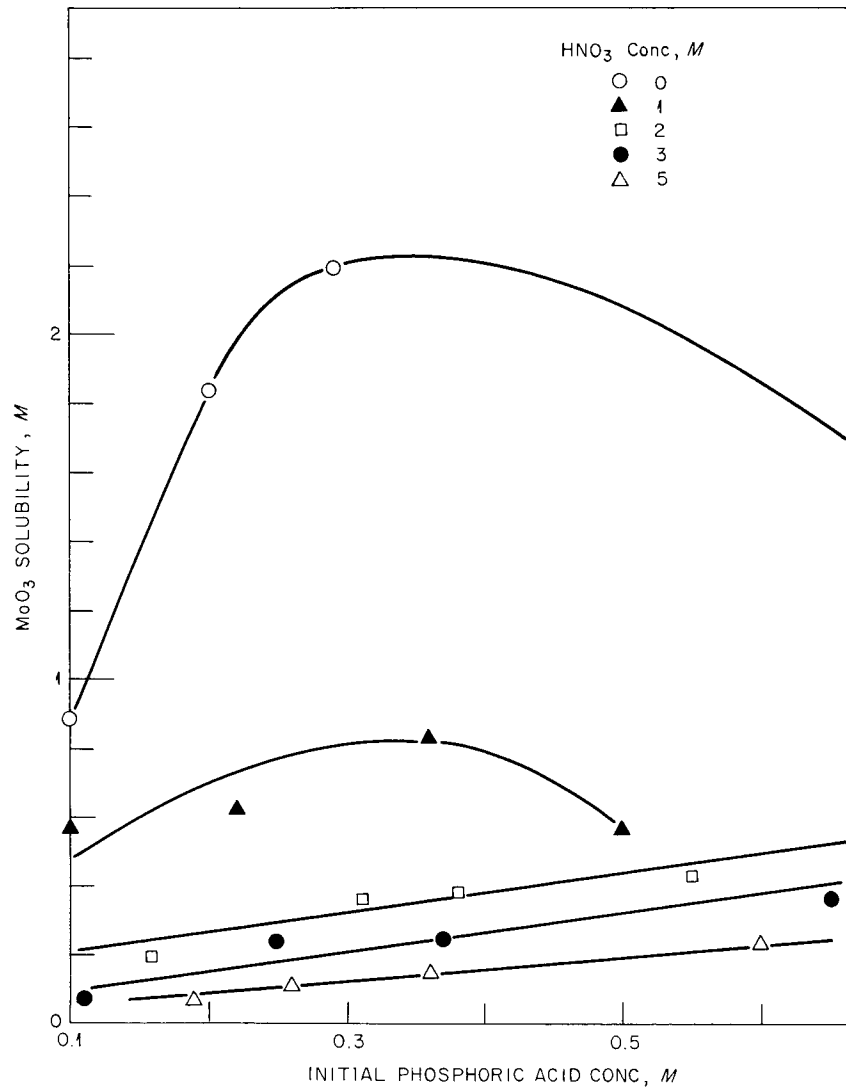


Fig. 12. Solubility of  $\text{MoO}_3$  in nitric acid—phosphoric acid solutions at  $26^\circ\text{C}$  as a function of initial acid concentrations.

Table 13. Solubility of MoO<sub>3</sub> in Nitric Acid-Phosphoric Acid Solutions at 26°C

Initial HNO <sub>3</sub> Conc, M	Initial H <sub>3</sub> PO <sub>4</sub> Conc, M	Mo Conc, M	Density of Saturated Solution, g/ml
0	0.1	0.893	1.1064
0	0.2	1.84	1.2293
0	0.29	2.20	1.2575
0	0.67	1.69	1.2318
1	0.1	0.577	1.1156
1	0.22	0.629	1.1391
1	0.36	0.837	1.1448
1	0.5	0.566	1.1656
2.16	0.16	0.204	1.0892
2.24	0.31	0.367	1.1170
2.40	0.38	0.391	1.1259
2.60	0.55	0.439	1.1394
3.20	0.11	0.079	1.1162
3.22	0.25	0.243	1.1235
3.33	0.37	0.254	1.1396
3.40	0.65	0.369	1.1632
5.04	0.19	0.076	1.1726
5.10	0.26	0.115	1.1808
5.10	0.36	0.148	1.1888
5.12	0.60	0.240	1.2079

Solubility of MoO<sub>3</sub> in Sodium Hydroxide Solutions. The solubility of sodium molybdate in sodium hydroxide solutions was measured to aid in determining the optimum conditions for storage of the basic waste solutions. At 26°C the solubility decreased from about 2.8 to 1.25 M as the sodium hydroxide concentration increased from 0 to about 4.5 M (Table 14). The solubility in water was estimated from data given by Seidell (22). The other solubility data are in excellent agreement with those reported by Bye (23).

### 3.4 Corrosion Studies

Titanium-45A, 304L stainless steel, Hastelloy F and Ni-o-nel appear suitable as materials of construction for processes involving nitric acid or nitric acid containing up to about 0.75 M Fe(NO<sub>3</sub>)<sub>3</sub>. Data given in

Table 14. Solubility of Sodium Molybdate in Sodium Hydroxide Solutions at 26°C

NaOH Conc, M	Na <sub>2</sub> MoO <sub>4</sub> Solubility, M	Density of Solution, g/ml
0 (ref. 22)	~2.8 (ref. 22)	~1.47 (ref. 22)
0.38	2.55	1.4395
0.76	2.44	1.4302
1.60	2.14	1.4151
2.72	1.86	1.3951
4.44	1.26	1.3635

Table 15 show that the rate of corrosion of titanium-45A was less than 0.1 mil/month in boiling 3 to 8 M HNO<sub>3</sub> containing 0.5 M Fe(NO<sub>3</sub>)<sub>3</sub>, 0.6 M UO<sub>2</sub>(NO<sub>3</sub>)<sub>2</sub>, and 0.02 M MoO<sub>3</sub> (24, 25). After 1000 hr in test, the general attack was negligible and there was no evidence of extensive localized attack. Work at HAPO (26) showed that corrosion of 304L stainless steel, Ni-o-nel, and vacuum-melted Hastelloy F was severe in solutions containing 1 M Fe(NO<sub>3</sub>)<sub>3</sub> but that rates were acceptably low when the ferric nitrate concentration was less than about 0.6 M.

Table 15. Corrosion of Titanium-45A in Boiling Nitric Acid Solutions Containing 0.5 M Fe(NO<sub>3</sub>)<sub>3</sub>, 0.6 M UO<sub>2</sub>(NO<sub>3</sub>)<sub>2</sub>, and 0.02 M MoO<sub>3</sub>

Duration of tests = 1000 hr

HNO <sub>3</sub> Conc, M	Corrosion Rate, mils/month <sup>a</sup>		
	Vapor	Interface	Solution
3.0	0.01	0.01	0.01
5.0	0.02	0.01	Gain
8.0	0.05	Gain	0.01

<sup>a</sup>Data from ref. 25.

#### 4.0 MISCELLANEOUS RELATED STUDIES

##### 4.1 Dissolution in Dilute Aqua Regia

Uranium-8.4% molybdenum alloy was readily dissolved in boiling 5 M  $\text{HNO}_3$  -- 2 M  $\text{HCl}$  to produce a solution which was stable only below 0.14 M uranium, showing that chloride was relatively ineffective as a complexing agent for molybdenum. Since the chloride must also be removed prior to solvent extraction, this method offers no advantages over direct dissolution in nitric acid.

##### 4.2 Other Potential Complexing Agents for Molybdenum

Scouting experiments showed that  $\text{Co(II)}$ ,  $\text{Cr(III)}$ ,  $\text{Ni(II)}$ ,  $\text{Al(III)}$ ,  $\text{Mg(II)}$ , and  $\text{As(V)}$  in 5 M  $\text{HNO}_3$  were less efficient than  $\text{Fe(III)}$  for complexing molybdenum.

##### 4.3 Reduction of Solvent Extraction Waste Volumes

Since solvent extraction waste volumes are expected to be high when ferric ion is used to complex molybdenum, scouting studies were performed to find means for producing concentrated waste solutions. One potential method involves neutralization of the iron-bearing waste solution with sodium hydroxide, allowing the ferric hydroxide to precipitate. If the supernatant solution is adjusted to about 1 M  $\text{NaOH}$ , the iron is essentially quantitatively precipitated with solubilization of 81 to 86% of the molybdenum. The molybdenum removal would probably have been higher if more refined experimental methods had been used. After the separation, the sodium molybdate solution can be concentrated by evaporation to about 2 M molybdenum (Table 14) and the ferric hydroxide dissolved in dilute nitric acid for storage or recycle as dissolvent.

In methods involving the use of nitric acid only, the solvent extraction raffinate will probably be about 3 M  $\text{HNO}_3$  containing up to 0.05 M molybdenum. These solutions, when boiled down to produce solutions boiling at about  $121^\circ\text{C}$ , result in the precipitation of most of the molybdenum and concentration of the nitric acid to 13 to 15 M. The residue from such a boildown is readily dissolved in dilute sodium hydroxide. Measurement of the temperature (which allows estimation of the nitric acid concentration) and the volume of the solution will define the amount of sodium hydroxide required to neutralize

the excess acid and dissolve the molybdic oxide. The caustic solutions would be concentrated before being admitted to waste tanks.

#### 4.4 Decladding of Zirconium-clad Fuels

Fuels like the Detroit Edison core (3) are composed of uranium-molybdenum alloy clad in zirconium or Zircaloy-2. The possibility of removing the cladding with boiling ammonium fluoride or  $\text{NH}_4\text{F}-\text{NH}_4\text{NO}_3$  solutions (Zirflex process (27)) was cursorily investigated. In boiling 6 M  $\text{NH}_4\text{F}$  or 6 M  $\text{NH}_4\text{F}-1$  M  $\text{NH}_4\text{NO}_3$  the rate of attack on alloys containing either 3 or 10% molybdenum was too great for a decladding procedure to be considered:

<u>Mo in Alloy, %</u>	<u>Solution</u>	<u>Rate of Attack,</u> <u>mg min<sup>-1</sup> cm<sup>-2</sup>,</u>
3	6 M $\text{NH}_4\text{F}$	2.5
3	6 M $\text{NH}_4\text{F}-1$ M $\text{NH}_4\text{NO}_3$	1.7
10	6 M $\text{NH}_4\text{F}$	2.1
10	6 M $\text{NH}_4\text{F}-1$ M $\text{NH}_4\text{NO}_3$	1.5

There is a possibility, however, that hydrofluoric acid could be used to declad U-10% Mo alloy fuels. Initial rates of dissolution of this alloy in 5-9 M HF were generally less than 0.2 mg min<sup>-1</sup> cm<sup>-2</sup> (28).

#### 5.0 REFERENCES

1. E. R. Irish and W. H. Reas, "The Purex Process: A Solvent Extraction Reprocessing Method for Irradiated Uranium," in "Symposium on the Reprocessing of Irradiated Fuels, Brussels, Belgium, May 20-25, 1957," TID-7534, p. 83.
2. J. W. Ullmann, "Reactor Data Sheet, Consumers Public Power," Aug. 22, 1960, personal compilation.
3. J. W. Ullmann, "Reactor Data Sheet, PRDC," June 29, 1959, personal compilation.
4. "Proceedings of the AEC Symposium for Chemical Processing of Irradiated Fuels from Power, Test, and Research Reactors, Richland, Washington, October 20-21, 1959," TID-7583.

- a) C. D. Watson, J. B. Adams, G. K. Ellis, G. A. West, F. L. Hannon, W. F. Schaffer, and B. B. Klima, "Mechanical Processing of Spent Reactor Fuel at Oak Ridge National Laboratory," p. 307.
- b) R. E. Blanco, L. M. Ferris, J. R. Flanary, F. G. Kitts, R. H. Rainey, and J. T. Roberts, "Chemical Processing of Power and Research Reactor Fuels at Oak Ridge National Laboratory," p. 234.
5. R. E. Blanco, "Processing of Power Reactor Fuels," Nucl. Sci. Eng., 1:409 (1956).
6. V. B. Shevchenko, N. S. Povitsky, and A. S. Salovkin, "Treatment of Irradiated Fuel Elements," P/2182, Proc. 2nd UN Internatl. Conf. on Peaceful Uses of Atomic Energy, Geneva, 1958, Vol. 17, p. 46, UN, New York.
7. W. W. Schulz, "Reprocessing Uranium-Molybdenum Alloy Fuels: Dissolution in Concentrated Nitric Acid," HW-64432 (March 17, 1960).
8. W. W. Schulz and E. M. Duke, "Reprocessing of Low-Enrichment Uranium-Molybdenum Alloy Fuels," HW-62086 (Sept. 15, 1959).
9. E. S. Occhipinti, "Reprocessing of Power Reactor Fuels, Third Quarterly Progress Report, April 1 - July 1, 1958," DP-318 (December, 1958).
10. T. A. Gens, "An Oxyhydrochlorination Process for Preparing Uranium-Molybdenum Reactor Fuels for Solvent Extraction," ORNL-3019 (in press).
11. R. E. Blanco, "Dissolution and Feed Adjustment," in "Symposium on the Reprocessing of Irradiated Fuels, Brussels, Belgium, May 20-25, 1957," TID-7534, p. 22.
12. K. S. Warren, "Survey of Potential Vapor-phase Explosions in Darex and Sulfex Processes," ORNL-2937 (Dec. 27, 1960).
13. L. H. Meyer, "Reprocessing of Power Reactor Fuels, Quarterly Progress Report, Oct. 1, 1959-Jan. 1, 1960," DP-479 (March, 1960).
14. D. H. Killeffer and A. Linz, "Molybdenum Compounds," Interscience, New York, 1952, p. 78.
15. A. F. Wells, "Structural Inorganic Chemistry," 2d ed, Oxford University Press, New York, 1950, p. 348.
16. H. J. Emeléus and J. S. Anderson, "Modern Aspects of Inorganic Chemistry," 2d ed, Van Nostrand, New York, 1952, p. 206.

17. M. L. Freedman, J. Am. Chem. Soc., 81: 3834 (1959).
18. L. M. Ferris, "The Solubility of Molybdc Oxide in Nitric Acid, Nitric Acid—Uranyl Nitrate, and Nitric Acid—Ferric Nitrate Solutions," ORNL-CF-60-2-94 (Rev. 1), paper submitted for publication.
19. P. Faugeras, Centre D'Etudes Nucléaires de Fontenay-Aux-Roses, France, personal communication, Sept. 22, 1960.
20. P. Cannon, J. Inorg. Nucl. Chem., 11: 124 (1959).
21. "Chemical Technology Division Annual Progress Report for Period Ending August 31, 1960," ORNL-2993, p. 50.
22. A. Seidell, "Solubilities of Inorganic and Metal Organic Compounds," 3rd ed, Van Nostrand, New York, 1940, p. 1272.
23. J. Bye, Bull. soc. chim., 10(5): 239 (1943).
24. Letter to F. L. Culler from W. E. Clark, ORNL, "PRFR Corrosion Status Summary and Proposed Program for Remainder of FY-1961," ORNL-CF-60-8-122 (Sept. 8, 1960).
25. Letter to W. E. Clark, ORNL, from L. Rice and D. N. Hess, ORNL, "Corrosion Program in Support of Power Reactor Fuel Reprocessing Program," Sept. 30, 1960.
26. R. F. Maness, "Power Reactor Fuels Reprocessing: Progress Report on Corrosion Aspects," HW-61662 (September 1959).
27. L. M. Ferris, "Zirflex Process for PWR Blanket Fuel. II. Revised Flowsheet," ORNL-2940 (Oct. 26, 1960).
28. W. E. Clark and A. H. Kibbey, "Hydrofluoric Acid Decladding of Zirconium-Clad Power Reactor Fuel Elements," ORNL-2460 (Oct. 15, 1958).

NOTEBOOK REFERENCES

1. J. F. Land, Unclassified Notebooks, A-2020, A-1765, and A-2148.
2. L. M. Ferris, Unclassified Notebook, A-1460.





ORNL-3068  
UC-10 - Chemical Separations  
Processes for Plutonium and Uranium  
TID-4500 (16th ed.)

INTERNAL DISTRIBUTION

- |   |                                 |
|---|---------------------------------|
| 1. Biology Library  | 52. M. T. Kelley                |
| 2-3. Central Research Library                                   | 53. J. F. Land                  |
| 4. Reactor Division Library                                     | 54. J. A. Lane                  |
| 5. ORNL - Y-12 Technical Library,<br>Document Reference Section | 55. T. A. Lincoln               |
| 6-25. Laboratory Records Department                             | 56. S. C. Lind                  |
| 26. Laboratory Records, ORNL R.C.                               | 57. R. S. Livingston            |
| 27. E. D. Arnold  | 58. J. T. Long                  |
| 28. R. E. Blanco  | 59. K. Z. Morgan                |
| 29. G. E. Boyd  | 60. J. P. Murray (K-25)         |
| 30. J. C. Bresee  | 61. M. L. Nelson                |
| 31. K. B. Brown   | 62. R. H. Rainey                |
| 32. F. R. Bruce   | 63. H. E. Seagren               |
| 33. C. E. Center  | 64. M. J. Skinner               |
| 34-35. F. L. Culler   | 65. J. A. Swartout              |
| 36. W. K. Eister  | 66. E. H. Taylor                |
| 37. D. E. Ferguson  | 67. J. W. Ullmann               |
| 38. L. M. Ferris  | 68. W. E. Unger                 |
| 39. J. R. Flanary   | 69. C. D. Watson                |
| 40. J. H. Frye, Jr.   | 70. A. M. Weinberg              |
| 41. J. H. Gillette  | 71. M. E. Whatley               |
| 42. H. E. Goeller   | 72. C. E. Winters               |
| 43. A. T. Gresky  | 73. R. G. Wymer                 |
| 44. W. R. Grimes  | 74. J. W. Youngblood            |
| 45. C. E. Guthrie   | 75. D. L. Katz (consultant)     |
| 46. C. W. Hancher   | 76. C. E. Larson (consultant)   |
| 47. C. S. Harrill   | 77. I. Perlman (consultant)     |
| 48. A. Hollaender   | 78. J. H. Rushton (consultant)  |
| 49. A. S. Householder   | 79. H. Worthington (consultant) |
| 50. R. G. Jordan (Y-12)   | 80. T. H. Pigford (consultant)  |
| 51. W. H. Jordan  |                                 |

EXTERNAL DISTRIBUTION

81. Division of Research and Development, AEC, ORO  
82-594. Given distribution as shown in TID-4500 (16th ed.) under Chemical Separations Processes for Plutonium and Uranium (75 copies - OTS)  
595. W. W. Schulz, Hanford Atomic Products Operation