

LOCKHEED MARTIN ENERGY RESEARCH LIBRARIES



3 4456 0515558 7

ORNL-4357
UC-4 - Chemistry

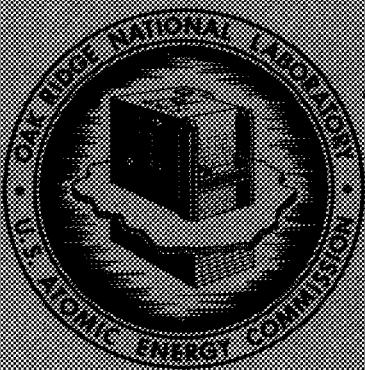
CURIUM DATA SHEETS

OAK RIDGE NATIONAL LABORATORY
CENTRAL RESEARCH LIBRARY
CIRCULATION SECTION
SECOND FLOOR 175

LIBRARY LOAN COPY

DO NOT TRANSFER TO ANOTHER PERSON
If you wish someone else to see this report, send in
name with report and the library will arrange a loan.

ORNL-143-6-27



OAK RIDGE NATIONAL LABORATORY

operated by

UNION CARBIDE CORPORATION

for the

U.S. ATOMIC ENERGY COMMISSION

Printed in the United States of America. Available from Clearinghouse for Federal

Scientific and Technical Information, National Bureau of Standards.

U.S. Department of Commerce, Springfield, Virginia 22151

Price: Printed Copy \$2.00; Microfiche \$0.65

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 liability 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.

ORNL-4357

Contract No. W-7405-eng-26

ISOTOPES DEVELOPMENT CENTER

CURIUM DATA SHEETS

S. J. Rimshaw
E. E. Ketchen

Isotopes Division

JANUARY 1969

OAK RIDGE NATIONAL LABORATORY
Oak Ridge, Tennessee
operated by
UNION CARBIDE CORPORATION
for the
U.S. ATOMIC ENERGY COMMISSION

LOCKHEED MARTIN ENERGY RESEARCH LIBRARIES



3 4456 0515558 7

11

CONTENTS

	<u>Page</u>
I. CURIUM-244	1
A. CURIUM-244 METAL	1
1. Composition	1
2. Specific Power	2
3. Radiation	2
4. Critical Mass	5
5. Compatibility with Materials of Containment	6
6. Thermophysical Properties	6
7. Mechanical Properties	7
8. Chemical Properties	8
9. Biological Tolerances	8
10. Shielding Data	9
B. CURIUM SESQUIOXIDE (Cm_2O_3)	13
1. Composition	13
2. Specific Power	13
3. Radiation	13
4. Critical Mass	13
5. Compatibility with Materials of Containment	13
6. Thermophysical Properties	13
7. Mechanical Properties	17
8. Chemical Properties	17
9. Biological Tolerances	18
10. Shielding Data	18
C. CURIUM OXYSULFIDE ($\text{Cm}_2\text{O}_2\text{S}$)	19
1. Composition	19
2. Specific Power	19
3. Radiation	19
4. Critical Mass	19
5. Compatibility with Materials of Containment	19
6. Thermophysical Properties	19
7. Mechanical Properties	21
8. Chemical Properties	21
9. Biological Tolerances	21
10. Shielding Data	21
D. CURIUM FLUORIDE (CmF_3)	22
1. Composition	22
2. Specific Power	22
3. Radiation	22
4. Critical Mass	22
5. Compatibility with Materials of Containment	22
6. Thermophysical Properties	22
7. Mechanical Properties	24
8. Chemical Properties	24
9. Biological Tolerances	24
10. Shielding Data	24

	<u>Page</u>
III. CURIUM-2 ⁴²	25
A. CURIUM-2 ⁴² OXIDE CERMET	25
1. Composition	25
2. Specific Power	26
3. Radiation	26
4. Critical Mass	30
5. Compatibility with Materials of Containment	30
6. Thermophysical Properties	30
7. Mechanical Properties	30
8. Chemical Properties	30
9. Biological Tolerances	30
10. Shielding Data	31
III. REFERENCES FOR CURIUM	34

CURIUM DATA SHEETSREFERENCE
COLUMNI. CURIUM-244A. CURIUM-244 METAL

Half-life: 18.1 y 1, 2

1. Composition

a. Radionuclidic abundance

3

The anticipated isotopic composition of ^{244}Cm product resulting from a large-scale production program is represented by the actual isotopic composition of the first product batch resulting from the 1968-1969 SRL campaign shown below. The mass spectrographic analysis yielding these data was performed 18 months after reactor discharge.

Isotopic Composition of Curium Product	
Isotopic mass	% Abundance
242	0.043
243	0.024
244	95.08
245	0.729
246	4.02
247	0.066
248	0.038

b. Radiochemical purity

The ^{244}Cm product will contain small amounts of fission products formed in the irradiation process and not removed in the chemical separations process. From the predicted performance of the chemical process and the calculated quantities of fission products associated with the curium after irradiation, the gamma activities of the fission-product impurities can be calculated and are tabulated in the table in Section I.A.3.c. (page 3) in the column labeled "From Impurities." Almost all of the gamma rays above 1 Mev are from ^{144}Pr and $^{152-154}\text{Eu}$, and those below 1 Mev are primarily from ^{95}Zr - ^{95}Nb , ^{144}Ce , ^{106}Rh , and ^{103}Ru - ^{103}Rh .

Fission products will also be formed by the spontaneous fission of ^{244}Cm and ^{252}Cf , which is present at an estimated maximum of 4×10^{-5} wt % of the ^{244}Cm . The gamma activities from these two isotopes are tabulated under the columns labeled "From Fission Products of ^{244}Cm and ^{252}Cf ."

Americium-243 and ^{252}Cf which are formed in the irradiation process are removed during chemical processing except for an estimated 0.5 wt % ^{243}Am and $<4 \times 10^{-5}$ wt % ^{252}Cf in the ^{244}Cm product.

CURIUM-2⁴⁴ METAL

c. Chemical purity

The following elemental impurity limits have been set as objectives for the ²⁴⁴Cm process by Savannah River Plant during the CY 1968-1969 campaign:

<u>Elemental impurities</u>	<u>Maximum wt % of Cm</u>
Li + Na + K (total)	0.1
Si + Zn (total)	0.1
Fe + Cr + Ni (total)	0.2
Any other single impurity (except Am, Zr)	0.2
Total of all impurities (except Am, Zr)	1.0
Total of all impurities (including Am, Zr)	3.0

Plutonium-240 as a decay product of ²⁴⁴Cm will be present in amounts increasing with time following chemical processing.

 2. Specific Power

- a. 2.84 w/g of ²⁴⁴Cm (100% ²⁴⁴Cm isotope) 4, 5, 6
- 2.70 w/g of pure curium element (95.08% ²⁴⁴Cm isotope)
- 2.62 w/g of curium product (97% chemically pure)
- b. Specific power values are based on 81.3 Ci/g of ²⁴⁴Cm (100%) and 34.95 w/kCi of ²⁴⁴Cm.

3. Radiation

a. Alpha particles

Nuclide	Max E, Mev	Av E, Mev	Abundance, %	w/kCi	Particles w ⁻¹ sec ⁻¹
²⁴⁴ Cm	5.801	5.801	76.7	{ 34.3	0.825 x 10 ¹²
	5.759	5.759	23.3		0.252 x 10 ¹²

The volume of helium from alpha decay as a function of decay time is given in the following table.

Volume of helium, ^a cm ³ of He per g of ²⁴⁴ Cm	Time Years	Time Half-lives
6.15	1.8	0.1
11.8	3.6	0.2
17.3	5.4	0.3
22.1	7.2	0.4
26.9	9.1	0.5
31.2	10.9	0.6
35.3	12.7	0.7
39.1	14.5	0.8

CURIUM-244 METAL

continued

cm ³ of He per g of ²⁴⁴ Cm	Time	
	Years	Half-lives
42.6	16.3	0.9
45.9	18.1	1.0
59.3	27.1	1.5
68.9	36.2	2.0
80.3	54.3	3.0
86.1	72.4	4.0
88.9	90.5	5.0
91.7	181.0	10.0

^aStandard conditions

b. Beta particles

None

c. Gamma

The significant gamma emissions of product-grade ²⁴⁴Cm are listed in the following table. The gamma emissions result from the following:

- (1) Decay of curium nuclides and of ²⁴³Am (based on 0.02 wt % ²⁴²Cm, 0.002 wt % ²⁴³Cm, 95.3 wt % ²⁴⁴Cm, and 0.5 wt % ²⁴³Am).
- (2) Spontaneous fission of ²⁴⁴Cm and ²⁵²Cf (based on a maximum of 4×10^{-5} wt % ²⁵²Cf).
- (3) Decay of fission products resulting from the spontaneous fission of ²⁴⁴Cm and ²⁵²Cf.
- (4) Decay of fission products formed in the irradiation process and not removed in the chemical separation process.

Significant Gamma Rays of Curium-244 Product

Energy, Mev	Abundance, photons/(sec)(w ²⁴⁴ Cm product)								Total	
	From decay of nuclides				From spontaneous fissions		From fission products of			
	²⁴² Cm	²⁴³ Cm	²⁴⁴ Cm	²⁴³ Am	²⁴⁴ Cm	²⁵² Cf	²⁴⁴ Cm	²⁵² Cf		
0.0-0.5	4.1×10^6	3.6×10^6	2.5×10^6	1.0×10^7	4.1×10^6	4.9×10^5	2.2×10^6	1.9×10^5	2.2×10^7	
0.5-1.0	3.0×10^4	—	2.2×10^6	—	2.6×10^6	2.5×10^5	6.7×10^6	6.0×10^5	7.5×10^6	
1.0-2.0	9.0×10^2	—	—	—	1.9×10^3	1.8×10^5	1.5×10^6	1.3×10^5	1.1×10^6	
2.0-3.0	—	—	—	—	6.0×10^5	4.9×10^4	6.0×10^5	5.2×10^2	3.7×10^5	
3.0-4.0	—	—	—	—	1.8×10^5	1.3×10^4	—	—	—	
4.0-5.0	—	—	—	—	6.0×10^4	3.7×10^3	—	—	6.4×10^4	
5.0-6.0	—	—	—	—	3.3×10^4	1.0×10^3	—	—	3.3×10^4	
6.0-7.0	—	—	—	—	5.6×10^3	1.5×10^2	—	—	5.6×10^3	

d. Bremsstrahlung

None

CURIUM-244 METAL

e. Neutrons

4.19×10^6 neutrons $\text{sec}^{-1} \text{w}^{-1}$ of ^{244}Cm from spontaneous fission (half-life for spontaneous fission is 1.4×10^7 y).

7, 8, 9

1.51×10^5 neutrons $\text{sec}^{-1} \text{w}^{-1}$ of ^{244}Cm from (α, n) reaction on oxygen in Cm_2O_3 .

7

3.5×10^5 neutrons $\text{sec}^{-1} \text{w}^{-1}$ of ^{244}Cm from estimated maximum of ^{252}Cf content of 4×10^{-5} wt %.

The energy distribution of spontaneous fission neutrons from ^{244}Cm is given in the following table. 4

Energy, Mev	Abundance, neutrons $\text{sec}^{-1} \text{w}^{-1}$ of ^{244}Cm
0.3-0.4	1.51×10^5
0.4-0.6	3.13×10^5
0.6-0.8	3.20×10^5
0.8-1.0	2.77×10^5
1.0-1.2	2.84×10^5
1.2-1.4	2.80×10^5
1.4-1.6	2.44×10^5
1.6-1.8	2.19×10^5
1.8-2.0	1.98×10^5
2.0-2.2	1.80×10^5
2.2-2.4	1.65×10^5
2.4-2.6	1.58×10^5
2.6-2.8	1.30×10^5
2.8-3.0	1.08×10^5
3.0-3.2	1.01×10^5
3.2-3.4	0.97×10^5
3.4-3.6	0.93×10^5
3.6-3.8	0.75×10^5
3.8-4.0	0.79×10^5
4.0-4.4	1.04×10^5
4.4-4.8	0.86×10^5
4.8-5.2	0.65×10^5
5.2-5.6	0.50×10^5
5.6-6.0	0.40×10^5
6.0-6.4	2.95×10^4
6.4-6.8	2.12×10^4
6.8-7.2	1.47×10^4
7.2-7.6	1.12×10^4
7.6-8.0	0.90×10^4
8.0-8.8	1.01×10^4
8.8-9.6	2.95×10^3
9.6-10.4	3.1×10^3
10.4-11.2	2.05×10^3
11.2-12.8	1.40×10^3

CURIUM-244 METAL

The energy distribution of neutrons occurring as a result of the collision of fast alpha particles from ^{244}Cm decay with oxygen atoms in Cm_2O_3 is given in the following table.

4

Neutrons from (α, n) Reactions with Oxygen

Energy, Mev	Abundance, neutrons sec $^{-1}$ w $^{-1}$ of ^{244}Cm
0.2	1.62×10^2
0.4	0.68×10^3
0.6	0.83×10^3
0.8	1.69×10^3
1.0	2.70×10^3
1.2	0.43×10^4
1.4	0.61×10^4
1.6	0.83×10^4
1.8	1.01×10^4
2.0	1.19×10^4
2.2	1.33×10^4
2.4	1.40×10^4
2.6	1.40×10^4
2.8	1.37×10^4
3.0	1.22×10^4
3.2	1.01×10^4
3.4	0.79×10^4
3.6	0.50×10^4
3.8	3.02×10^3
4.0	2.01×10^3
4.2	1.37×10^3
4.4	0.86×10^3
4.6	0.72×10^2

4. Critical Mass

The critical mass of unreflected and reflected spheres of ^{244}Cm and $^{244}\text{Cm}_2\text{O}_3$ have been calculated by C. W. Craven, Jr., at ORNL using the cross-section data available as of November 1965. The results, shown in the following table, agree within 10% of the results obtained in the replacement experiment at Los Alamos. The critical mass equations used are

10

$$\text{Cm: } M_C = 2450/\rho^2 \text{ (kg)}$$

$$\text{Cm}_2\text{O}_3: M_C = 2370/\rho^2 \text{ (kg)}$$

CURIUM-2⁴⁴ METALREFERENCE
COLUMNCalculated Critical Mass of ^{244}Cm and $^{244}\text{Cm}_2\text{O}_3$

Mixture	Core Density, g/cm ³	Reflector		Critical mass, kg	Critical radius, cm
		Mixture	Thickness, cm		
Cm ^a	14.0	Bare	—	12.5	5.97
Cm ₂ O ₃ ^a	10.60	Bare	—	21.1	7.8031
Cm ₂ O ₃	9.01	Bare	—	29.2	9.1803
Cm ₂ O ₃	10.60	Au-H ₂ O	4.0-15.0	11.9	6.4540
Cm ₂ O ₃	10.60	Au-H ₂ O	2.0-7.5	13.5	6.7188
Cm ₂ O ₃	10.60	Au-H ₂ O	0.5-2.0	16.5	7.1966

^aAssumed composition: 98.07 wt % ^{244}Cm and 1.93 wt % ^{241}Pu .

5. Compatibility with Materials of Containment6. Thermophysical Properties

a. Density

13.51 g/cm³ of metal 11

b. Coefficient of thermal expansion 12 (Am)

$$\alpha_a = 7.5 \times 10^{-6}/^\circ\text{C}$$

$$\alpha_c = 6.2 \times 10^{-6}/^\circ\text{C}$$

c. Specific heat and enthalpy

(1) Specific heat

0.0270 cal g⁻¹ °C⁻¹ 13 (Nd)

(2) Enthalpy in cal/mole

$$H_t - H_0 = 6.48 + 3.53 \times 10^{-3} t^2 + 1.49 \times 10^{-6} t^3 \quad 14 \text{ (Nd)}$$

(Nd temperature is 0-400°C and t is in °C)

d. Temperatures of phase transformations

(1) Melting point

1340 ± 40°C 15

1350 ± 60°C 16

(2) Boiling point

3525 ± 130°C 17

CURIUM- 244 METAL

e.	Latent heats of phase transformations	
	ΔH fusion 3.2 kcal/mole (Richard's rule)	17
	$\Delta H_{1600^{\circ}\text{K}}$ vaporization 82.6 ± 2.7 kcal/mole	17
f.	Vapor pressure	
	$\log_{10} P = 4.74 \pm 0.37 - [(18,060 \pm 590)/T]$ (T is in $^{\circ}\text{K}$ and P is in atm)	17
g.	Thermal conductivity	
	0.021 cal cm^{-1} sec^{-1} $^{\circ}\text{C}^{-1}$ at 26°C	18 (Cd)
h.	Thermal diffusivity	
	0.575 cm^2/sec at 26°C	
	This value was calculated by dividing the product of the specific heat and density into the thermal conductivity.	
i.	Viscosity	
	2.5 centipoises at 1340°C	19
j.	Surface tension	
	500 dyn/cm	20
k.	Total hemispherical emittance	
	0.37 at 89°C	21 (Pu)
l.	Spectral emissivity	
	0.55	22 (Er)
	A higher value of 0.9 can be assumed if the metal surface is oxidized or if impurities are present.	
m.	Crystallography	
	Double hexagonal close packed	11
	$a = 3.496 \pm 0.003 \text{ \AA}$ $c = 11.331 \pm 0.005 \text{ \AA}$	
n.	Solubilities	
	Reacts strongly with water	23
o.	Diffusion rates	
7.	<u>Mechanical Properties</u>	
a.	Hardness	
	Vicker's 97.7	24 (Dy)

CURIUM-244 METAL

b. Crush strength

7741 kg/cm²

24 (Tm)

8. Chemical Properties

a. Heat and free energy of formation, entropy

(1) Heat of formation

Zero - by definition of standard state

(2) Free energy of formation

Zero - by definition of standard state

(3) Entropy

S°₂₉₈ = 18 cal °C⁻¹ mole⁻¹

15

b. Chemical reactions and reaction rates

(1) Oxygen at room temperature - slow

11

(2) Oxygen at elevated temperature - fast

11

(3) Nitrogen at room temperature - very slow

25

(4) Nitrogen at elevated temperature - slow

25

(5) Water at room temperature - fast

23

(6) Hydrogen at room temperature - slow

23

(7) Hydrogen at elevated temperature - fast

23

9. Biological Tolerances

Maximum permissible body burdens and maximum permissible concentrations of ²⁴⁴Cm in air and water are shown in the following table.

26

 Maximum Permissible Body Burdens and Maximum Permissible Concentrations
 for Radionuclides in Air and in Water for Occupational Exposure²⁶

Radionuclide and type of decay	Organ of reference (critical organ underscored)	Max permissible burden in total body, q(uCi)	Max permissible concentrations, uCi/cm ³			
			For 40-hr week		For 168-hr week	
			Water	Air	Water	Air
²⁴⁴ Cm (α, γ) (Sol)	Bone	0.1	2 × 10 ⁻⁴	9 × 10 ⁻¹²	7 × 10 ⁻⁵	3 × 10 ⁻¹²
	Liver	0.2	3 × 10 ⁻⁴	10 ⁻¹¹	9 × 10 ⁻⁵	4 × 10 ⁻¹²
	Kidney	0.2	4 × 10 ⁻⁴	2 × 10 ⁻¹¹	10 ⁻⁴	6 × 10 ⁻¹²
	Total Body	0.3	6 × 10 ⁻⁴	3 × 10 ⁻¹¹	2 × 10 ⁻⁴	9 × 10 ⁻¹²
	GI (LLI) ^a		8 × 10 ⁻⁴	2 × 10 ⁻⁷	3 × 10 ⁻⁴	6 × 10 ⁻⁸
(Insol)	Lung			10 ⁻¹⁰		3 × 10 ⁻¹¹
	GI (LLI) ^a		8 × 10 ⁻⁴	10 ⁻⁷	3 × 10 ⁻⁴	5 × 10 ⁻⁸

^aThe abbreviations GI and LLI refer to the gastrointestinal tract and lower large intestine, respectively.

CURIUM-244 METALREFERENCE
COLUMN10. Shielding Data

Gamma dose rates with water, iron, lead, and uranium shielding are given in Figs. 1-5 for ^{244}Cm power sources of 100, 200, 500, 1000, 2000, 5000, 10,000, and 20,000 w. Neutron dose rates with water shielding are given in Fig. 6. Neutron dose rates on shielding with Be, CH, CH_2 , or LiH can be estimated by using Fig. 6 in conjunction with Fig. 7.

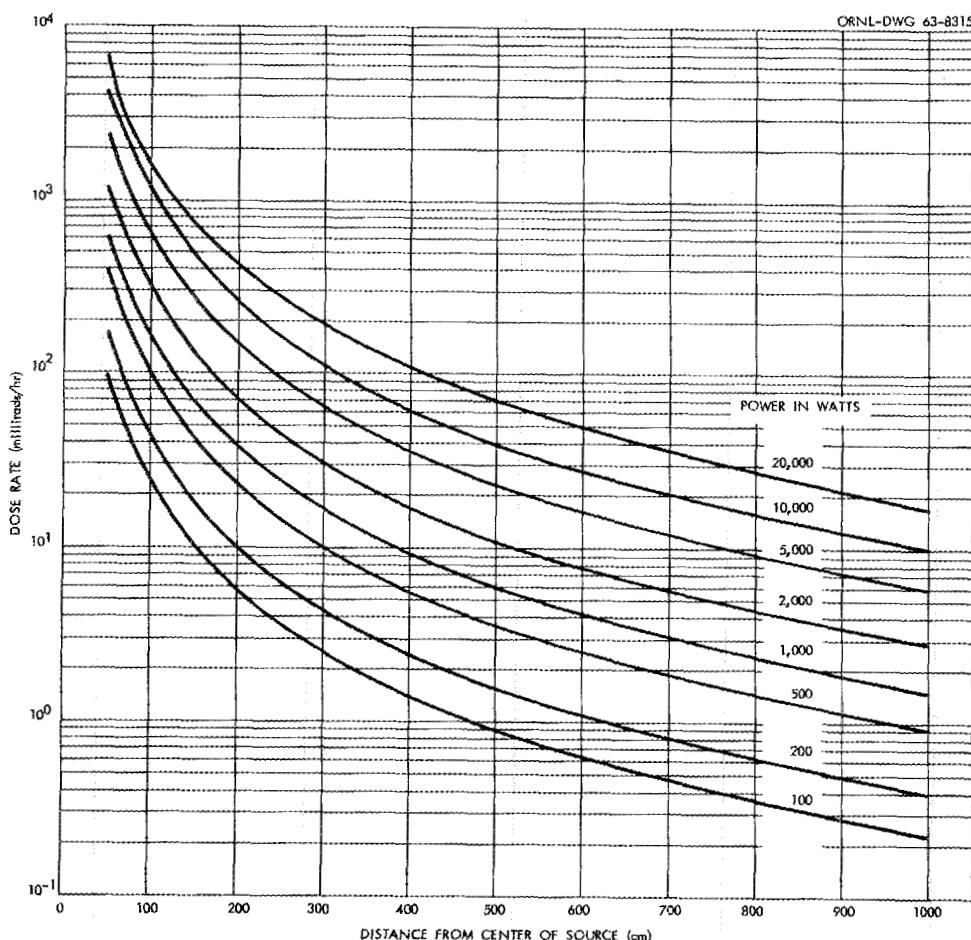


Fig. 1. Gamma Dose Rates from Unshielded Isotopic Power Sources of Curium-244 as a Function of Distance from Center of Source.

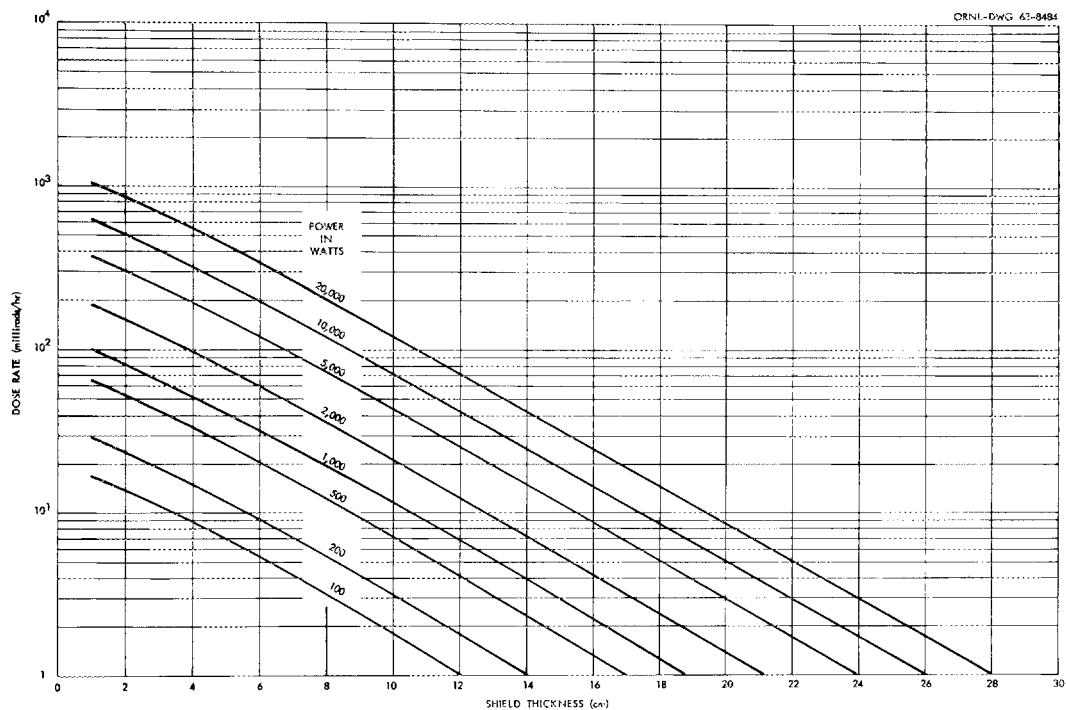


Fig. 2. Gamma Dose Rates from Iron-Shielded Isotopic Power Sources of Curium-244. Center of source to dose point separation distance is 100 cm.

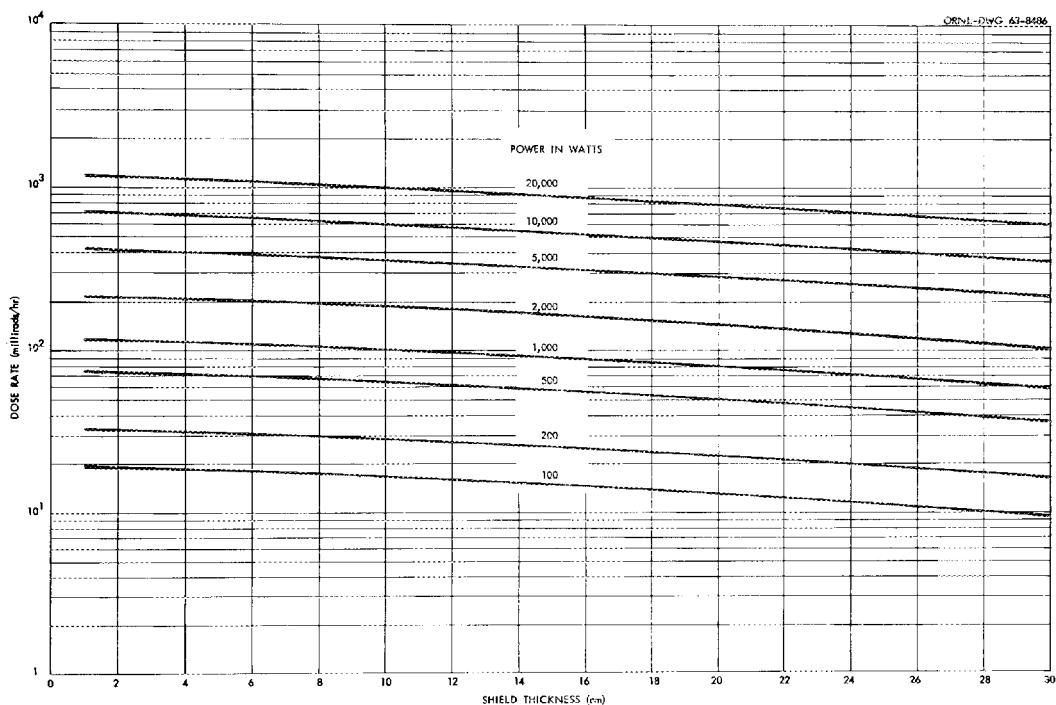


Fig. 3. Gamma Dose Rates from Water-Shielded Isotopic Power Sources of Curium-244. Center of source to dose point separation distance is 100 cm.

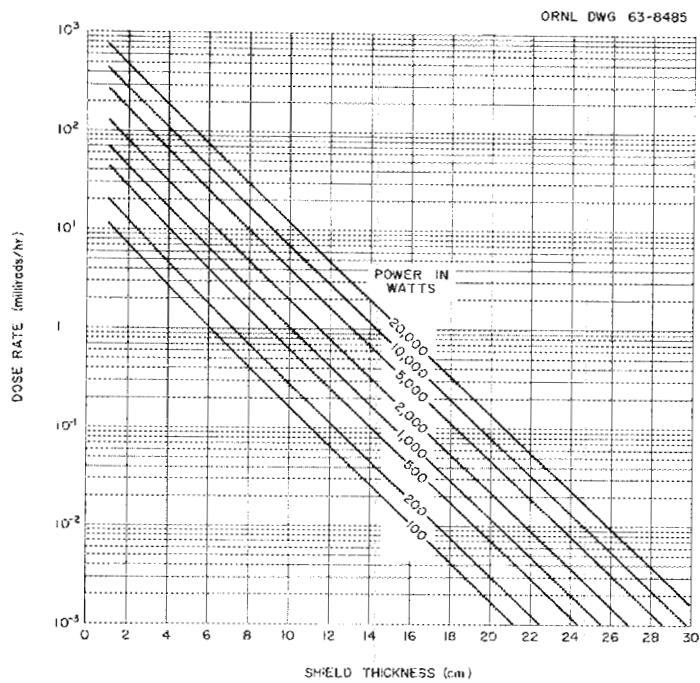


Fig. 4. Gamma Dose Rates from Lead-Shielded Isotopic Power Sources of Curium-244. Center of source to dose point separation distance is 100 cm.

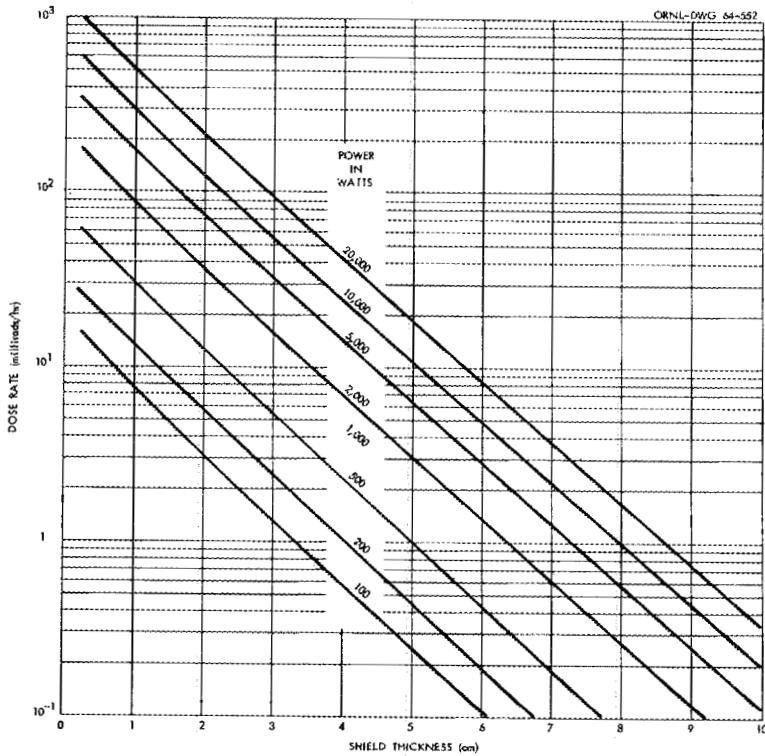


Fig. 5. Gamma Dose Rates from Uranium-Shielded Isotopic Power Sources of Curium-244. Center of source to dose point separation distance is 100 cm.

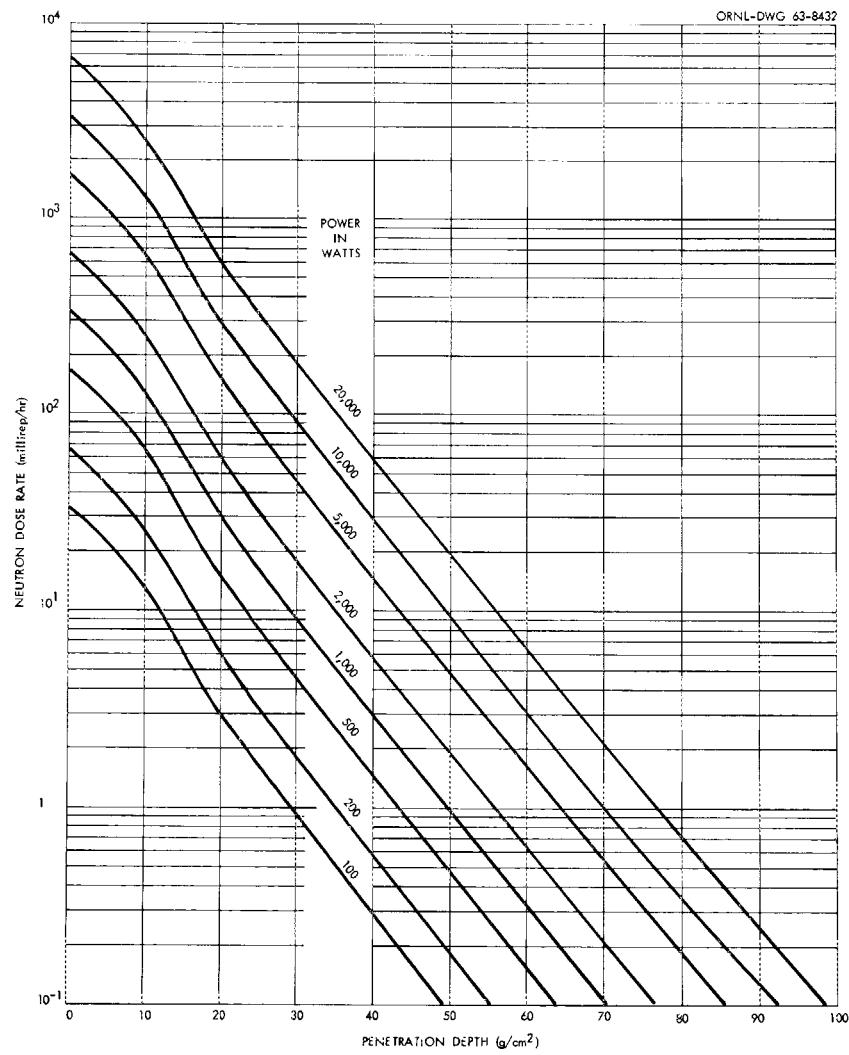


Fig. 6. Neutron Dose Rates from Water-Shielded Isotopic Power Sources of Curium-244 as a Function of Penetration Depth of Shielding Material. Center of source to dose point separation distance is 100 cm.

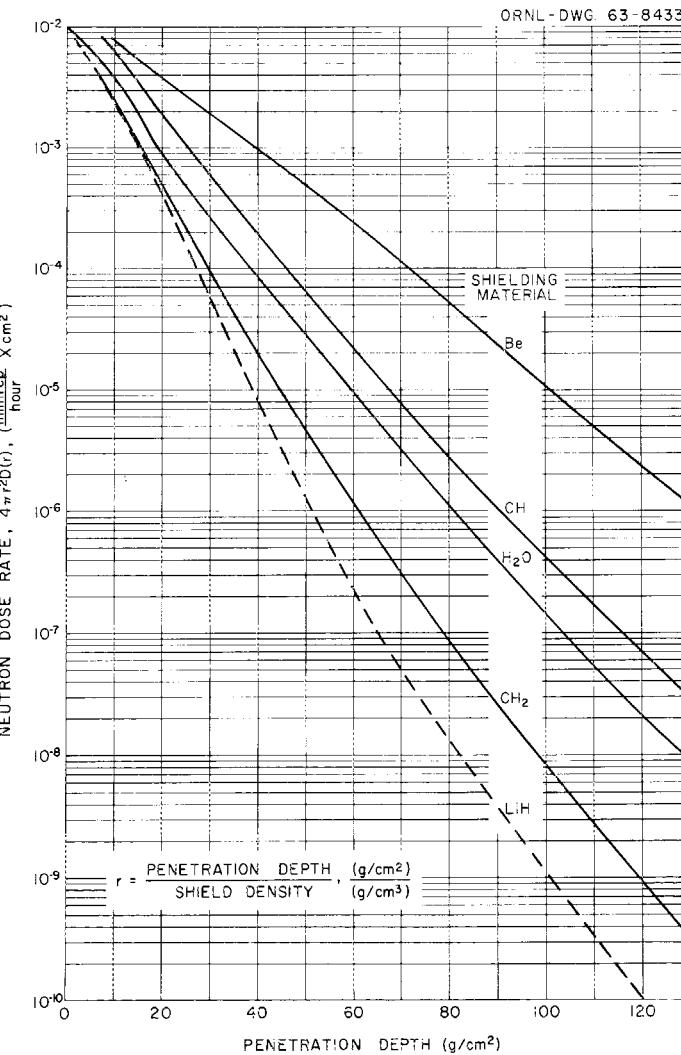


Fig. 7. Fast Neutron Dose Rate (Multiplied by $4\pi r^2$) in Various Materials as a Function of Penetration Depth from a Unit Point Isotropic Fission Source.

B. CURIUM SESQUIOXIDE (Cm_2O_3)

1. Composition

- a. Radionuclidic abundance

See Section I.A.1.a.

- b. Radiochemical purity

See Section I.A.1.b.

- c. Chemical purity

See Section I.A.1.c.

- d. Phase composition of various curium oxides

27-32

$\text{CmO}_{2.00}$

$\text{CmO}_{1.52-1.62}$ (C-type)

$\text{CmO}_{1.83}$ (Cm_6O_{11})

$\text{CmO}_{1.50}$ (B- and A-type)

$\text{CmO}_{1.71}$ (Cm_7O_{12})

2. Specific Power

- a. 2.46 w/g of 100% pure Cm_2O_3 (86.5% ^{244}Cm isotope)

4, 5, 6

2.38 w/g of 97% pure Cm_2O_3 (83.9% ^{244}Cm isotope)

- b. Specific power values are based on 81.3 Ci/g of ^{244}Cm (100%) isotope and 34.95 w/kCi of ^{244}Cm .

3. Radiation

The radiation is given under Section I.A.3.

4. Critical Mass

See Section I.A.4.

5. Compatibility with Materials of Containment

See ORNL-4359, Compatibility Data Sheets for Cerium-144, Cesium-137, Curium, and Strontium-90 (classified).

6. Thermophysical Properties

- a. Density (calculated from lattice parameters)

C-type (bcc)	10.70 g/cm ³ at room temperature	29
B-type (monoclinic)	11.67 g/cm ³ at room temperature	29
A-type (hex)	11.4 g/cm ³ at 1640°C	29
	11.1 g/cm ³ at 1750°C	
	11.0 g/cm ³ at 1910°C	

CURIUM SESQUOXIDE (Cm_2O_3)

- b. Coefficient of thermal expansion
 B-type (monoclinic) Cm_2O_3 29, 30, 31
 0.60% over 25-680°C
 0.85% over 25-940°C
 1.4% over 25-1300°C
- c. Specific heat and enthalpy
 (1) Specific heat in cal $\text{g}^{-1} \text{ °C}^{-1}$
 $3.7 \times 10^{-2} + 2.91 \times 10^{-5}\text{T } (\text{°K})$ 33 (Am_2O_3)
 (2) Enthalpy
- d. Temperatures of phase transformations
 (1) Phase transitions
 C-type Cm_2O_3 (bcc) transforms irreversibly to
 B-type Cm_2O_3 (monoclinic) between 800-1300°C 29, 31, 32,
 depending on degree of crystallinity of cubic
 form. 34, 35
 B-type Cm_2O_3 (monoclinic) transforms
 reversibly to A-type Cm_2O_3 (hex) at 1600°C. 29, 31, 32,
 34, 35
 (2) Melting point
 $2265 \pm 20^\circ\text{C}$ 16, 36
 $2180 \pm 20^\circ\text{C}$ (production oxide) 16, 36
 $2230 \pm 25^\circ\text{C}$ (SRL production, Batch 1) 3
 (3) Boiling point
 $3700 \pm 200^\circ\text{C}$ 32, 37
 (4) Thermogravimetric analytical work on the
 decomposition of CmO_2 to Cm_2O_3 at SRL con-
 firmed previous work at ORNL and indicated
 some variations from earlier SRL results.
 Stable oxide phases and temperature ranges
 of stability in various atmospheres are:

Phase	Composition	Range of stability in indicated atmosphere, °C		
		Oxygen	Helium	$4\% \text{ H}_2 -$ $96\% \text{ He}$
CmO_2	$\text{CmO}_{2.0}$	<400	<400	<250
$\text{Cm}_{0.11}$	$\text{CmO}_{1.83}$	440-450	Unstable	Unstable
$\text{Cm}_{0.12}$	$\text{CmO}_{1.71}$	460-720	450-520	Unstable
Cm_2O_3				
C-type	$\text{CmO}_{1.62-1.52}$	750-870 ^a	600-900	530-750
B-type	$\text{CmO}_{1.50}$	>850	>950	>850

^aHeating terminated at 870°C.

CURIUM SESQUIOXIDE (Cm_2O_3)

e. Latent heats of phase transformations

ΔH fusion	29 kcal/mole (Richard's rule)	37
ΔH° vaporization	133.0 \pm 2.7 kcal/mole of gas between 1600 and 2250°C	32, 37

f. Vapor pressure

$$\log_{10} P_E \text{ (atm)} = 7.32 (\pm 0.26) - \frac{29,050 (\pm 590)}{T \text{ (°K)}} \quad 32, 35, 37$$

at 1800-2600°K, where P_E is an artificial pressure of $\text{Cm}_2\text{O}_3(g)$. The vapor species in the vaporization reaction are unknown. P_E is related to the actual vapor pressure by a proportionality constant, k , given as follows for vaporization to other gaseous species:

$$\begin{aligned} k &= 1.415 \text{ for } \text{CmO}(g) + \text{CmO}_2(g) \\ k &= 1.567 \text{ for } \text{CmO}(g) + \text{O}(g) \\ k &= 1.869 \text{ for } \text{Cm}(g) + \text{O}(g) \\ k &= 1.414 \text{ for } \text{Cm}(g) + \text{CmO}_2(g) \end{aligned}$$

g. Thermal conductivity

38

Thermal conductivity, cal cm^{-1} sec $^{-1}$ °C $^{-1}$	Molar heat capacity, cal °C $^{-1}$ mole $^{-1}$	Temp, °C
0.00347	31.50	27
0.00378	35.00	127
0.00389	38.32	327
0.00392	40.26	527
0.00399	41.80	727
0.00393	43.06	927
0.00393	44.02	1127
0.00388	44.80	1327
0.00384	45.14	1427

These values were calculated from the thermal diffusivities of Chikalla.³⁸ The density of the hot-pressed pellet was assumed to remain constant at 11.24 g/cm³. Values of the molar heat capacities assumed for ²⁴⁴Cm are those of Pu_2O_3 .

h. Thermal diffusivity

39

Thermal diffusivity, cm 2 /sec	Temperature, °C
0.0053	27
0.0052	127
0.0049	327
0.0047	527
0.0046	727
0.0044	927
0.0043	1127
0.0042	1327
0.0041	1427

CURIUM SESQUIOXIDE (Cm_2O_3)

h. Thermal diffusivity (continued)

Preliminary data on a hot-pressed (30 min at 1380°C) Cm_2O_3 pellet having a density of 11.24 g/cm^3 .

i. Viscosity

j. Surface tension

k. Total hemispherical emittance

l. Spectral emissivity

0.20 to 0.57 22 (ThO_2)

The precise value is a function of the roughness of the material surface, the presence of impurities, and the effects of radiation.

m. Crystallography

Body-centered cubic, C-type rare-earth oxide structure with variable lattice parameter, produced by reducing CmO_2 . 40, 41

$a = 10.922 \pm 0.001$ ($\text{CmO}_{1.62}$) to 10.997 ± 0.003 ($\text{CmO}_{1.52}$) at room temperature

Hexagonal, H-type rare-earth oxide structure, obtained by irreversible transformation or reaction of B-type Cm_2O_3 on noble metals on heating at 1500°C . 30

$a = 3.863 \pm 0.01 \text{ \AA}$ at 1750°C on Ir
 $c = 6.174 \pm 0.001 \text{ \AA}$ at 1750°C on Ir

Cubic with superstructure, related to bcc structure of C-type Cm_2O_3 , possibly $\text{CmO}_{1.72}$, obtained by heating CmO_2 between 470 and 700°C in oxygen (1 atm). 29

$a = 5.450 \pm 0.001 \text{ \AA}$ at room temperature after 470°C in O_2
 $a = 5.461 \pm 0.001 \text{ \AA}$ at room temperature after 700°C in O_2

Cubic, related to fcc structure of CmO_2 , possibly $\text{CmO}_{1.83}$, obtained on heating at 450°C in oxygen. 29

$a = 5.435 \pm 0.001 \text{ \AA}$ at room temperature

Supporting data on crystallography of Cm_2O_3 . 30, 32

n. Solubilities

Insoluble in water

23

o. Diffusion rates

Helium release from $^{244}\text{Cm}_2\text{O}_3$ microspheres and pellets has been correlated to obtain the "apparent diffusion coefficients" for microspheres in the range of 390 to 1200°C and for pellets in the range of 1100 to 1400°C . 42

CURIUM SESQUIOXIDE (Cm_2O_3)

o. Diffusion rates (continued)

Helium diffusion coefficient, cm^2/sec	Temperature, $^{\circ}\text{C}$
---	------------------------------------

Microspheres

4.1×10^{-10}	390
1.74×10^{-9}	600
2.7×10^{-9}	800
4.1×10^{-9}	1000
2.58×10^{-9}	1100
2.77×10^{-9}	1200
3.45×10^{-9}	1200

Cylindrical Pellets

5×10^{-8}	1100
5.8×10^{-8}	1200
7×10^{-8}	1300
5×10^{-8}	1400

7. Mechanical Properties

a. Hardness

b. Crush strength

1900 kg/cm²43 (CeO_2)8. Chemical Properties

a. Heat and free energy of formation, entropy

(1) Heat of formation

Phase $-\Delta H_f^\circ$ (298°K), kcal/mole

$\text{CmO}_{1.50}$	215 ± 7	37
$\text{CmO}_{1.71}$	218 ± 10 (est)	44
$\text{CmO}_{1.83}$	225 ± 10 (est)	
$\text{CmO}_{2.00}$	236 ± 8 (est)	44

(2) Free energy of formation of $\text{CmO}_{1.50}$ ΔG_f° (298°K) = -205 ± 7 kcal/mole (calc)

32, 37

(3) Entropy of $\text{CmO}_{1.50}$ $S_{298}^\circ = 19.4$ eu

45

CURIUM SESQUIOXIDE (Cm_2O_3)

b. Chemical reactions and reaction rates

- (1) Oxidation of B-type Cm_2O_3 in oxygen (1 atm) 16, 30, 31,
 Heating m: 32, 35

<u>Heating temp, °C</u>	<u>Time</u>	<u>Result</u>
75	4 hr	No oxidation
175	4 hr	No oxidation
220	8 hr	80-90% oxidation to CmO_2
290	2 hr	Complete oxidation to CmO_2
360	10 min	Oxidation to 40% Cm_7O_{12} -60% CmO_2
360	30 min	Complete oxidation to CmO_2
450	15 min	Complete oxidation to $\text{CmO}_{1.71}$ ^a
480	15 min	Complete oxidation to $\text{CmO}_{1.71}$
505	4 hr	70-80% oxidation to $\text{CmO}_{1.71}$
530	2 hr	<10% oxidation to $\text{CmO}_{1.71}$
570	2 hr	No oxidation
750	4 hr	No oxidation

^aThe intermediate phase $\text{CmO}_{1.71}$ is produced at somewhat lower temperatures (450°C) by reoxidation of Cm_2O_3 rather than by thermal decomposition of CmO_2 (470°C).

- | | |
|---|----|
| (2) Nitrogen - no reaction | 23 |
| (3) Water - possible hydration reaction | 23 |
| (4) Inorganic acids - soluble in acids | 23 |

9. Biological Tolerances

Maximum permissible body burdens and maximum permissible concentrations of ^{244}Cm in air and in water are given under Section I.A.9. 26

10. Shielding Data

The dose rates are given under Section I.A.10.

C. CURIUM OXYSULFIDE ($\text{Cm}_2\text{O}_2\text{S}$)

1. Composition

- a. Radionuclidic abundance

See Section I.A.1.a.

- b. Radiochemical purity

See Section I.A.1.b.

- c. Chemical purity

See Section I.A.1.c.

2. Specific Power

- a. 2.39 w/g of 100% pure $\text{Cm}_2\text{O}_2\text{S}$ (84.1% ^{244}Cm isotope) 4, 5, 6
 2.32 w/g of 97% pure $\text{Cm}_2\text{O}_2\text{S}$ (81.6% ^{244}Cm isotope)

- b. Specific power values are based on 81.3 Ci/g of
 ^{244}Cm and 34.95 w/kCi of ^{244}Cm .

3. Radiation

The radiation is given under Section I.A.3.

4. Critical Mass

See Section I.A.4.

5. Compatibility with Materials of Containment

6. Thermophysical Properties

- a. Density

9.95 g/cm³

46 ($\text{Pu}_2\text{O}_2\text{S}$)

- b. Coefficient of thermal expansion

0.60% over 25-680°C

29, 30, 31
(Cm_2O_3)

0.85% over 25-940°C

1.4% over 25-1300°C

- c. Specific heat and enthalpy

- (1) Specific heat in cal g⁻¹ °C⁻¹

$$3.62 \times 10^{-2} + 2.83 \times 10^{-5}T \text{ (°K)}$$

33 (Am_2O_3)

- (2) Enthalpy

CURIUM OXYSULFIDE ($\text{Cm}_2\text{O}_2\text{S}$)REFERENCE
COLUMN

d.	Temperatures of phase transformations	
(1)	Melting point	
	2000°C	47 ($\text{Ce}_2\text{O}_2\text{S}$)
(2)	Boiling point	
	3130°C	33 (Am_2O_3)
e.	Latent heats of phase transformations	
	ΔH fusion	17 kcal/mole
	ΔH vaporization	85 kcal/mole
f.	Vapor pressure	
g.	Thermal conductivity	
	Values for the thermal conductivity of curium oxysulfide have not been derived but probably are in the same order of magnitude as the values for Cm_2O_3 which are given in Section I.B.6.g.	38 (Cm_2O_3)
h.	Thermal diffusivity	
	Values for the thermal diffusivity of curium oxysulfide have not been derived but probably are in the same order of magnitude as the values for Cm_2O_3 which are given in Section I.B.6.h.	39 (Cm_2O_3)
i.	Viscosity	
j.	Surface tension	
k.	Total hemispherical emittance	
l.	Spectral emissivity	
	0.20 to 0.57	22 (ThO_2)
	The emissivity value depends on the roughness of the material surface, radiation effects, and the presence of impurities.	
m.	Crystallography	
	Hexagonal	46 ($\text{Pu}_2\text{O}_2\text{S}$)
	$a = 4.008 \text{ \AA}$	$c = 6.769 \text{ \AA}$
	The cell constants should be slightly smaller for $\text{Cm}_2\text{O}_2\text{S}$ due to the actinide contraction.	
n.	Solubilities	
(1)	Soluble in strong acids ($\text{Ce}_2\text{O}_2\text{S}$)	48
(2)	Insoluble in acetic acid ($\text{Ce}_2\text{O}_2\text{S}$)	48

CURIUM OXYSULFIDE ($\text{Cm}_2\text{O}_2\text{S}$)

o. Diffusion rates	
7. <u>Mechanical Properties</u>	
a. Hardness	
b. Crush strength	
8. <u>Chemical Properties</u>	
a. Heat and free energy of formation, entropy	
(1) Heat of formation	
$\Delta H_f^\circ = -430 \text{ kcal/mole}$ (approximated by $\text{Ce}_2\text{O}_2\text{S}$)	47
(2) Free energy of formation	
$\Delta F_f^\circ = -414 \text{ kcal/mole}$ (calculated by)	
$\Delta F_f^\circ = \Delta H_f^\circ - T\Delta S_f^\circ$)	
(3) Entropy	
$S_{298}^\circ = 34.3 \text{ eu}$ (calculated by Latimer's method)	49
b. Chemical reactions and reaction rates	46
(1) Air at room temperature — slow	
(2) Air at elevated temperature — fast	
(3) Water at room temperature — no reaction	
(4) Inorganic acids at room temperature — reacts	
9. <u>Biological Tolerances</u>	
Maximum permissible body burdens and maximum permissible concentrations of ^{244}Cm in air and in water are given in the ^{244}Cm Metal Source Form, Section I.A.9.	26
10. <u>Shielding Data</u>	
The radiation dose rates are given under the ^{244}Cm Metal Source Form, Section I.A.10.	

D. CURIUM FLUORIDE (CmF_3)

1. Composition

- a. Radionuclidic abundance

See Section I.A.1.a.

- b. Radiochemical purity

See Section I.A.1.b.

- c. Chemical purity

See Section I.A.1.c.

2. Specific Power

- a. 2.19 w/g of 100% pure CmF_3 (77.1% ^{244}Cm isotope)

4, 5, 6

- 2.12 w/g of 97% pure CmF_3 (74.8% ^{244}Cm isotope)

b. Specific power values are based on 81.3 Ci/g of

^{244}Cm (100%) and 34.95 w/kCi of ^{244}Cm .

3. Radiation

The radiation is given under Section I.A.3.

4. Critical Mass

See Section I.A.4.

5. Compatibility with Materials of Containment

6. Thermophysical Properties

- a. Density

9.80 g/cm³

50

- b. Coefficient of thermal expansion

$$a = 19.74 \times 10^{-6} + 2.62 \times 10^{-8} t + 0.15 \times 10^{-10} t^2$$

51 (BaF_2)

(t is in °C with a temperature range of 26-296°C)

- c. Specific heat and enthalpy

- (1) Specific heat in cal g⁻¹ °C⁻¹

$$7.21 \times 10^{-2} + 2.06 \times 10^{-5} T (\text{°K})$$

33 (AmF_3)

- (2) Enthalpy in cal

$$H_f - H_{298} = 21 T + 3.5 \times 10^{-3} T^2$$

33

(estimated from other actinide data)

CURIUM FLUORIDE (CmF_3)

d. Temperatures of phase transformations

(1) Melting point

 1406°C

52

(2) Boiling point

 2330°C

14

This value is the average of the boiling points of LaF_3 , CeF_3 , and PrF_3 .

e. Latent heats of phase transformations

 ΔH fusion 9 kcal/mole33 (CeF_3) ΔH vaporization 62 kcal/mole33 (CeF_3)

f. Vapor pressure

<u>Vapor pressure, torr</u>	<u>Temperature, $^\circ\text{C}$</u>	53
5.07×10^{-7}	853	
7.18×10^{-6}	943	
1.486×10^{-4}	1033	
1.725×10^{-3}	1141	
5.820×10^{-3}	1196	

Vapor pressures are for AmF_3 .

g. Thermal conductivity

<u>Thermal conductivity,</u> <u>cal $\text{cm}^{-1} \text{ sec}^{-1} \text{ }^\circ\text{C}^{-1}$</u>	<u>Temperature,</u> <u>$^\circ\text{C}$</u>	54 (BaF_2)
0.0296	0	
0.0251	100	

h. Thermal diffusivity

<u>Thermal diffusivity,</u> <u>cm^2/sec</u>	<u>Temperature,</u> <u>$^\circ\text{C}$</u>
0.0399	0
0.0321	100

Calculated by dividing the product of the specific heat and the room temperature density into the thermal conductivity.

i. Viscosity

j. Surface tension

k. Total hemispherical emittance

A value of 0.9 can be assumed.

l. Spectral emissivity

CURIUM FLUORIDE (CmF_3)

m.	Crystalllography	
	Hexagonal, LaF_3 -type, space group $P\bar{6}_3/mmc$	55
	$a = 4.041 \pm 0.001 \text{ \AA}$ $c = 7.179 \pm 0.002 \text{ \AA}$	
n.	Solubilities	
o.	Diffusion rates	
7.	<u>Mechanical Properties</u>	
a.	Hardness	
b.	Crush strength	
8.	<u>Chemical Properties</u>	
a.	Heat and free energy of formation, entropy	
(1)	Heat of formation	
	$\Delta H_f^\circ = -357 \text{ kcal/mole}$	15
	(estimated from other actinide data)	
(2)	Free energy of formation	
	$\Delta F_f^\circ = -339 \text{ kcal/mole}$	15
(3)	Entropy	
	$S_{298}^\circ = 29 \text{ eu}$ (estimated from other actinide data)	33
b.	Chemical reactions and reaction rates	23
(1)	Air at room temperature — no reaction	
(2)	Air at elevated temperature — forms oxyfluoride	
(3)	Nitrogen — no reaction	
(4)	Water — insoluble	
(5)	Inorganic acids — insoluble in weak inorganic acid	
9.	<u>Biological Tolerances</u>	
	Maximum permissible body burdens and maximum permissible concentrations of ^{244}Cm in air and in water are given under Section I.A.9.	26
10.	<u>Shielding Data</u>	
	The dose rates are given under Section I.A.10.	

II. CURIUM-242

A. CURIUM-242 OXIDE CERMET

Half-life: 163 d

1, 5, 7, 9

1. Composition

a. Radionuclidic abundance

The composition of the product will depend on the irradiation history of the ^{241}Am target as well as the cooling time after removal from the pile. The product is expected to be at least 40% ^{242}Cm . If it is not this high, an Am-Cm separation is used to lower the americium content of the product.

The analysis for a typical batch of ^{241}Am with a total integrated neutron dose of 1.3×10^{21} and with a 90-day cooling and processing period (with all the plutonium removed) is as follows:

<u>Isotope</u>	<u>% Abundance</u>
^{241}Am	47.5
^{242}Am	1.6
^{243}Am	7.7
^{242}Cm	41.9
^{243}Cm	0.5
^{244}Cm	0.8

The decay of ^{242}Cm to ^{238}Pu is illustrated in the following table. A 10-g product with 40% ^{242}Cm and 60% other actinides is assumed. The decay is shown for 163 days.

<u>Time, days</u>	<u>$^{242}\text{Cm}, \text{g}$</u>	<u>$^{238}\text{Pu}, \text{g}$</u>
0	4.00	0
16	3.73	0.27
32	3.48	0.52
65	3.03	0.97
81.5	2.83	1.17
98	2.46	1.54
163	2.00	2.00

The oxide mixture will be AmO_2 and Cm_2O_3 . This will be suspended in a neutral matrix material to give the prescribed power density:

30 vol % oxide mixture
 70 vol % matrix material

CURIUM-242 OXIDE CERMETREFERENCE
COLUMN

b. Radiochemical purity

The only important heat contributor to the ^{242}Cm source material is ^{242}Cm . The following table shows the contribution of each isotope to 10 g of the oxide mixture.

Heat Contribution of Each Isotope of the ^{242}Cm Product

Nuclide	Half-life	Specific activity, w/g	% of nuclide	Heat contribution watts	%
^{241}Am	458 y	0.106	47.5	0.50	0.1
^{242}Am	152 y	0.034	1.6	0.05	0.01
^{243}Am	7650 y	0.006	7.7	0.005	0.001
^{242}Cm	163 d	120.0	41.9	502.8	99.8
^{243}Cm	32 y	1.44	0.5	0.07	0.015
^{244}Cm	18.1 y	2.78	0.8	0.22	0.05
^{238}Pu	89 y	0.55			

The ^{242}Cm product has contained up to 30 Ci of ^{144}Ce per 8.35 g of ^{242}Cm . Trace amounts of ^{103}Ru - ^{106}Ru and ^{95}Zr - ^{95}Nb have been found in the feed but do not contribute materially to the power or radiation of the source.

The above contaminations can be reduced to much lower levels by additional processing.

2. Specific Power

a. 42.8 w/g of $\text{AmO}_2\text{-Cm}_2\text{O}_3$ (35.7% ^{242}Cm metal)

It is assumed that there are 3320 Ci/g of ^{242}Cm and 36.1 w/kCi of ^{242}Cm .

b. 1.186 Ci of ^{242}Cm per gram of $\text{AmO}_2\text{-Cm}_2\text{O}_3$ (35.7% ^{242}Cm metal)

3. Radiation

a. Alpha particles

Nuclide	Max E, Mev	Av E, Mev	Abundance, %	w/kCi	Particles $\text{w}^{-1} \text{sec}^{-1}$
^{242}Cm	6.11	6.11	73.7	{ 36.10	0.755×10^{12}
	6.066	6.066	26.3		0.270×10^{12}

CURIUM-242 OXIDE CERMET

REFERENCE
COLUMN

a. Alpha particles (continued)

The amount of helium produced by alpha decay of ^{242}Cm as a function of time is given in the following table.

Volume of helium, ^a cm ³ of He per g of ^{242}Cm	Time Days	Time Half-lives
11.9	32.6	0.2
22.4	65.2	0.4
31.5	97.8	0.6
39.4	130.4	0.8
46.3	163	1.0
59.8	244	1.5
69.4	326	2.0
80.9	489	3.0
86.7	652	4.0
89.6	815	5.0
92.4	1630	10.0

^aStandard conditions.

b. Beta particles

None

c. Gamma

In addition to the gammas from the alpha decay, there are the prompt and fission-product gammas from the spontaneous fission of ^{242}Cm ($T_{1/2} = 7.2 \times 10^6$ y). The gamma-emission rates are given in the following table.

Gamma-emission rate, ^a photons sec ⁻¹ g ⁻¹	Photon energy, Mev
<u>Curium-242 Gammas</u>	
0.27×10^{14}	0.044
1.42×10^{10}	0.10
2.83×10^9	0.158
<u>Prompt Gammas</u>	
2.44×10^7	1.0
6.09×10^6	1.5
6.45×10^6	2.3
1.15×10^6	3.0
1.52×10^6	5.0

CURIUM-242 OXIDE CERMET

Gamma-emission rate, ^a photons sec ⁻¹ g ⁻¹	Photon energy, Mev
<u>Fission-Product Gammas</u>	
2.16 x 10 ⁷	0.63
7.99 x 10 ⁶	1.1
9.36 x 10 ⁶	1.55
1.80 x 10 ⁶	2.38
2.66 x 10 ⁶	2.75

^a120-w source.

d. Bremsstrahlung

None

e. Neutrons

2.30 x 10⁷ neutrons sec⁻¹ g⁻¹ of ²⁴²Cm from spontaneous fission (120-w source)

7

2.0 x 10⁷ neutrons sec⁻¹ g⁻¹ of ²⁴²Cm from (α ,n) reaction of oxygen in Cm_2O_3 (120-w source)

The energy distribution of spontaneous fission neutrons from ²⁴²Cm is given in the following table. 4

Spontaneous Fission Neutrons from Curium-242

Energy, Mev	Abundance, ^a neutrons sec ⁻¹ g ⁻¹ of ²⁴² Cm
0.3-0.4	7.7 x 10 ⁵
0.4-0.6	1.6 x 10 ⁶
0.6-0.8	1.6 x 10 ⁶
0.8-1.0	1.4 x 10 ⁶
1.0-1.2	1.4 x 10 ⁶
1.2-1.4	1.4 x 10 ⁶
1.4-1.6	1.2 x 10 ⁶
1.6-1.8	1.1 x 10 ⁶
1.8-2.0	1.0 x 10 ⁶
2.0-2.2	9.1 x 10 ⁵
2.2-2.4	8.4 x 10 ⁵
2.4-2.6	8.0 x 10 ⁵
2.6-2.8	6.5 x 10 ⁵
2.8-3.0	5.5 x 10 ⁵
3.0-3.2	5.0 x 10 ⁵
3.2-3.4	5.0 x 10 ⁵
3.4-3.6	4.7 x 10 ⁵
3.6-3.8	3.7 x 10 ⁵
3.8-4.0	4.1 x 10 ⁵

CURIUM-242 OXIDE CERMETREFERENCE
COLUMN

Spontaneous Fission Neutrons (continued)

Energy, Mev	Abundance, ^a neutrons sec ⁻¹ g ⁻¹ of ²⁴² Cm
4.0-4.4	5.2 x 10 ⁵
4.4-4.8	4.4 x 10 ⁵
4.8-5.2	3.2 x 10 ⁵
5.2-5.6	2.5 x 10 ⁵
5.6-6.0	1.9 x 10 ⁵
6.0-6.4	1.5 x 10 ⁵
6.4-6.8	1.1 x 10 ⁵
6.8-7.2	7.5 x 10 ⁴
7.2-7.6	5.6 x 10 ⁴
7.6-8.0	4.7 x 10 ⁴
8.0-8.8	5.0 x 10 ⁴
8.8-9.6	1.5 x 10 ⁴
9.6-10.4	1.6 x 10 ⁴
10.4-11.2	1.0 x 10 ⁴
11.2-12.8	7.1 x 10 ³

^a120-w source.

The energy distribution of neutrons occurring as a result of a collision of fast alpha particles from ²⁴²Cm decay with oxygen atoms in Cm_2O_3 is given in the following table.

4

Neutrons from (α ,n) Reactions with Oxygen

Energy, Mev	Abundance, ^a neutrons sec ⁻¹ g ⁻¹ of ²⁴² Cm
0.2	5.0 x 10 ³
0.4	1.0 x 10 ⁴
0.6	2.0 x 10 ⁴
0.8	2.5 x 10 ⁴
1.0	7.6 x 10 ⁴
1.2	1.5 x 10 ⁵
1.4	2.8 x 10 ⁵
1.6	4.5 x 10 ⁵
1.8	7.6 x 10 ⁵
2.0	1.0 x 10 ⁶
2.2	1.3 x 10 ⁶
2.4	1.7 x 10 ⁶
2.6	2.0 x 10 ⁶
2.8	2.1 x 10 ⁶
3.0	2.1 x 10 ⁶
3.2	2.1 x 10 ⁶
3.4	1.8 x 10 ⁶

CURIUM-242 OXIDE CERMETNeutrons from (α, n) Reactions (continued)

Energy, Mev	Abundance, ^a neutrons sec ⁻¹ g ⁻¹ of ^{242}Cm
3.6	1.5×10^6
3.8	1.0×10^6
4.0	6.5×10^5
4.2	4.0×10^5
4.4	2.8×10^5
4.6	1.8×10^5
4.8	5.0×10^4

^a120-w source.4. Critical Mass5. Compatibility with Materials of Containment

See ORNL-4359, Compatibility Data Sheets for Cerium-144, Cesium-137, Curium, and Strontium-90 (classified).

6. Thermophysical Properties

The thermophysical properties of a ^{242}Cm oxide cermet will depend on the individual properties of the oxide and matrix material. With the selection of the proper refractory metal as the matrix material, it should be possible to reach a $0.06 \text{ cal cm}^{-1} \text{ sec}^{-1} {^\circ}\text{C}^{-1}$ thermal conductivity for the cermet.

7. Mechanical Properties

The mechanical properties of the metal-oxide mixture will depend strongly on the properties of the matrix material.

8. Chemical Properties

The chemical properties will depend on the individual properties of the oxide and matrix material.

9. Biological Tolerances

The ^{242}Cm tolerances taken from reference 26 are given in the table on the following page.

CURTUM-242 OXIDE CERMETREFERENCE
COLUMN

Maximum Permissible Body Burdens and Maximum Permissible Concentrations
for Radionuclides in Air and in Water for Occupational Exposure²⁶

Radionuclide and type of decay	Organ of reference (critical organ underscored)	Max permissible burden in total body, g(μ Ci)	Max permissible concentrations, μ Ci/cm ³			
			For 40-hr week		For 168-hr week	
			Water	Air	Water	Air
^{242}Cm ^{96}Cm (α , γ)	GI (LLI) ^a (Sol)	7 $\times 10^{-4}$	2 $\times 10^{-7}$	2 $\times 10^{-4}$	5 $\times 10^{-8}$	
		Liver	0.05	3 $\times 10^{-3}$	10 ⁻¹⁰	9 $\times 10^{-4}$
		Bone	0.09	5 $\times 10^{-3}$	2 $\times 10^{-10}$	2 $\times 10^{-3}$
		Kidney	0.2	9 $\times 10^{-3}$	4 $\times 10^{-10}$	3 $\times 10^{-3}$
	Total Body	0.2	0.01	6 $\times 10^{-10}$	5 $\times 10^{-3}$	2 $\times 10^{-10}$
	(Insol)	Lung		2 $\times 10^{-10}$		6 $\times 10^{-11}$
		GI (LLI) ^a	7 $\times 10^{-4}$	10 ⁻⁷	3 $\times 10^{-4}$	4 $\times 10^{-8}$

^aThe abbreviations GI and LLI refer to the gastrointestinal tract and lower large intestine, respectively.

10. Shielding Data

Gamma dose rates with water, iron, lead, and uranium shielding are given in Figs. 8-12 for ^{242}Cm power sources of 100, 200, 500, 1000, 2000, 5000, 10,000, and 20,000 w. Neutron dose rates with water shielding are given in Fig. 13. Neutron dose rates on shielding with Be, Ch, CH₂, or LiH can be estimated by using Fig. 13 in conjunction with Fig. 7.

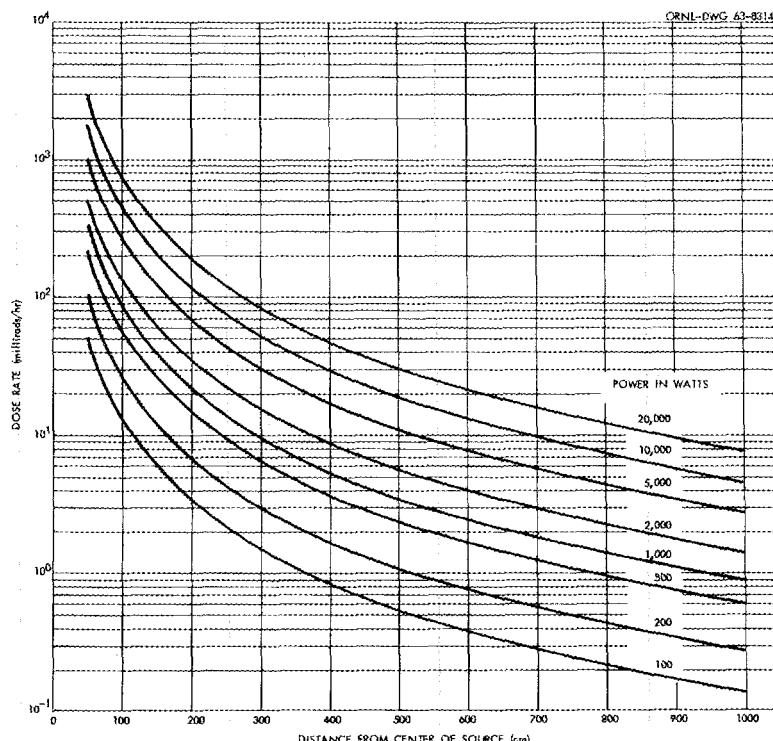


Fig. 8. Gamma Dose Rates from Unshielded Isotopic Power Sources of Curium-242 as a Function of Distance from Center of Source.

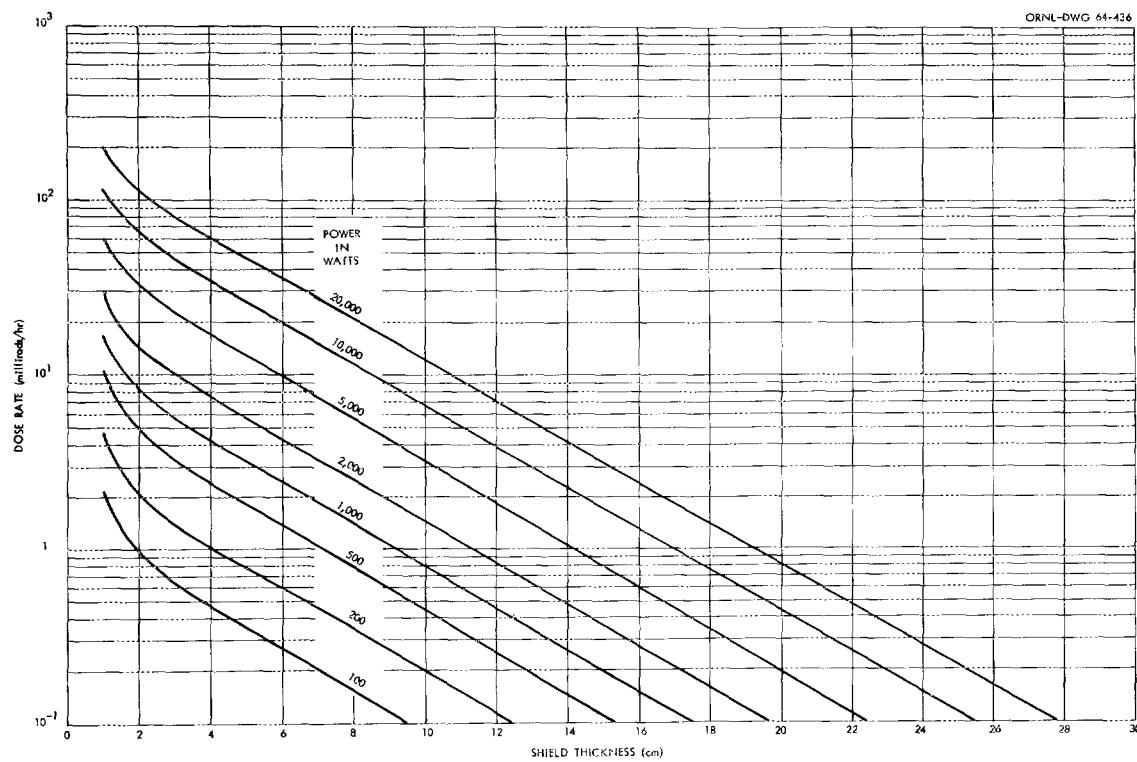


Fig. 9. Gamma Dose Rates from Iron-Shielded Isotopic Power Sources of Curium-242. Center of source to dose point separation distance is 100 cm.

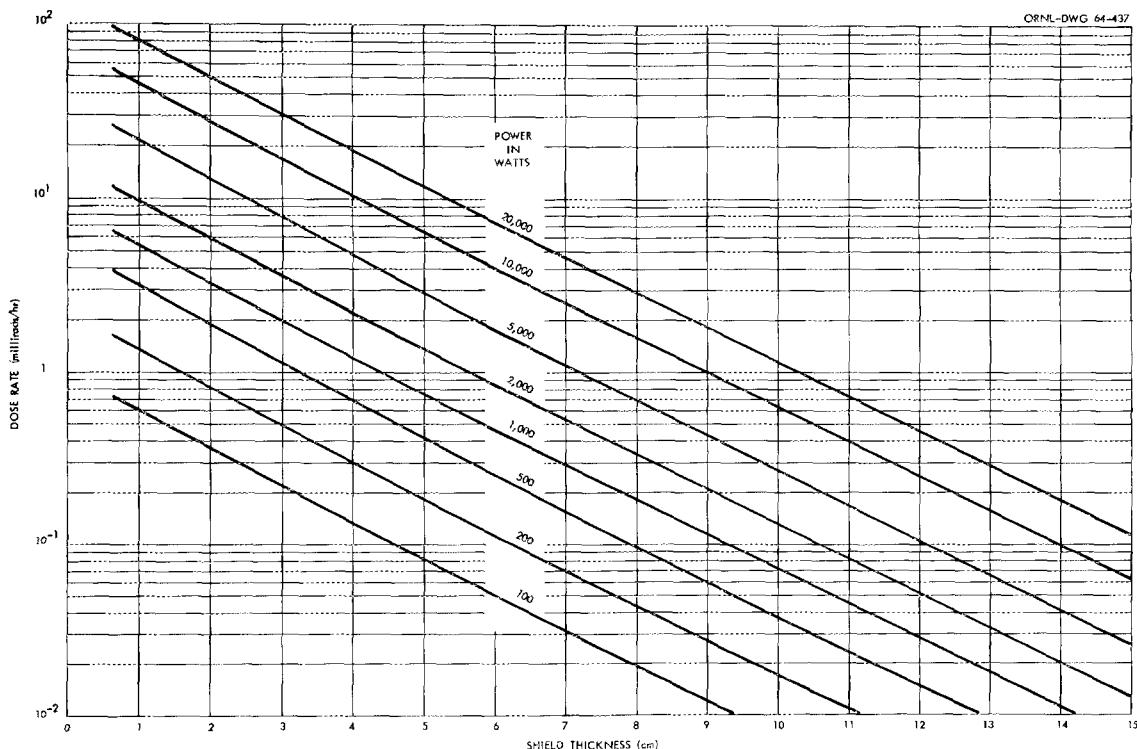


Fig. 10. Gamma Dose Rate from Lead-Shielded Isotopic Power Sources of Curium-242. Center of source to dose point separation distance is 100 cm.

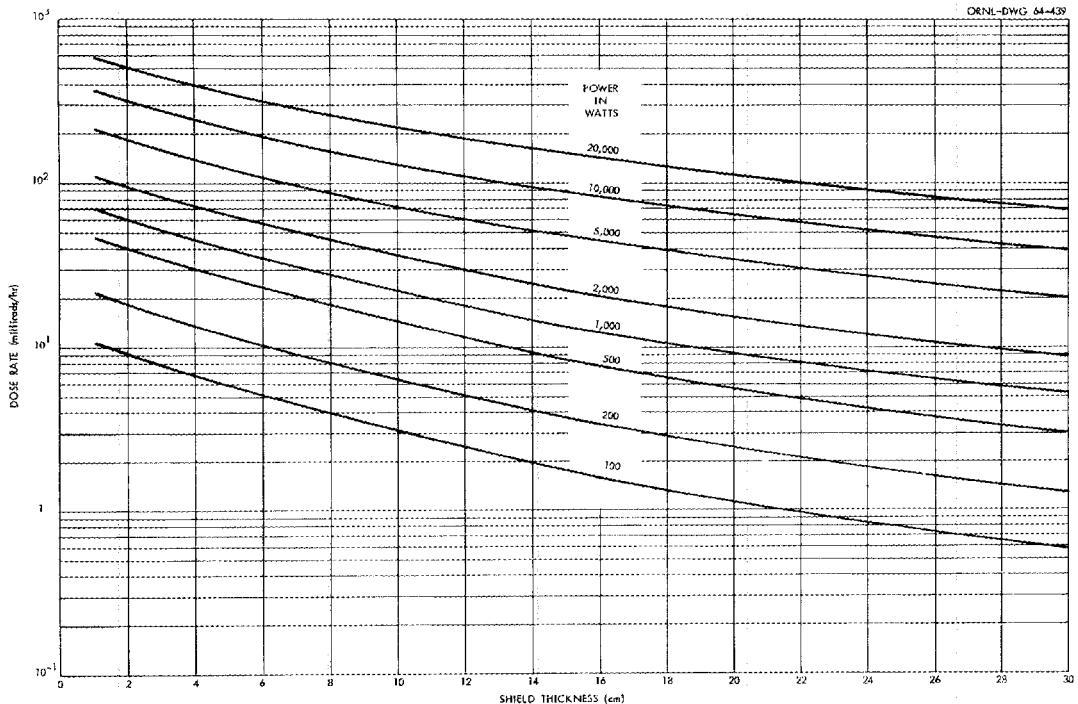


Fig. 11. Gamma Dose Rates from Water-Shielded Isotopic Power Sources of Curium-242. Center of source to dose point separation distance is 100 cm.

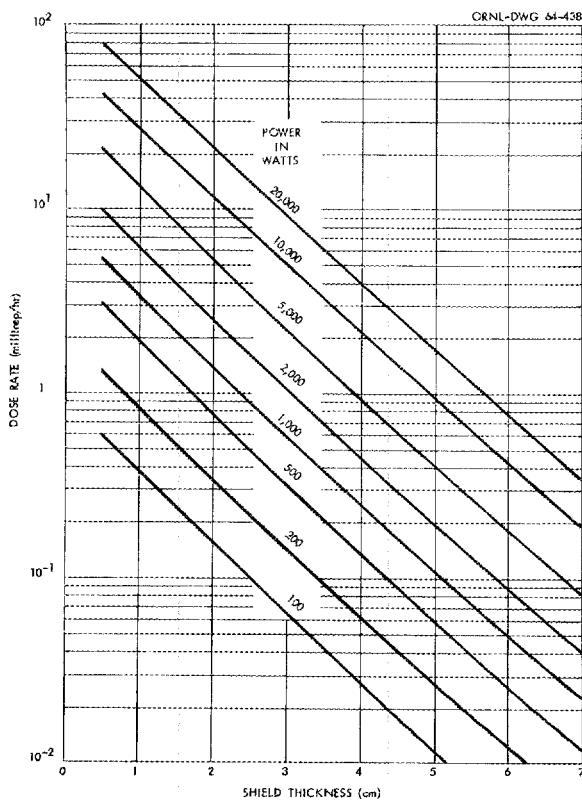


Fig. 12. Gamma Dose Rates from Uranium-Shielded Isotopic Power Sources of Curium-242. Center of source to dose point separation distance is 100 cm.

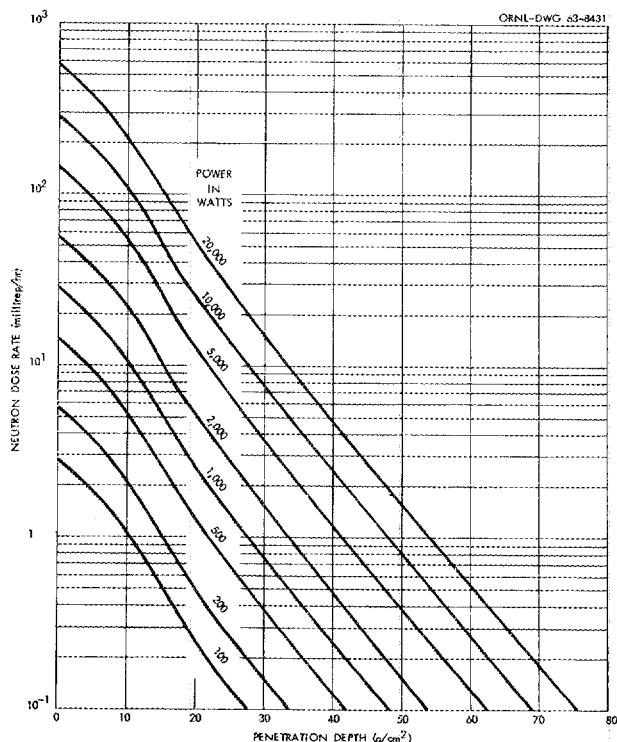


Fig. 13. Neutron Dose Rates from Water-Shielded Isotopic Power Sources of Curium-242 as a Function of Penetration Depth of Shielding Material. Center of source to dose point separation distance is 100 cm. Refer to Fig. 7 for dose rates through other materials.

III. REFERENCES FOR CURIUM

1. Knolls Atomic Power Laboratory, General Electric Company, Chart of the Nuclides, 7th ed., revised to June 1964.
2. W. C. Bentley, Alpha Half-life of ^{244}Cm , J. Inorg. Nucl. Chem. 30: 2007-9 (1968).
3. D. G. Ebenhack, Savannah River Laboratory, private communication, November 1968.
4. D. H. Stoddard, Radiation Properties of ^{244}Cm Produced for Isotopic Power Generators, USAEC Rpt. DP-939, Savannah River Laboratory (November 1964).
5. B. S. Dzhelepov and L. K. Peker, Decay Schemes of Radioactive Nuclei, Pergamon Press, London and New York, 1961.
6. D. Metta et al., Nuclear Constants of Nine Transplutonium Nuclides, Abstracts of Papers, 148th Am. Chem. Soc. Meeting, Chicago, 1964, p. 8R.
7. E. D. Arnold, Handbook of Shielding Requirements and Radiation Characteristics of Isotopic Power Sources for Terrestrial, Marine, and Space Applications, USAEC Rpt. ORNL-3576, Oak Ridge National Laboratory (April 1964).
8. S. J. Rimshaw, Isotopic Power Data Sheets, USAEC Rpt. TID-7698 (Pt. 1), Conf. on Isotopic Power Development, Germantown, Maryland, May 18-19, 1964.
9. C. A. Rohrman, Radioisotopic Heat Sources, USAEC Rpt. HW-76323, Hanford Laboratories (February 1963).
10. R. A. Robinson et al., Brayton-Cycle Radioisotope Heat Source Design Study, Phase I (Conceptual Design) Report, USAEC Rpt. ORNL-TM-1691-del., Oak Ridge National Laboratory (August 1967), pp. 107-13.
11. B. B. Cunningham and J. C. Wallmann, Crystal Structure and Melting Point of Curium Metal, J. Inorg. Nucl. Chem. 26: 271-5 (1964).
12. D. B. McWhan, B. B. Cunningham, and J. C. Wallmann, Crystal Structure, Thermal Expansion, and Melting Point of Americium Metal, J. Inorg. Nucl. Chem. 24: 1025-38 (1962).
13. K. A. Gschneidner, Solid State Physics, F. Seitz and D. Turnbull, eds., Vol. 16, Academic Press, 1964.
14. J. A. Gibson et al., The Properties of the Rare Earth Metals and Compounds, Battelle Memorial Institute, Columbus, Ohio, May 1959.
15. B. B. Cunningham, Compounds of the Actinides, in Preparative Inorganic Reactions, Vol. 3, William L. Jolly (ed.), Interscience Publishers, New York, 1966, pp. 79-121.

16. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, January-March 1968, USAEC Rpt. DP-1155-II, Savannah River Laboratory (classified).
17. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, April-June 1968, USAEC Rpt. DP-1169-II, Savannah River Laboratory (classified).
18. S. N. Lvov, V. F. Nemchenko, and Yu. B. Paderno, Thermal Conductivities of the Alkaline Earth and Rare Earth Hexaborides, Dokl. Akad. Nauk SSSR 146(6): 1371-2 (1963).
19. E. N. da C. Andrade, A Theory of the Viscosity of Liquids — Part II, Phil. Mag. 17: 698-732 (1934).
20. O. Flint, Surface Tension of Liquid Metals (a review), British Rpt. AERE-M-1479 (November 1964).
21. R. N. Karlsson, The Total Hemispherical Emittance of Polished and Oxidized Alpha Plutonium, USAEC Rpt. RFP-616, The Dow Chemical Company (July 1965).
22. R. C. Weast, S. M. Selby, and C. D. Hodgman, Handbook of Chemistry and Physics, 45th ed., The Chemistry Rubber Company, Cleveland, Ohio, 1964.
23. J. L. Burnett, Oak Ridge National Laboratory, private communication, November 1965.
24. T. Lyman, ed., Metals Handbook, 8th Ed., Vol. 1, Properties and Selection of Metals, American Society for Metals, Metals Park, Novelty, Ohio, 1961.
25. R. M. Dell and M. Allbutt, The Nitrides of Uranium, Thorium, and Plutonium. A Review of Present Knowledge, British Rpt. AERE-R-4253 (March 1963).
26. U. S. Department of Commerce, National Bureau of Standards, Maximum Permissible Body Burdens and Maximum Permissible Concentrations of Radionuclides in Air and in Water for Occupational Exposure, Handbook 69 (June 5, 1959).
27. J. C. Posey, P. R. Kuehn, and R. E. McHenry, The Influence of Temperature and Oxygen Pressure on the Composition of Curium Oxide, presented at the Am. Chem. Soc. Meeting, Atlantic City, New Jersey, September 1965.
28. T. D. Chikalla and L. Eyring, The Curium-Oxygen System, USAEC Rpt. BNWL-CC-1569, Pacific Northwest Laboratory (February 13, 1968).
29. W. C. Mosley, B-Type $^{244}\text{Cm}_2\text{O}_3$ — A Candidate Isotopic Power Fuel, Abstracts of Papers, 155th National Meeting, Am. Chem. Soc., San Francisco, California, April 1968, No. O-173.

30. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, January-March 1967, USAEC Rpt. DP-1105-II, Savannah River Laboratory (classified).
31. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, April-June 1967, USAEC Rpt. DP-1120-II, Savannah River Laboratory (classified).
32. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, October-December 1967, USAEC Rpt. DP-1143-II, Savannah River Laboratory (classified).
33. A. Glassner, The Thermochemical Properties of the Oxides, Fluorides, and Chlorides to 2500°K, USAEC Rpt. ANL-5750, Argonne National Laboratory (1957).
34. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, July-September 1966, USAEC Rpt. DP-1088, Savannah River Laboratory (classified).
35. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, July-September 1967, USAEC Rpt. DP-1129-II, Savannah River Laboratory (classified).
36. P. K. Smith, Melting Point of Cm_2O_3 , J. Inorg. Nucl. Chem. (to be published).
37. P. K. Smith and D. E. Peterson, High Temperature Evaporation and Thermodynamic Properties of Cm_2O_3 , Abstracts of Papers, 155th National Meeting, Am. Chem. Soc., San Francisco, California, April 1968, No. O-031.
38. J. C. Posey, Oak Ridge National Laboratory, unpublished data, April 1968.
39. T. D. Chikalla, Pacific Northwest Laboratory, unpublished data, April 1968.
40. H. S. Hilborn (comp.), Savannah River Laboratory Isotopic Power and Heat Sources, Quarterly Progress Report, October-December 1966, USAEC Rpt. DP-1094, Savannah River Laboratory.
41. Herman O. Haug, Curium Sesquioxide Cm_2O_3 , J. Inorg. Nucl. Chem. 29: 2753-58 (1967).
42. P. Angelini and R. E. McHenry, Diffusion of Helium in Curium- 244 Fuel Forms, Abstracts of Papers, 155th National Meeting, Am. Chem. Soc., San Francisco, California, April 1968, No. O-174.
43. J. R. Hague et al. (eds.), Refractory Ceramics for Aerospace, American Ceramics Society, Columbus, Ohio, 1964.

44. W. R. McDonell et al., Curium-244 for Radioisotopic Heat Sources - Work at Savannah River Laboratory, USAEC Rpt. DP-1066, Vol. II, Savannah River Laboratory (May 1966).
45. E. F. Westrum, Jr., and Fredrik Gronvold, Thermochemical Data of Actinide Element Chalcogenides, in Thermodynamics of Nuclear Materials, International Atomic Energy Agency, Vienna, 1962, p. 28.
46. W. H. Zachariasen, Crystal Chemical Studies of the 5-f Series of Elements, VII. The Crystal Structure of $\text{Ce}_2\text{O}_2\text{S}$, $\text{La}_2\text{O}_2\text{S}$, and $\text{Pu}_2\text{O}_2\text{S}$, Acta Cryst. 2: 60-2 (1949).
47. E. D. Eastman et al., Preparation and Properties of the Oxide-Sulfides of Cerium, Zirconium, Thorium, and Uranium, J. Am. Chem. Soc. 73: 3896-9 (1951).
48. G. V. Samsonov and S. V. Radzikovskaya, The Chemistry of Rare Earth and Actinide Sulfides, Russian Chemical Reviews, 30: 28-41 (1961).
49. W. Latimer, Oxidation Potentials, Prentice-Hall, Inc., Englewood Cliffs, New Jersey, 1952.
50. A. N. Hodge, The Fluorides of the Actinide Elements in Advances in Fluorine Chemistry, Vol. 2, Butterworth, London, 1961, pp. 138-82.
51. D. B. Sirdeshmukh and V. T. Desphande, Temperature Variation of the Lattice Constants and the Coefficients of Thermal Expansion of Some Fluoride Type Crystals, Indian J. Pure Appl. Phys. 2(12): 405-7 (1964).
52. J. Burnett, Melting Points of CmF_3 and AmF_3 , J. Inorg. Nucl. Chem. 28: 2454-6 (1966).
53. S. C. Carnigler and B. B. Cunningham, Vapor Pressures of Americium Trifluoride and Plutonium Trifluoride, Heats and Free Energies of Sublimation, J. Am. Chem. Soc. 77: 1451-3 (1955).
54. K. A. McCarthy and S. S. Ballard, Thermal Conductivity of Eight Halide Crystals in the Temperature Range 220°K to 390°K, J. Appl. Phys. 31(8): 1410-2 (1960).
55. L. B. Asprey, T. K. Keenan, and F. H. Kruse, Crystal Structures of the Trifluorides, Trichlorides, Tribromides, and Triiodides of Americium and Curium, J. Inorg. Chem. 4(7): 985-6 (1965).
56. H. A. O'Brien, Jr., Oak Ridge National Laboratory, private communication, April 24, 1963.

ORNL-4357
UC-4 — Chemistry

INTERNAL DISTRIBUTION

- | | |
|-------------------------------------|-------------------------|
| 1. Biology Library | 72-73. E. E. Ketchen |
| 2-4. Central Research Library | 74. R. H. Lafferty, Jr. |
| 5-6. ORNL — Y-12 Technical Library | 75. E. Lamb |
| Document Reference Section | 76. C. E. Larson |
| 7-41. Laboratory Records Department | 77. R. J. Lauer |
| 42. Laboratory Records, ORNL R.C. | 78. K. H. Lin |
| 43. R. G. Affel | 79. H. G. MacPherson |
| 44. P. S. Baker | 80. H. A. Mahlman |
| 45. Russell Baldock | 81. H. F. McDuffie |
| 46. R. D. Baybarz | 82. R. E. McHenry |
| 47. E. E. Beauchamp | 83. J. R. McWherter |
| 48. C. E. Bemis, Jr. | 84. F. L. Moore |
| 49. J. E. Bigelow | 85. R. G. Niemeyer |
| 50. D. S. Billington | 86. P. B. Orr |
| 51. F. T. Binford | 87. C. L. OttINGER |
| 52. G. E. Boyd | 88. J. J. Pinajian |
| 53. W. D. Burch | 89. J. K. Poggenburg |
| 54. J. Burnett | 90-114. S. J. Rimshaw |
| 55. T. A. Butler | 115. R. A. Robinson |
| 56. F. N. Case | 116. P. S. Rudolph |
| 57. T. G. Clark | 117. A. F. Rupp |
| 58. J. W. Cleland | 118. H. T. Russell |
| 59. T. F. Connolly | 119. R. W. Schaich |
| 60. W. Davis, Jr. | 120. J. R. Sites |
| 61. J. R. DiStefano | 121. M. J. Skinner |
| 62. C. Feldman | 122. H. M. Smith |
| 63. F. J. Furman | 123. D. A. Sundberg |
| 64. J. H. Gillette | 124. J. R. Tarrant |
| 65. K. W. Haff | 125. W. A. Thomas |
| 66. F. E. Harrington | 126. W. E. Unger |
| 67. J. D. Hoeschele | 127. A. M. Weinberg |
| 68. R. W. Horton | 128. R. C. Weir |
| 69. H. B. Hupf | 129. J. C. White |
| 70. H. Inouye | 130. J. P. Witherspoon |
| 71. D. Jacobs | 131. R. G. Wymer |

EXTERNAL DISTRIBUTION

- 132. Director, Division of Space Nuclear Systems, AEC, Washington, D.C.
- 133. Director, Division of Reactor Development and Technology, AEC, Washington, D.C.
- 134-136. Director, Division of Isotopes Development, AEC, Washington, D.C.
- 137. Director, Division of Production, AEC, Washington, D.C.
- 138. Assistant Director, Space Electric Power Office, SNS-AEC, Washington, D.C.
- 139. Chief, Isotope Power Systems Branch, SEPO-SNS-AEC, Washington, D.C.

140. Chief, Advanced Engine Branch, SNS-AEC, Washington, D.C.
141. Chief, Safety Branch, SEPO-SNS-AEC, Washington, D.C.
142. Chief, Space Isotopes and Materials Branch, SEPO-SNS-AEC, Washington, D.C.
143. Chief, Isotopic Auxiliary Power Branch, DRD and T-AEC, Washington, D.C.
144. Assistant Director, Nuclear Safety Office, DRD and T-AEC, Washington, D.C.
145. J. A. Swartout, Union Carbide Corporation, New York
146. Laboratory and University Division, AEC, ORO
- 147-356. Given distribution as shown in TID-4500 under Chemistry category
(25 copies -- CFSTI)