MASTER

ORNL-2548
Chemistry - General
TID-4500 (15th ed.)

PHASE DIAGRAMS OF NUCLEAR REACTOR MATERIALS

R. E. Thoma, Editor

OAK RIDGE NATIONAL LABORATORY

operated by

UNION CARBIDE CORPORATION

for the

U.S. ATOMIC ENERGY COMMISSION

Printed in USA. Price \$3.50. 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.

DISCLAIMER

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

DISCLAIMER

Portions of this document may be illegible in electronic image products. Images are produced from the best available original document.

ORNL-2548
Chemistry - General
TID-4500 (15th ed.)

Contract No. W-7405-eng-26

REACTOR CHEMISTRY DIVISION

PHASE DIAGRAMS OF NUCLEAR REACTOR MATERIALS

R. E. Thoma, Editor

DATE ISSUED

1964

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

-

-

-

CONTENTS

INTRODUCTION	1
1. METAL SYSTEMS	2
1.1. The System Silver–Zirconium	2
1.2. The System Indium–Zirconium	4
1.3. The System Antimony–Zirconium	6
1.4. The System Lead–Zirconium.....	8
2. METAL–FUSED-SALT SYSTEMS.....	9
2.1. The Sodium Metal–Sodium Halide Systems	9
2.2. The Potassium Metal–Potassium Halide Systems	10
2.3. The Rubidium Metal–Rubidium Halide Systems	11
2.4. The Cesium Metal–Cesium Halide Systems	12
2.5. The Alkali Metal–Alkali Metal Fluoride Systems	13
3. FUSED-SALT SYSTEMS.....	14
3.1. The System LiF–NaF	14
3.2. The System LiF–KF	15
3.3. The System LiF–RbF	16
3.4. The System LiF–CsF	17
3.5. The System NaF–KF	18
3.6. The System NaF–RbF	19
3.7. The System KF–RbF	20
3.8. The System LiF–NaF–KF	21
3.9. The System LiF–NaF–RbF	22
3.10. The System LiF–KF–RbF	23
3.11. The System NaF–KF–RbF	24
3.12. The System NaBF ₄ –KBF ₄	25
3.13. The System NaF–FeF ₂	26
3.14. The System NaF–NiF ₂	27
3.15. The System RbF–CaF ₂	28
3.16. The System LiF–NaF–CaF ₂	29
3.17. The System NaF–MgF ₂ –CaF ₂	30
3.18. The System NaF–KF–AlF ₃	32
3.19. The System LiF–BeF ₂	33
3.20. The System NaF–BeF ₂	34
3.21. The System KF–BeF ₂	36
3.22. The System RbF–BeF ₂	38
3.23. The System CsF–BeF ₂	40
3.24. The System LiF–NaF–BeF ₂	42
3.25. The System LiF–RbF–BeF ₂	44
3.26. The System NaF–KF–BeF ₂	46
3.27. The System NaF–RbF–BeF ₂	47
3.28. The System NaF–ZnF ₂	48
3.29. The System KF–ZnF ₂	49
3.30. The System RbF–ZnF ₂	50
3.31. The System LiF–YF ₃	51
3.32. The System LiF–ZrF ₄	52
3.33. The System NaF–ZrF ₄	54

3.34. The System $\text{KF}-\text{ZrF}_4$	56
3.35. The System $\text{RbF}-\text{ZrF}_4$	57
3.36. The System $\text{CsF}-\text{ZrF}_4$	58
3.37. The System $\text{LiF}-\text{NaF}-\text{ZrF}_4$	60
3.38. The System $\text{NaF}-\text{KF}-\text{ZrF}_4$	62
3.39. The System $\text{NaF}-\text{RbF}-\text{ZrF}_4$	64
3.40. The System $\text{NaF}-\text{CrF}_2-\text{ZrF}_4$: The Section $\text{NaF}\cdot\text{CrF}_2 - \text{ZrF}_4$	66
3.41. The System $\text{NaF}-\text{CeF}_3-\text{ZrF}_4$	67
3.42. The System $\text{LiF}-\text{CeF}_3$	68
3.43. The System $\text{NaF}-\text{HfF}_4$	70
3.44. The System $\text{LiF}-\text{ThF}_4$	72
3.45. The System $\text{NaF}-\text{ThF}_4$	73
3.46. The System $\text{KF}-\text{ThF}_4$	74
3.47. The System $\text{RbF}-\text{ThF}_4$	76
3.48. The System $\text{BeF}_2-\text{ThF}_4$	77
3.49. The System BeF_2-UF_4	78
3.50. The System $\text{MgF}_2-\text{ThF}_4$	79
3.51. The System $\text{LiF}-\text{BeF}_2-\text{ThF}_4$	80
3.52. The System $\text{NaF}-\text{ZrF}_4-\text{ThF}_4$	82
3.53. The System $\text{LiF}-\text{UF}_3$	84
3.54. The System $\text{LiF}-\text{UF}_4$	85
3.55. The System $\text{NaF}-\text{UF}_3$	86
3.56. The System $\text{NaF}-\text{UF}_4$	88
3.57. The System $\text{KF}-\text{UF}_4$	90
3.58. The System $\text{RbF}-\text{UF}_4$	91
3.59. The System $\text{CsF}-\text{UF}_4$	92
3.60. The System ZrF_4-UF_4	93
3.61. The System SnF_2-UF_4	94
3.62. The System PbF_2-UF_4	95
3.63. The System ThF_4-UF_4	96
3.64. The System $\text{LiF}_4-\text{NaF}-\text{UF}_4$	98
3.65. The System $\text{LiF}-\text{KF}-\text{UF}_4$	101
3.66. The System $\text{LiF}-\text{RbF}-\text{UF}_4$	102
3.67. The System $\text{NaF}-\text{KF}-\text{UF}_4$	103
3.68. The System $\text{NaF}-\text{RbF}-\text{UF}_4$	104
3.69. The System $\text{LiF}-\text{BeF}_2-\text{UF}_4$	108
3.70. The System $\text{NaF}-\text{BeF}_2-\text{UF}_4$	110
3.71. The System $\text{NaF}-\text{PbF}_2-\text{UF}_4$	113
3.72. The System $\text{KF}-\text{PbF}_2-\text{UF}_4$	114
3.73. The System $\text{NaF}-\text{ZrF}_4-\text{UF}_4$	116
3.74. The System $\text{LiF}-\text{ThF}_4-\text{UF}_4$	119
3.75. The System $\text{NaF}-\text{ThF}_4-\text{UF}_4$: The Section $2\text{NaF}\cdot\text{ThF}_4-2\text{NaF}\cdot\text{UF}_4$	124
3.76. The System $\text{LiF}-\text{PuF}_3$	125
3.77. The System $\text{LiCl}-\text{FeCl}_2$	126
3.78. The System $\text{KCl}-\text{FeCl}_2$	127
3.79. The System $\text{NaCl}-\text{ZrCl}_4$	128
3.80. The System $\text{KCl}-\text{ZrCl}_4$	130
3.81. The System $\text{LiCl}-\text{UCl}_3$	131
3.82. The System $\text{LiCl}-\text{UCl}_4$	132
3.83. The System $\text{NaCl}-\text{UCl}_3$	133
3.84. The System $\text{NaCl}-\text{UCl}_4$	134

3.85. The System $\text{KCl}-\text{UCl}_4$	135
3.86. The System $\text{RbCl}-\text{UCl}_3$	136
3.87. The System $\text{CsCl}-\text{UCl}_4$	137
3.88. The System $\text{K}_3\text{CrF}_6-\text{Na}_3\text{CrF}_6-\text{Li}_3\text{CrF}_6$	138
4. OXIDE AND HYDROXIDE SYSTEMS	139
4.1. The System $\text{SiO}_2-\text{ThO}_2$	139
4.2. The System $\text{LiOH}-\text{NaOH}$	140
4.3. The System $\text{LiOH}-\text{KOH}$	141
4.4. The System $\text{NaOH}-\text{KOH}$	142
4.5. The System $\text{NaOH}-\text{RbOH}$	143
4.6. The System $\text{Ba}(\text{OH})_2-\text{Sr}(\text{OH})_2$	144
5. AQUEOUS SYSTEMS	145
5.1. The System $\text{UO}_2-\text{P}_2\text{O}_5-\text{H}_2\text{O}$, 25°C Isotherm	145
5.2. The System $\text{UO}_3-\text{H}_3\text{PO}_4-\text{H}_2\text{O}$, 25°C Isotherm	146
5.3. The System $\text{U}(\text{HPO}_4)_2-\text{Cl}_2\text{O}_7-\text{H}_2\text{O}$, 25°C Isotherm	147
5.4. The System $\text{UO}_3-\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, 26°C Isotherm	148
5.5. Portions of the System $\text{ThO}_2-\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, 25°C Isotherm	150
5.6. The System $\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, 25°C Isotherm	153
5.7. Solubility of Uranyl Sulfate in Water	156
5.8. Two-Liquid-Phase Region of UO_2SO_4 in Ordinary and Heavy Water	158
5.9. The System $\text{UO}_3-\text{SO}_3-\text{H}_2\text{O}$	160
5.10. Coexistence Curves for Two Liquid Phases in the System $\text{UO}_2\text{SO}_4-\text{H}_2\text{SO}_4-\text{H}_2\text{O}$	164
5.11. Two-Liquid-Phase Region of the System $\text{UO}_2\text{SO}_4-\text{H}_2\text{SO}_4-\text{H}_2\text{O}$	166
5.12. Two-Liquid-Phase Coexistence Curves for UO_3 -Rich Solutions in the System $\text{UO}_3-\text{SO}_3-\text{H}_2\text{O}$ With and Without HNO_3	167
5.13. Solubility of UO_3 in $\text{H}_2\text{SO}_4-\text{H}_2\text{O}$ Mixtures	168
5.14. Effect of Excess H_2SO_4 on the Phase Equilibria in Very Dilute UO_2SO_4 Solutions	169
5.15. Second-Liquid-Phase Temperatures of $\text{UO}_2\text{SO}_4-\text{Li}_2\text{SO}_4$ Solutions	170
5.16. Second-Liquid-Phase Temperatures for UO_2SO_4 Solutions Containing Li_2SO_4 or BeSO_4	171
5.17. Second-Liquid-Phase Temperatures for BeSO_4 Solutions Containing UO_3	172
5.18. The System $\text{NiSO}_4-\text{UO}_2\text{SO}_4-\text{H}_2\text{O}$ at 25°C	173
5.19. Phase-Transition Temperatures in Solutions Containing CuSO_4 , UO_2SO_4 , and H_2SO_4	174
5.20. The Effect of CuSO_4 and NiSO_4 on Phase-Transition Temperatures: 0.04 <i>m</i> UO_2SO_4 ; 0.01 <i>m</i> H_2SO_4	175
5.21. The System $\text{UO}_3-\text{CuO}-\text{NiO}-\text{SO}_3-\text{D}_2\text{O}$ at 300°C; 0.06 <i>m</i> SO_3	176
5.22. Solubility of $\text{Nd}_2(\text{SO}_4)_3$ in 0.02 <i>m</i> UO_2SO_4 Solution Containing 0.005 <i>m</i> H_2SO_4 (180–300°C)	177
5.23. Solubility of $\text{La}_2(\text{SO}_4)_3$ in UO_2SO_4 Solutions	178
5.24. Solubility of CdSO_4 in UO_2SO_4 Solutions	179
5.25. Solubility of Cs_2SO_4 in UO_2SO_4 Solutions	180
5.26. Solubility of $\text{Y}_2(\text{SO}_4)_3$ in UO_2SO_4 Solutions	181
5.27. Solubility of Ag_2SO_4 in UO_2SO_4 Solutions	182
5.28. Solubility at 250°C of BaSO_4 in $\text{UO}_2\text{SO}_4-\text{H}_2\text{O}$ Solutions	183
5.29. Solubility of H_2WO_4 in 0.126 and 1.26 <i>M</i> UO_2SO_4	184
5.30. Phase Stability of H_2WO_4 in 1.26 <i>M</i> UO_2F_2	186

5.31. The System $\text{UO}_2(\text{NO}_3)_2\text{--H}_2\text{O}$	188
5.32. Phase Equilibria of UO_3 and HF in Stoichiometric Concentrations (Aqueous System).....	190
5.33. Solubility of Uranium Trioxide in Orthophosphoric Acid Solutions	191
5.34. Solubility of Uranium Trioxide in Phosphoric Acid at 250°C	192
5.35. Solubility of UO_3 in H_3PO_4 Solution	193
5.36. The System $\text{UO}_2\text{CrO}_4\text{--H}_2\text{O}$	194
5.37. Variation of Li_2CO_3 Solubility with UO_2CO_3 Concentration at Constant CO_2 Pressure (250°C)	195
5.38. The System $\text{Li}_2\text{O--UO}_3\text{--CO}_2\text{--H}_2\text{O}$ at 250°C and 1500 psi.....	196
5.39. The System $\text{Th}(\text{NO}_3)_4\text{--H}_2\text{O}$	197
5.40. Hydrolytic Stability of Thorium Nitrate–Nitric Acid and Uranyl Nitrate Solutions	198
5.41. The System $\text{ThO}_2\text{--CrO}_3\text{--H}_2\text{O}$ at 25°C	199
5.42. Phase Stability of $\text{ThO}_2\text{--H}_3\text{PO}_4\text{--H}_2\text{O}$ Solutions at High Concentrations of ThO_2	200

PHASE DIAGRAMS OF NUCLEAR REACTOR MATERIALS

INTRODUCTION

This compilation presents the phase diagrams for possible materials for nuclear reactors developed at the Oak Ridge National Laboratory over the period 1950–59. It represents the efforts of certain personnel in what are now the Chemical Technology, Metallurgy, Chemistry, and Reactor Chemistry Divisions of ORNL. This document also contains diagrams originated by other installations under contract to ORNL during that interval. In addition, a few diagrams from the unclassified literature have been included when they form part of completed systems sequences.

No general discussion of the principles of heterogeneous phase equilibrium has been included since excellent discussions are available in many well-known publications. Equilibria in several of the fused-salt systems presented below have been described in some detail in a recent publication by Ricci.¹ Nor has any attempt been made to assemble the phase equilibrium data on reactor materials available from many other sources. For such information the reader is referred to other recently published compilations of phase diagrams^{2–6} and to the several new works dealing specifically with nuclear reactor materials.^{7–9}

While some of the diagrams presented below are the result of fundamental researches, most were obtained in support of the ORNL programs in development of fluid-fueled reactors. Others have been developed as a consequence of ORNL interests in reprocessing of nuclear fuels, reactor metallurgy, production of uranium and thorium from raw materials, and studies of mechanisms of corrosion.

Much of the research effort was devoted to searches for systems of direct use as fluid fuels or blanket materials; unpromising systems were often given only casual attention. The diagrams in this collection, accordingly, vary widely in the degree of completeness of examination. A brief description of the status of the work is presented in each case; in a few cases, where no detailed description of the system has been published, the available data have been included with the diagram.

¹J. E. Ricci, *Guide to the Phase Diagrams of the Fluoride Systems*, ORNL-2396 (Nov. 19, 1958).

²J. F. Hogerton and R. C. Grass (eds.), *The Reactor Handbook*, vol 2, AECD-3646 (May 1955).

³T. Lyman (ed.), *Metals Handbook*, American Society for Metals, Cleveland, Ohio, 1948.

⁴F. A. Rough and A. A. Bauer (eds.), *Constitution of Uranium and Thorium Alloys*, BMI-1300 (June 2, 1958).

⁵E. M. Levin, H. F. McMurdie, and F. P. Hall, *Phase Diagrams for Ceramists*, The American Ceramic Society, Columbus, Ohio, 1956.

⁶E. M. Levin, H. F. McMurdie, and F. P. Hall, *Supplement to Phase Diagrams for Ceramists*, The American Ceramic Society, Columbus, Ohio, 1959.

⁷B. Yates, *Materials for Use in Nuclear Reactors. Information Bibliography*, IGRL-IB/R-15 (2nd ed.), 1958.

⁸H. H. Hausner and S. B. Roboff, *Materials for Nuclear Power Reactors*, Reinhold Publishing Corp., New York, 1955.

⁹B. Kopelman (ed.), *Materials for Nuclear Reactors*, McGraw-Hill Book Co., New York, 1959.

1. METAL SYSTEMS

1.1. The System Silver-Zirconium

J. O. Betterton, Jr., and D. S. Easton, "The Silver-Zirconium System," *Trans. Met. Soc. AIME* 212, 470-75 (1958).

A detailed investigation was made of the phase diagram of silver-zirconium, particularly in the region 0 to 36 at. % Ag. The system was found to be characterized by two intermediate phases, Zr_2Ag and $ZrAg$, and a eutectoid reaction in which the β -zirconium solid solution decomposes into α -zirconium and Zr_2Ag . It was found that impurities in the range 0.05% from the iodide-type zirconium were sufficient to introduce deviations from binary behavior; with partial removal of these impurities an increase in the α -phase solid solubility limit from 0.1 to 1.1 at. % Ag was observed.

Fig. 1.1a. The System Silver-Zirconium in the Eutectoid Region (0-6 at. % Ag).

Fig. 1.1b. The System Silver-Zirconium in the Region 0-36 at. % Ag.

1.2. The System Indium–Zirconium

J. O. Betterton, Jr., and W. K. Noyce, "The Equilibrium Diagram of Indium–Zirconium in the Region 0–26 at. pct In," *Trans. Met. Soc. AIME* **212**, 340–42 (1958).

The zirconium-rich portion of the indium–zirconium phase diagram was determined as a study of the effect of alloying a trivalent B-subgroup element, indium, with zirconium in group IVA. The temperature of the allotropic transition was found to rise with indium, terminating in a peritectoid reaction, $\beta(9.3\% \text{ In}) + \gamma(22.4\% \text{ In}) \rightleftharpoons \alpha(10.1\% \text{ In})$ at 1003°C . In this respect the effect of indium is analogous to that of aluminum in zirconium and titanium alloys.

Fig. 1.2a. The System Indium–Zirconium in the Peritectoid Region (0–13 at. % In).

BLANK

BLANK

Fig. 1.3b. The System Antimony-Zirconium in the Peritectoid Region (0-6 at. % Sb).

1.4. The System Lead-Zirconium

G. D. Kneip, Jr., and J. O. Betterton, Jr., cited by E. T. Hayes and W. L. O'Brien, "Zirconium Equilibrium Diagrams," p 443-83 in *The Metallurgy of Zirconium*, ed. by B. Lustman and F. Kerze, Jr., McGraw-Hill Book Co., New York, 1955.

Fig. 1.4. The System Lead-Zirconium in the Peritectoid Region (0-12 at. % Pb).

2. METAL-FUSED-SALT SYSTEMS

2.1. The Sodium Metal-Sodium Halide Systems

M. A. Bredig and J. E. Sutherland, "High-Temperature Region of the Sodium-Sodium Halide Systems," *Chem. Ann. Prog. Rep.* June 20, 1957, ORNL-2386, p 119. Subsequent report on these systems to be published.

UNCLASSIFIED
ORNL-LR-DWG. 21894A

Fig. 2.1. The Sodium Metal-Sodium Halide Systems.

2.2. The Potassium Metal–Potassium Halide Systems

J. W. Johnson and M. A. Bredig, "Miscibility of Metals with Salts in the Molten State. III. The Potassium–Potassium Halide Systems," *J. Phys. Chem.* **62**, 604 (1958).

Complete miscibility of metal and salt exists in the $KX-K$ systems, $X = F, Cl, Br, \text{ and } I$, at and above 904, 790, 727, and 717°C, that is, as close as +47, +18, -7, and +22°, respectively, to the melting points of the salts. The asymmetry of the liquid-liquid coexistence areas, especially of the fluoride and chloride systems, as indicated by the consolute compositions (20, 39, 44, and 50 mole % metal, respectively), is largely explained by the difference in molar volumes of salt and metal. At the monotectic temperature, the solubility of the solid salts in the liquid metal, as well as of the metal in the liquid salt phase, is much larger than in the case of the sodium systems.

Fig. 2.2. The Potassium Metal–Potassium Halide Systems.

2.3. The Rubidium Metal-Rubidium Halide Systems

M. A. Bredig and J. W. Johnson, "Rubidium-Rubidium Halide Systems," *Chem. Ann. Prog. Rep.* June 20, 1957, ORNL-2386, p 122; M. A. Bredig, J. E. Sutherland, and A. S. Dworkin, "Molten Salt-Metal Solutions. Phase Equilibria," *Chem. Ann. Prog. Rep.* June 20, 1958, ORNL-2584, p 73. Subsequent report on these systems to be published.

Fig. 2.3. The Rubidium Metal-Rubidium Halide Systems.

2.4. The Cesium Metal-Cesium Halide Systems

M. A. Bredig, H. R. Bronstein, and W. T. Smith, Jr., "Miscibility of Liquid Metals with Salts. II. The Potassium-Potassium Fluoride and Cesium-Cesium Halide Systems," *J. Am. Chem. Soc.* **77**, 1454 (1955).

Miscibility in all proportions of cesium metal with cesium halides at and above the melting points of the pure salts, and of potassium metal with potassium fluoride 50° above the melting point of the salt, occurs. The temperature-concentration range of coexistence of two liquid phases decreases in going from sodium to potassium systems and disappears altogether for the cesium systems. The solubility of the solid halides in the corresponding liquid alkali metals, at a given temperature, increases greatly with increase of atomic number of the metal.

Fig. 2.4. The Cesium Metal-Cesium Halide Systems.

2.5. The Alkali Metal-Alkali Metal Fluoride Systems

M. A. Bredig, J. E. Sutherland, and A. S. Dworkin, "Molten Salt-Metal Solutions. Phase Equilibria," *Chem. Ann. Prog. Rep.* June 20, 1958, ORNL-2584, p 73.

Fig. 2.5. The Alkali Metal-Alkali Metal Fluoride Systems.

3. FUSED-SALT SYSTEMS

3.1. The System LiF–NaF

A. G. Bergman and E. P. Dergunov, "Fusion Diagram of LiF–KF–NaF," *Compt. rend. acad. sci. U.R.S.S.* **31**, 753–54 (1941).

The system LiF–NaF contains a single eutectic at 60 LiF–40 NaF (mole %), m.p. 652°C.

Fig. 3.1. The System LiF–NaF.

3.2. The System LiF-KF

A. G. Bergman and E. P. Dergunov, "Fusion Diagram of LiF-KF-NaF," *Compt. rend. acad. sci. U.R.S.S.* **31**, 753-54 (1941).

The system LiF-KF contains a single eutectic at 50 LiF-50 KF (mole %), m.p. 492°C.

Fig. 3.2. The System LiF-KF.

3.3. The System LiF-RbF

C. J. Barton, T. N. McVay, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-54.

Preliminary diagram.

Invariant Equilibria			
Mole % LiF in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
44	470	Eutectic	$L \rightleftharpoons \text{RbF} + \text{LiF} \cdot \text{RbF}$
47	475	Peritectic	$L + \text{LiF} \rightleftharpoons \text{LiF} \cdot \text{RbF}$

General characteristics of the system have been reported by E. P. Dergunov, "Fusion Diagrams of the Ternary Systems of the Fluorides of Lithium, Sodium, Potassium, and Rubidium," *Doklady Akad. Nauk S.S.S.R.* **58**, 1369-72 (1947).

Fig. 3.3. The System LiF-RbF.

3.4. The System LiF–CsF

C. J. Barton, L. M. Bratcher, T. N. McVay, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952–54.

Preliminary diagram.

Invariant Equilibria			
Mole % CsF in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
55	495 ± 5	Peritectic	$\text{LiF} + L \rightleftharpoons \text{LiF} \cdot \text{CsF}$
63	475 ± 5	Eutectic	$L \rightleftharpoons \text{LiF} \cdot \text{CsF} + \text{CsF}$

Fig. 3.4. The System LiF–CsF.

3.5. The System NaF-KF

A. G. Bergman and E. P. Dergunov, "Fusion Diagram of LiF-KF-NaF," *Compt. rend. acad. sci. U.R.S.S.* **31**, 753-54 (1941).

The system NaF-KF contains a single eutectic at 40 NaF-60 KF (mole %), m.p. 710°C.

Fig. 3.5. The System NaF-KF.

3.6. The System NaF-RbF

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system NaF-RbF contains a single eutectic at 27 NaF-73 RbF (mole %), m.p. $675 \pm 10^\circ\text{C}$.

General characteristics of the system have been reported by E. P. Dergunov, "Fusion Diagrams of the Ternary Systems of the Fluorides of Lithium, Sodium, Potassium, and Rubidium," *Doklady Akad. Nauk S.S.S.R.* **58**, 1369-72 (1947).

Fig. 3.6. The System NaF-RbF.

3.7. The System KF-RbF

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system KF-RbF contains a solid solution minimum at 28 KF-72 RbF (mole %), m.p. $770 \pm 10^\circ\text{C}$.

Fig. 3.7. The System KF-RbF.

3.8. The System LiF-NaF-KF

A. G. Bergman and E. P. Dergunov, "Fusion Diagram of LiF-KF-NaF," *Compt. rend. acad. sci. U.R.S.S.* 31, 753-54 (1941).

The system LiF-NaF-KF contains a single eutectic at 46.5 LiF-11.5 NaF-42.0 KF (mole %), m.p. 454°C.

UNCLASSIFIED
ORNL-LR-DWG 1199A

Fig. 3.8. The System LiF-NaF-KF.

3.9. The System LiF-NaF-RbF

C. J. Barton, L. M. Bratcher, J. P. Blakely, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system LiF-NaF-RbF contains a single eutectic at 42 LiF-6 NaF-52 RbF (mole %), m.p. $430 \pm 10^\circ\text{C}$. The composition and temperature of the ternary peritectic point have not yet been determined.

General characteristics of the system have been reported by E. P. Dergunov, "Fusion Diagrams of the Ternary systems of the Fluorides of Lithium, Sodium, Potassium, and Rubidium," *Doklady Akad. Nauk S.S.S.R.* **58**, 1369-72 (1947). Dergunov lists the composition and temperature of the ternary eutectic as 46.5 LiF-6.5 NaF-47 RbF (mole %), m.p. 426°C .

UNCLASSIFIED
ORNL-LR-DWG 1200A

Fig. 3.9. The System LiF-NaF-RbF.

3.10. The System LiF-KF-RbF

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system LiF-KF-RbF contains a single eutectic at 40 LiF-32.5 KF-27.5 RbF (mole %), $m.p. 440 \pm 10^\circ\text{C}$.

General characteristics of the system have been reported by E. P. Dergunov, "Fusion Diagrams of the Ternary Systems of the Fluorides of Lithium, Sodium, Potassium, and Rubidium," *Doklady Akad. Nauk S.S.S.R.* **58**, 1369-72 (1947).

Fig. 3.10. The System LiF-KF-RbF.

3.11. The System NaF-KF-RbF

C. J. Barton, L. M. Bratcher, J. P. Blakely, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system NaF-KF-RbF contains a single eutectic at 21 NaF-5 KF-74 RbF (mole %), m.p. $621 \pm 10^\circ\text{C}$.

Fig. 3.11. The System NaF-KF-RbF.

3.12. The System NaBF_4 - KBF_4

R. E. Moore, J. G. Surak, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1957.

Preliminary diagram. The system NaBF_4 - KBF_4 contains a single eutectic at 88 NaBF_4 -12 KBF_4 (mole %), m.p. 355°C . The dotted line was obtained from thermal effects representing incompletely understood solid-phase transformations of KBF_4 and NaBF_4 .

Fig. 3.12. The System NaBF_4 - KBF_4 .

3.13. The System NaF-FeF₂

R. E. Thoma, H. A. Friedman, B. S. Landau, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1957.

Preliminary diagram.

Invariant Equilibria

Mole % FeF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
30	680	Eutectic	$L \rightleftharpoons \text{NaF} + \text{NaF} \cdot \text{FeF}_2$
50	783	Congruent melting point	$L \rightleftharpoons \text{NaF} \cdot \text{FeF}_2$
63	745	Eutectic	$L \rightleftharpoons \text{NaF} \cdot \text{FeF}_2 + \text{FeF}_2$

Fig. 3.13. The System NaF-FeF₂.

3.14. The System NaF–NiF₂

R. E. Thoma, H. A. Friedman, B. S. Landau, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1957.

Preliminary diagram.

Invariant Equilibria			
Mole % NiF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
23	795	Eutectic	$L \rightleftharpoons \text{NaF} + \text{NaF} \cdot \text{NiF}_2$
50	1045	Congruent melting point	$L \rightleftharpoons \text{NaF} \cdot \text{NiF}_2$
57	1040	Eutectic	$L \rightleftharpoons \text{NaF} \cdot \text{NiF}_2 + \text{NiF}_2$

Fig. 3.14. The System NaF–NiF₂

3.15. The System RbF-CaF₂

C. J. Barton, L. M. Bratcher, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956.

Preliminary diagram.

Invariant Equilibria

Mole % CaF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
9	760	Eutectic	$L \rightleftharpoons \text{RbF} + \text{RbF} \cdot \text{CaF}_2$
50	1110	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot \text{CaF}_2$
57	1090	Eutectic	$L \rightleftharpoons \text{RbF} \cdot \text{CaF}_2 + \text{CaF}_2$

Fig. 3.15. The System RbF-CaF₂

3.16. The System LiF-NaF-CaF₂

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram. The system LiF-NaF-CaF₂ contains a single eutectic at 53 LiF-36 NaF-11 CaF₂ (mole %), m.p. 616°C.

UNCLASSIFIED
ORNL-LR-DWG 12779R

Fig. 3.16. The System LiF-NaF-CaF₂

3.17. The System NaF-MgF₂-CaF₂

C. J. Barton, L. M. Bratcher, J. P. Blakely, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Temperature
NaF	CaF ₂	MgF ₂			
65	23	12	745	Eutectic	NaF, NaF•MgF ₂ , and CaF ₂
35	28	37	905	Eutectic	CaF ₂ , MgF ₂ , and NaF•MgF ₂

Fig. 3.17a. The System NaF-MgF₂-CaF₂

The minimum temperature along the quasi-binary section $\text{NaF} \cdot \text{MgF}_2 - \text{CaF}_2$ occurs at the composition 36 NaF –36 MgF_2 –28 CaF_2 (mole %), at 910°C.

The existence of the compound $\text{NaF} \cdot \text{MgF}_2$ was previously reported by A. G. Bergman and E. P. Dergunov, *Compt. rend. acad. sci. U.R.S.S.* **31**, 755 (1941).

Ternary systems of alkali fluorides with MgF_2 have been reported by Bergman *et al.* as follows: the system $\text{LiF} - \text{NaF} - \text{MgF}_2$ by A. G. Bergman and E. P. Dergunov, *Compt. rend. acad. sci. U.R.S.S.* **31**, 755 (1941); the system $\text{LiF} - \text{KF} - \text{MgF}_2$ by A. G. Bergman and S. P. Parlenko, *Compt. rend. acad. sci. U.R.S.S.* **31**, 818–19 (1941); the system $\text{NaF} - \text{KF} - \text{MgF}_2$ by A. G. Bergman and E. P. Dergunov, *Compt. rend. acad. sci. U.R.S.S.* **48**, 330 (1945).

Fig. 3.17b. The Subsystem $\text{NaF} \cdot \text{MgF}_2 - \text{CaF}_2$

3.18. The System NaF-KF-AlF₃

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-52.

Preliminary diagram. No study has been made at the Oak Ridge National Laboratory of the phase relationships occurring within the system NaF-KF-AlF₃. Phase diagrams have been reported for the AlF₃ binary systems, NaF-AlF₃ [P. P. Fedotieff and W. P. Iljinsky, *Z. anorg. Chem.* **80**, 121 (1913); also N. A. Pushin and A. V. Baskow, *ibid.* **81**, 350 (1913)] and KF-AlF₃ [P. P. Fedotieff and K. Timofeef, *Z. anorg. u. allgem. Chem.* **206**, 265 (1932)].

Fig. 3.18. The System NaF-KF-AlF₃.

3.19. The System LiF-BeF₂

This phase diagram is a composite from several published sources¹⁻³ and from unpublished data derived at the Oak Ridge National Laboratory (R. E. Moore, C. J. Barton, R. E. Thoma, and T. N. McVay) and at the Mound Laboratory (J. F. Eichelberger, C. R. Hudgens, L. V. Jones, and T. B. Rhinehammer). Thermal gradient quenching data (ORNL) and differential thermal analysis data (Mound Laboratory) have served to corroborate this composite diagram.

Invariant Equilibria

Mole % BeF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
33.5*	454	Peritectic	$L + \text{LiF} \rightleftharpoons 2\text{LiF} \cdot \text{BeF}_2$
52	355	Eutectic	$L \rightleftharpoons 2\text{LiF} \cdot \text{BeF}_2 + \text{BeF}_2$
—	280	Upper temperature of stability for LiF·BeF ₂	$2\text{LiF} \cdot \text{BeF}_2 + \text{BeF}_2 \rightleftharpoons \text{LiF} \cdot \text{BeF}_2$

*Ref 3.

Roy, Roy, and Osborn have reported the presence of the compound LiF·2BeF₂ in the system LiF-BeF₂ although neither the Mound Laboratory nor the ORNL data have indicated the existence of this compound.

¹D. M. Roy, R. Roy, and E. F. Osborn, *J. Am. Ceram. Soc.* 37, 300 (1954).

²A. V. Novoselova, Yu. P. Simanov, and E. I. Yarembash, *J. Phys. Chem. (U.S.S.R.)* 26, 1244 (1952).

³John L. Speirs, *The Binary and Ternary Systems Formed by Calcium Fluoride, Lithium Fluoride, and Beryllium Fluoride: Phase Diagrams and Electrolytic Studies*, Ph.D. thesis, University of Michigan, May 29, 1952.

Fig. 3.19. The System LiF-BeF₂.

3.20. The System NaF–BeF₂

D. M. Roy, R. Roy, and E. F. Osborn, "Fluoride Model Systems: III. The System NaF–BeF₂ and the Polymorphism of Na₂BeF₄ and BeF₂," *J. Am. Ceram. Soc.* **36**, 185 (1953).

Invariant Equilibria*			
Mole % BeF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
31	570	Eutectic	$L \rightleftharpoons \text{NaF} + \alpha\text{-}2\text{NaF}\cdot\text{BeF}_2$
33.3	600	Congruent melting point	$L \rightleftharpoons \alpha\text{-}2\text{NaF}\cdot\text{BeF}_2$
–	320	Inversion	$\alpha\text{-}2\text{NaF}\cdot\text{BeF}_2 \rightleftharpoons \alpha'\text{-}2\text{NaF}\cdot\text{BeF}_2$
–	225	Inversion	$\alpha'\text{-}2\text{NaF}\cdot\text{BeF}_2 \rightleftharpoons \gamma\text{-}2\text{NaF}\cdot\text{BeF}_2$
43	340	Eutectic	$L \rightleftharpoons \alpha\text{-}2\text{NaF}\cdot\text{BeF}_2 + \beta\text{-NaF}\cdot\text{BeF}_2$
50	376 ± 5	Congruent melting point	$L \rightleftharpoons \beta'\text{-NaF}\cdot\text{BeF}_2$
55	365	Eutectic	$L \rightleftharpoons \beta'\text{-NaF}\cdot\text{BeF}_2 + \text{BeF}_2$

*Roy, Roy, and Osborn did not list invariant equilibria; those shown are estimates made from the reported work and from experiments performed at the Oak Ridge National Laboratory.

Fig. 3.20. The System NaF-BeF₂

3.21. The System $\text{KF}-\text{BeF}_2$

R. E. Moore, C. J. Barton, L. M. Bratcher, T. N. McVay, G. D. White, R. J. Sheil, W. R. Grimes, R. E. Meadows, and L. A. Harris, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram.

Invariant Equilibria

Mole % BeF_2 In Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
19	720	Eutectic	$L \rightleftharpoons \text{KF} + 3\text{KF} \cdot \text{BeF}_2$
25	740	Congruent melting point	$L \rightleftharpoons 3\text{KF} \cdot \text{BeF}_2$
27	730	Eutectic	$L \rightleftharpoons 3\text{KF} \cdot \text{BeF}_2 + \alpha\text{-}2\text{KF} \cdot \text{BeF}_2$
33.3	787	Congruent melting point	$L \rightleftharpoons \alpha\text{-}2\text{KF} \cdot \text{BeF}_2$
—	685	Inversion	$\alpha\text{-}2\text{KF} \cdot \text{BeF}_2 \rightleftharpoons \beta\text{-}2\text{KF} \cdot \text{BeF}_2$
52	390	Peritectic	$\beta\text{-}2\text{KF} \cdot \text{BeF}_2 + L \rightleftharpoons \text{KF} \cdot \text{BeF}_2$
59	330	Eutectic	$L \rightleftharpoons \text{KF} \cdot \text{BeF}_2 + \beta\text{-KF} \cdot 2\text{BeF}_2$
66.7	358	Congruent melting point	$L \rightleftharpoons \alpha\text{-KF} \cdot 2\text{BeF}_2$
—	334	Inversion	$\alpha\text{-KF} \cdot 2\text{BeF}_2 \rightleftharpoons \beta\text{-KF} \cdot 2\text{BeF}_2$
72.5	323	Eutectic	$L \rightleftharpoons \beta\text{-KF} \cdot 2\text{BeF}_2 + \text{BeF}_2$

This system has been reported by M. P. Borzenkova, A. V. Novoselova, Yu. P. Simanov, V. I. Chernikh, and E. I. Yarembash, "Thermal and X-Ray Analysis of the $\text{KF}-\text{BeF}_2$ System," *Zhur. Neorg. Khim.* **1**, 2071-82 (1956).

Fig. 3.21. The System KF-BeF₂.

3.22. The System RbF-BeF₂

R. E. Moore, C. J. Barton, L. M. Bratcher, T. N. McVay, G. D. White, R. J. Sheil, W. R. Grimes, and R. E. Meadows, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram.

Invariant Equilibria

Mole % BeF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
16	675	Eutectic	$L \rightleftharpoons \text{RbF} + 3\text{RbF} \cdot \text{BeF}_2$
25	725	Congruent melting point	$L \rightleftharpoons 3\text{RbF} \cdot \text{BeF}_2$
27	720	Eutectic	$L \rightleftharpoons 3\text{RbF} \cdot \text{BeF}_2 + 2\text{RbF} \cdot \text{BeF}_2$
33.3	800	Congruent melting point	$L \rightleftharpoons 2\text{RbF} \cdot \text{BeF}_2$
50.5	442	Peritectic	$L + 2\text{RbF} \cdot \text{BeF}_2 \rightleftharpoons \text{RbF} \cdot \text{BeF}_2$
61	383	Eutectic	$L \rightleftharpoons \text{RbF} \cdot \text{BeF}_2 + \text{RbF} \cdot 2\text{BeF}_2$
66.7	464	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot 2\text{BeF}_2$
81	397	Eutectic	$L \rightleftharpoons \text{RbF} \cdot 2\text{BeF}_2 + \text{BeF}_2$

This system has also been reported by R. G. Grebenshchikov, "Investigation of the Phase Diagram of the RbF-BeF₂ System and of Its Relationship to the BaO-SiO₂ System," *Doklady Akad. Nauk S.S.S.R.* **114**, 316 (1957).

Fig. 3.22. The System $\text{RbF}-\text{BeF}_2$

3.23. The System CsF-BeF₂

O. N. Breusov, A. V. Novoselova, and Yu. P. Simanov, "Thermal and X-Ray Phase Analysis of the System CsF-BeF₂ and Its Interrelationships with MeF and BeF₂ Systems," *Doklady Akad. Nauk S.S.S.R.* **118**, 935-37 (1958).

Invariant Equilibria

Mole % BeF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
14	598	Eutectic	$L \rightleftharpoons \text{CsF} + \beta\text{-3CsF}\cdot\text{BeF}_2$
23.5	659	Peritectic	$\alpha\text{-2CsF}\cdot\text{BeF}_2 + L \rightleftharpoons \alpha\text{-3CsF}\cdot\text{BeF}_2$
—	617	Inversion	$\alpha\text{-3CsF}\cdot\text{BeF}_2 \rightleftharpoons \beta\text{-3CsF}\cdot\text{BeF}_2$
33.3	793	Congruent melting point	$L \rightleftharpoons \alpha\text{-2CsF}\cdot\text{BeF}_2$
—	404	Inversion	$\alpha\text{-2CsF}\cdot\text{BeF}_2 \rightleftharpoons \beta\text{-2CsF}\cdot\text{BeF}_2$
48	449	Eutectic	$L \rightleftharpoons \alpha\text{-2CsF}\cdot\text{BeF}_2 + \alpha\text{-CsF}\cdot\text{BeF}_2$
50	475	Congruent melting point	$L \rightleftharpoons \alpha\text{-CsF}\cdot\text{BeF}_2$
—	360	Inversion	$\alpha\text{-CsF}\cdot\text{BeF}_2 \rightleftharpoons \beta\text{-CsF}\cdot\text{BeF}_2$
—	140	Inversion	$\beta\text{-CsF}\cdot\text{BeF}_2 \rightleftharpoons \gamma\text{-CsF}\cdot\text{BeF}_2$
58.4	393	Eutectic	$L \rightleftharpoons \alpha\text{-CsF}\cdot\text{BeF}_2 + \beta\text{-CsF}\cdot 2\text{BeF}_2$
66.7	480	Congruent melting point	$L \rightleftharpoons \alpha\text{-CsF}\cdot 2\text{BeF}_2$
—	450	Inversion	$\alpha\text{-CsF}\cdot 2\text{BeF}_2 \rightleftharpoons \beta\text{-CsF}\cdot 2\text{BeF}_2$
77.5	367	Eutectic	$L \rightleftharpoons \beta\text{-CsF}\cdot 2\text{BeF}_2 + \text{BeF}_2$

Fig. 3.23. The System CsF-BeF₂

3.24. The System LiF–NaF–BeF₂

R. E. Moore, C. J. Barton, W. R. Grimes, R. E. Meadows, L. M. Bratcher, G. D. White, T. N. McVay, and L. A. Harris, unpublished work performed at the Oak Ridge National Laboratory, 1951–58.

Preliminary diagram.

Invariant Equilibria*

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
LiF	NaF	BeF ₂			
15	58	27	480	Eutectic	NaF, LiF, and 2NaF·BeF ₂
23	41	36	328	Eutectic	LiF, 2NaF·BeF ₂ , and 2NaF·LiF·2BeF ₂
20	40	40	355	Congruent melting point	2NaF·LiF·2BeF ₂
5	53	42	318	Eutectic	NaF·BeF ₂ , 2NaF·BeF ₂ , and 2NaF·LiF·2BeF ₂
31.5	31	37.5	315	Eutectic	2LiF·BeF ₂ , LiF, and 2NaF·LiF·2BeF ₂

*Invariant equilibria shown in the phase diagram by the intersections of dotted-line boundary curves have not been determined with sufficient precision to be listed in the table.

Minimum Temperature on Alkemade Lines

Composition of Liquid (mole %)			Temperature (°C)	Alkemade Line
LiF	NaF	BeF ₂		
16	56	28	485	2NaF·BeF ₂ –LiF
26	37	37	340	2NaF·LiF·2BeF ₂ –LiF
11	44	45	332	NaF·BeF ₂ –2NaF·LiF·2BeF ₂
16	45	39	343	2NaF·BeF ₂ –2NaF·LiF·2BeF ₂
30.5	31	38.5	316	2LiF·BeF ₂ –2NaF·LiF·2BeF ₂

Phase relationships of two of the ternary compounds in this system have been reported by W. Jahn [“Silicate Models. V. NaLi(BeF₄), a Model Substance for Monticellite, CaMg(SiO₄)₂,” *Z. anorg. u. allgem. Chem.* **276**, 113–27 (1954); “Silicate Models. VI. Na₃Li(BeF₄)₂, a New Compound in the Ternary System NaF–LiF–BeF₂, and Its Relation to Merwinite, Ca₃Mg(SiO₄)₂,” *Z. anorg. u. allgem. Chem.* **277**, 274–86 (1954)].

Fig. 3.24. The System LiF-NaF-BeF₂

3.25. The System LiF–RbF–BeF₂

T. B. Rhinehammer, D. E. Etter, C. R. Hudgens, N. E. Rogers, and P. A. Tucker, unpublished work performed at the Mound Laboratory.

Preliminary diagram.

Invariant Equilibria and Singular Points

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solid Phases Present
LiF	RbF	BeF ₂			
40.0	20.0	40.0	485 ± 5	Congruent melting point	2LiF·RbF·2BeF ₂
33.3	33.3	33.3	544 ± 5	Incongruent melting point of LiF·RbF·BeF ₂	
33.0	8.0	59.0	295 ± 5	Eutectic	2LiF·BeF ₂ , 2LiF·RbF·2BeF ₂ , and BeF ₂
26.0	12.0	62.0	305–320	Peritectic	2LiF·RbF·2BeF ₂ , RbF·2BeF ₂ , and BeF ₂
12.0	29.0	59.0	334 ± 5	Quasi-binary eutectic	2LiF·RbF·2BeF ₂ and RbF·2BeF ₂
11.0	34.0	55.0	300 ± 5	Eutectic	2LiF·RbF·2BeF ₂ , RbF·BeF ₂ , and RbF·2BeF ₂
11.5	35.5	53.0	335 ± 5	Peritectic	2LiF·RbF·2BeF ₂ , LiF·RbF·BeF ₂ , and RbF·BeF ₂
9.0	40.0	51.0	380 ± 5	Peritectic	LiF·RbF·BeF ₂ , 2RbF·BeF ₂ , and RbF·BeF ₂
50.0	8.0	42.0	426 ± 5	Peritectic	LiF, 2LiF·RbF·2BeF ₂ , and 2LiF·BeF ₂
41.5	19.5	39.0	473 ± 5	Quasi-binary eutectic	LiF and 2LiF·RbF·2BeF ₂
35.0	25.5	39.5	470 ± 5	Peritectic	LiF, 2LiF·RbF·2BeF ₂ , and LiF·RbF·BeF ₂
23.0	40.0	37.0	530 ± 5	Peritectic	LiF, 2RbF·BeF ₂ , and LiF·RbF·BeF ₂
30.0	35.0	35.0	544 ± 5	Maximum on the boundary	LiF and LiF·RbF·BeF ₂
41.0	39.5	19.5	640 ± 5	Quasi-binary eutectic	LiF and 2RbF·BeF ₂
43.0	50.0	7.0	527 ± 5	Peritectic	LiF, 3RbF·BeF ₂ , and 2RbF·BeF ₂
43.0	54.0	3.0	470 ± 5	Peritectic	LiF, LiF·RbF, and 3RbF·BeF ₂
43.0	55.0	2.0	462 ± 5	Eutectic	LiF·RbF, RbF, and 3RbF·BeF ₂

Fig. 3.25. The System LiF-RbF-BeF_2 .

3.26. The System NaF-KF-BeF₂

R. E. Moore, C. J. Barton, L. M. Bratcher, T. N. McVay, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952-55.

Preliminary diagram. Phase relationships within the system NaF-KF-BeF₂ have not yet become so well defined at the Oak Ridge National Laboratory that the compositions and temperatures of the invariant points may be listed.

UNCLASSIFIED
ORNL-LR-DWG 38114

Fig. 3.26. The System NaF-KF-BeF₂

3.27. The System NaF-RbF-BeF₂

R. E. Moore, C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1954-57.

Preliminary diagram.

Invariant Equilibria and Singular Points

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Temperature
NaF	RbF	BeF ₂			
20.5	68.5	11.0	610	Eutectic	RbF, NaF, and 3RbF·BeF ₂
50	37.5	12.5	640	Quasi-binary eutectic	3RbF·BeF ₂ and NaF
46	38	16	635	Eutectic	3RbF·BeF ₂ , 2RbF·BeF ₂ , and NaF
45	37	18	650	Quasi-binary eutectic	2RbF·BeF ₂ and NaF
37	33	30	570	Peritectic	2RbF·BeF ₂ , NaF, and 2NaF·4RbF·3BeF ₂
52	15	33	477	Eutectic	2NaF·4RbF·3BeF ₂ , NaF, and 2NaF·BeF ₂
33	33.7	33.3	580	Peritectic	2RbF·BeF ₂ and 2NaF·4RbF·3BeF ₂
51.7	15	33.3	480	Quasi-binary eutectic	2NaF·4RbF·3BeF ₂ and 2NaF·BeF ₂

Fig. 3.27. The System NaF-RbF-BeF₂.

3.28. The System NaF–ZnF₂

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952.

Preliminary diagram.

Invariant Equilibria			
Mole % ZnF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
33	635	Eutectic	$L \rightleftharpoons \text{NaF} + \text{NaF} \cdot \text{ZnF}_2$
50	748 ± 10	Congruent melting point	$L \rightleftharpoons \text{NaF} \cdot \text{ZnF}_2$
69	685	Eutectic	$L \rightleftharpoons \text{NaF} \cdot \text{ZnF}_2 + \text{ZnF}_2$

Fig. 3.28. The System NaF–ZnF₂

3.29. The System $\text{KF}-\text{ZnF}_2$

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952.

Preliminary diagram.

Invariant Equilibria

Mole % ZnF_2 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
21	670	Eutectic	$L \rightleftharpoons \text{KF} + 2\text{KF} \cdot \text{ZnF}_2$
30	720 ± 10	Peritectic	$L + \text{KF} \cdot \text{ZnF}_2 \rightleftharpoons 2\text{KF} \cdot \text{ZnF}_2$
50	850 ± 10	Congruent melting point	$L \rightleftharpoons \text{KF} \cdot \text{ZnF}_2$
80	740 ± 5	Eutectic	$L \rightleftharpoons \text{KF} \cdot \text{ZnF}_2 + \text{ZnF}_2$

Fig. 3.29. The System $\text{KF}-\text{ZnF}_2$

3.30. The System RbF–ZnF₂

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952.

Preliminary diagram.

Invariant Equilibria			
Mole % ZnF ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
21	595 ± 10	Eutectic	$L \rightleftharpoons \text{RbF} + 2\text{RbF} \cdot \text{ZnF}_2$
32	620 ± 10	Peritectic	$L + \text{RbF} \cdot \text{ZnF}_2 \rightleftharpoons 2\text{RbF} \cdot \text{ZnF}_2$
50	730 ± 10	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot \text{ZnF}_2$
70	650	Eutectic	$L \rightleftharpoons \text{RbF} \cdot \text{ZnF}_2 + \text{ZnF}_2$

Fig. 3.30. The System RbF–ZnF₂

3.31. The System LiF-YF₃

R. E. Thoma, C. F. Weaver, and H. A. Friedman, unpublished work performed at the Oak Ridge National Laboratory, 1958-59.

Preliminary diagram.

Invariant Equilibria			
Mole % YF ₃ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
19	695	Eutectic	$L \rightleftharpoons \text{LiF} + \text{LiF} \cdot \text{YF}_3$
49	815	Peritectic	$L + \text{YF}_3 \rightleftharpoons \text{LiF} \cdot \text{YF}_3$

Fig. 3.31. The System LiF-YF₃.

3.32. The System LiF–ZrF₄

R. E. Moore, F. F. Blankenship, W. R. Grimes, H. A. Friedman, C. J. Barton, R. E. Thoma, and H. Insley, unpublished work performed at the Oak Ridge National Laboratory, 1951–56. Preliminary diagram.

Invariant Equilibria

Mole % ZrF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
21	598	Eutectic	$L \rightleftharpoons \text{LiF} + \alpha\text{-3LiF}\cdot\text{ZrF}_4$
25	662	Congruent melting point	$L \rightleftharpoons \alpha\text{-3LiF}\cdot\text{ZrF}_4$
–	475	Inversion	$\alpha\text{-3LiF}\cdot\text{ZrF}_4 \rightleftharpoons \beta\text{-3LiF}\cdot\text{ZrF}_4$
–	470	Decomposition	$\beta\text{-3LiF}\cdot\text{ZrF}_4 \rightleftharpoons \text{LiF} + 2\text{LiF}\cdot\text{ZrF}_4$
29.5	570	Eutectic	$L \rightleftharpoons \alpha\text{-3LiF}\cdot\text{ZrF}_4 + 2\text{LiF}\cdot\text{ZrF}_4$
33.3	596	Congruent melting point	$L \rightleftharpoons 2\text{LiF}\cdot\text{ZrF}_4$
49	507	Eutectic	$L \rightleftharpoons 2\text{LiF}\cdot\text{ZrF}_4 + 3\text{LiF}\cdot 4\text{ZrF}_4$
51.5	520	Peritectic	$L + \text{ZrF}_4 \rightleftharpoons 3\text{LiF}\cdot 4\text{ZrF}_4$
–	466	Decomposition	$3\text{LiF}\cdot 4\text{ZrF}_4 \rightleftharpoons 2\text{LiF}\cdot\text{ZrF}_4 + \text{ZrF}_4$

Fig. 3.32. The System LiF-ZrF₄.

3.33. The System NaF–ZrF₄

C. J. Barton, W. R. Grimes, H. Insley, R. E. Moore, and R. E. Thoma, "Phase Equilibria in the Systems NaF–ZrF₄, UF₄–ZrF₄, and NaF–ZrF₄–UF₄," *J. Phys. Chem.* **62**, 665–76 (1958).

Invariant Equilibria

Mole % ZrF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
20	747	Eutectic	$L \rightleftharpoons \text{NaF} + 3\text{NaF} \cdot \text{ZrF}_4$
25	850	Congruent melting point	$L \rightleftharpoons 3\text{NaF} \cdot \text{ZrF}_4^*$
–	523	Inversion	$\alpha\text{-}5\text{NaF} \cdot 2\text{ZrF}_4 \rightleftharpoons \beta\text{-}5\text{NaF} \cdot 2\text{ZrF}_4$
30.5	500	Eutectoid	$\alpha\text{-}5\text{NaF} \cdot 2\text{ZrF}_4^{ss} \rightleftharpoons \beta\text{-}5\text{NaF} \cdot 2\text{ZrF}_4 + \gamma\text{-}2\text{NaF} \cdot \text{ZrF}_4$
34	640	Peritectic	$L + 3\text{NaF} \cdot \text{ZrF}_4(ss, \text{ca. } 27.5 \text{ mole \% ZrF}_4) \rightleftharpoons \alpha\text{-}5\text{NaF} \cdot 2\text{ZrF}_4$
39.5	544	Peritectic	$L + \alpha\text{-}5\text{NaF} \cdot 2\text{ZrF}_4(ss, 30 \text{ mole \% ZrF}_4) \rightleftharpoons \alpha\text{-}2\text{NaF} \cdot \text{ZrF}_4$
–	533	Inversion	$\alpha\text{-}2\text{NaF} \cdot \text{ZrF}_4 \rightleftharpoons \beta\text{-}2\text{NaF} \cdot \text{ZrF}_4$
–	505	Inversion	$\beta\text{-}2\text{NaF} \cdot \text{ZrF}_4 \rightleftharpoons \gamma\text{-}2\text{NaF} \cdot \text{ZrF}_4$
40.5	500	Eutectic	$L \rightleftharpoons \gamma\text{-}2\text{NaF} \cdot \text{ZrF}_4 + 7\text{NaF} \cdot 6\text{ZrF}_4^{ss}$
46.2	525	Congruent melting point	$L \rightleftharpoons 7\text{NaF} \cdot 6\text{ZrF}_4$
49.5	512	Eutectic	$L \rightleftharpoons 7\text{NaF} \cdot 6\text{ZrF}_4 + 3\text{NaF} \cdot 4\text{ZrF}_4$
56.5	537	Peritectic	$L + \text{ZrF}_4 \rightleftharpoons 3\text{NaF} \cdot 4\text{ZrF}_4$

*A determination of the crystal structure of 3NaF·ZrF₄ has been reported by L. A. Harris, "The Crystal Structures of Na₃ZrF₇ and Na₃HfF₇," *Acta Cryst.* **12**, 172 (1959).

Fig. 3.33. The System NaF-ZrF₄.

3.34. The System $\text{KF}-\text{ZrF}_4$

C. J. Barton, H. Insley, R. P. Metcalf, R. E. Thoma, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-55.

Preliminary diagram.

Invariant Equilibria			
Mole % ZrF_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
14	765	Eutectic	$L \rightleftharpoons \text{KF} + 3\text{KF} \cdot \text{ZrF}_4$
25	910	Congruent melting point	$L \rightleftharpoons 3\text{KF} \cdot \text{ZrF}_4$
36	590	Peritectic	$3\text{KF} \cdot \text{ZrF}_4 + L \rightleftharpoons 2\text{KF} \cdot \text{ZrF}_4$
40.5	412	Peritectic	$2\text{KF} \cdot \text{ZrF}_4 + L \rightleftharpoons 3\text{KF} \cdot 2\text{ZrF}_4$
42	390	Eutectic	$L \rightleftharpoons 3\text{KF} \cdot 2\text{ZrF}_4 + \text{KF} \cdot \text{ZrF}_4$
50	475	Congruent melting point	$L \rightleftharpoons \text{KF} \cdot \text{ZrF}_4$
55	440	Eutectic	$L \rightleftharpoons \text{KF} \cdot \text{ZrF}_4 + \text{ZrF}_4$

Fig. 3.34. The System $\text{KF}-\text{ZrF}_4$.

3.35. The System RbF-ZrF₄

R. E. Moore, R. E. Thoma, C. J. Barton, W. R. Grimes, H. Insley, B. S. Landau, and H. A. Friedman, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram.

Invariant Equilibria

Mole % ZrF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
10	710	Eutectic	$L \rightleftharpoons \text{RbF} + 3\text{RbF} \cdot \text{ZrF}_4$
25	897	Congruent melting point	$L \rightleftharpoons 3\text{RbF} \cdot \text{ZrF}_4$
34	620	Peritectic	$L + 3\text{RbF} \cdot \text{ZrF}_4 \rightleftharpoons \alpha\text{-}2\text{RbF} \cdot \text{ZrF}_4$
—	460	Inversion	$\alpha\text{-}2\text{RbF} \cdot \text{ZrF}_4 \rightleftharpoons \beta\text{-}2\text{RbF} \cdot \text{ZrF}_4$
—	370	Lowered inversion and decomposition	$\alpha\text{-}2\text{RbF} \cdot \text{ZrF}_4 \rightleftharpoons \beta\text{-}2\text{RbF} \cdot \text{ZrF}_4$
42	410	Eutectic	$L \rightleftharpoons \alpha\text{-}2\text{RbF} \cdot \text{ZrF}_4 + 5\text{RbF} \cdot 4\text{ZrF}_4$
44.4	445	Congruent melting point	$L \rightleftharpoons 5\text{RbF} \cdot 4\text{ZrF}_4$
48	390	Eutectic	$L \rightleftharpoons 5\text{RbF} \cdot 4\text{ZrF}_4 + \beta\text{-RbF} \cdot \text{ZrF}_4$
50	423	Congruent melting point	$L \rightleftharpoons \alpha\text{-RbF} \cdot \text{ZrF}_4$
—	391	Inversion	$\alpha\text{-RbF} \cdot \text{ZrF}_4 \rightleftharpoons \beta\text{-RbF} \cdot \text{ZrF}_4$
54	400	Eutectic	$L \rightleftharpoons \alpha\text{-RbF} \cdot \text{ZrF}_4 + \text{RbF} \cdot 2\text{ZrF}_4$
57	447	Peritectic	$L + \text{ZrF}_4 \rightleftharpoons \text{RbF} \cdot 2\text{ZrF}_4$

Fig. 3.35. The System RbF-ZrF₄.

3.36. The System CsF–ZrF₄

C. J. Barton, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1952–53.

Preliminary diagram.

Invariant Equilibria

Mole % ZrF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
9	640	Eutectic	$L \rightleftharpoons \text{CsF} + 3\text{CsF} \cdot \text{ZrF}_4$
25	775	Congruent melting point	$L \rightleftharpoons 3\text{CsF} \cdot \text{ZrF}_4$
35	520	Peritectic	$L + 3\text{CsF} \cdot \text{ZrF}_4 \rightleftharpoons 2\text{CsF} \cdot \text{ZrF}_4$
40	420	Eutectic	$L \rightleftharpoons 2\text{CsF} \cdot \text{ZrF}_4 + \alpha\text{-CsF} \cdot \text{ZrF}_4$
50	515	Congruent melting point	$L \rightleftharpoons \alpha\text{-CsF} \cdot \text{ZrF}_4$
—	320	Inversion	$\alpha\text{-CsF} \cdot \text{ZrF}_4 \rightleftharpoons \beta\text{-CsF} \cdot \text{ZrF}_4$
59	470	Eutectic	$L \rightleftharpoons \alpha\text{-CsF} \cdot \text{ZrF}_4 + \text{ZrF}_4$

Fig. 3.36. The System CsF-ZrF₄.

3.37. The System LiF-NaF-ZrF₄

F. F. Blankenship, H. A. Friedman, H. Insley, R. E. Thoma, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953-59.

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Temperature (°C)	Type of Invariant	Solids Present at Invariant Temperature
LiF	NaF	ZrF ₄			
37	52	11	604	Eutectic	NaF, LiF, and 3NaF·ZrF ₄
55	22	23	590	Eutectic	α-3LiF·ZrF ₄ , 3NaF·ZrF ₄ ss, and LiF
32	36	32	450	Peritectic	β-3LiF·ZrF ₄ , 3NaF·ZrF ₄ ss, and 2LiF·ZrF ₄
31	35	34	448	Peritectic	3NaF·ZrF ₄ , α-5NaF·2ZrF ₄ ss, and 2LiF·ZrF ₄
31	34.5	34.5	445	Peritectic	2NaF·ZrF ₄ , α-5NaF·2ZrF ₄ ss, and 2LiF·ZrF ₄
26	37	37	425	Eutectic	2NaF·ZrF ₄ , 7NaF·6ZrF ₄ , and 2LiF·ZrF ₄
29	25.5	45.5	449	Eutectic	7NaF·6ZrF ₄ , 3NaF·4ZrF ₄ ss, and 2LiF·ZrF ₄
29	24.5	46.5	453	Peritectic	3NaF·4ZrF ₄ ss, 3LiF·4ZrF ₄ ss, and 2LiF·ZrF ₄
28	24	48	475	Peritectic	3NaF·4ZrF ₄ ss, 3LiF·4ZrF ₄ ss, and ZrF ₄

Fig. 3.37. The System LiF-NaF-ZrF_4 .

3.38. The System NaF-KF-ZrF₄

R. E. Thoma, C. J. Barton, H. Insley, H. A. Friedman, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-55.

Preliminary diagram.

Invariant Equilibria*

Composition of Liquid (mole %)**			Invariant Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
NaF	KF	ZrF ₄			
34	58	8	695	Eutectic	NaF, KF, and 3KF·ZrF ₄
45	38	17	720	Eutectic	NaF, 3KF·ZrF ₄ , and 3NaF·3KF·2ZrF ₄
61	19	20	710	Eutectic	NaF, 3NaF·ZrF ₄ , and 3NaF·3KF·2ZrF ₄
52	17	31		Peritectic	3NaF·ZrF ₄ , 3NaF·3KF·2ZrF ₄ , and NaF·KF·ZrF ₄
48	18	34		Peritectic	3NaF·ZrF ₄ , NaF·KF·ZrF ₄ , and 5NaF·2ZrF ₄
33	32	35		Peritectic	3NaF·3KF·2ZrF ₄ , 3KF·ZrF ₄ , and NaF·KF·ZrF ₄
12	51	37		Peritectic	3KF·ZrF ₄ , 2KF·ZrF ₄ , and NaF·KF·ZrF ₄
40	21	39		Peritectic	5NaF·2ZrF ₄ , 2NaF·ZrF ₄ , and NaF·KF·ZrF ₄
39	21	40		Peritectic	2NaF·ZrF ₄ , 7NaF·6ZrF ₄ , and NaF·KF·ZrF ₄
11	49	40		Peritectic	2KF·ZrF ₄ , 3KF·2ZrF ₄ , and NaF·KF·ZrF ₄
10	48	42	385	Eutectic	3KF·2ZrF ₄ , KF·ZrF ₄ , and NaF·KF·ZrF ₄
15	40	45		Eutectic	KF·ZrF ₄ , NaF·KF·ZrF ₄ , and 2NaF·3KF·5ZrF ₄
22	33	45		Peritectic	NaF·KF·ZrF ₄ , 7NaF·6ZrF ₄ , and 2NaF·3KF·5ZrF ₄
25	25	50		Eutectic	7NaF·6ZrF ₄ , 3NaF·4ZrF ₄ , and 2NaF·3KF·5ZrF ₄
21	29	51		Eutectic	3NaF·4ZrF ₄ , ZrF ₄ , and 2NaF·3KF·5ZrF ₄
20	30	50	432	Congruent melting point	2NaF·3KF·5ZrF ₄

*Three solid phases have been observed routinely in the composition region 30-50 mole % ZrF₄ which have not been identified as to composition. Whether these exhibit primary phases at the liquidus surface has not yet been determined.

**Compositions of invariant points shown by the intersection of dotted boundary curves are approximate.

Fig. 3.38. The System NaF-KF-ZrF₄. The isotherms for the temperatures 600, 650, and 700°C between the 30 and 40 mole % ZrF₄ compositions have been omitted in order not to obscure the phase relationships in this polythermal projection.

3.39. The System NaF-RbF-ZrF₄

R. E. Thoma, H. Insley, H. A. Friedman, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-56.

Preliminary diagram.

Invariant Equilibria					
Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
NaF	RbF	ZrF ₄			
23	72	5	643	Eutectic	NaF, RbF, and 3RbF·ZrF ₄
50	27	23	720	Eutectic	NaF, 3RbF·ZrF ₄ , and 3NaF·ZrF ₄
37	32	31	605	Eutectic	3NaF·ZrF ₄ , 3RbF·ZrF ₄ , and NaF·RbF·ZrF ₄
39	26	35	545	Peritectic	3NaF·ZrF ₄ , 5NaF·2ZrF ₄ , and NaF·RbF·ZrF ₄
36.3	24.2	39.5	427	Peritectic	5NaF·2ZrF ₄ , 2NaF·ZrF ₄ , and NaF·RbF·ZrF ₄
34.5	24	41.5	424	Peritectic	2NaF·ZrF ₄ , NaF·RbF·ZrF ₄ , and 3NaF·3RbF·4ZrF ₄
34	23.5	42.5	422	Peritectic	2NaF·ZrF ₄ , 7NaF·6ZrF ₄ , and 3NaF·3RbF·4ZrF ₄
33	23.5	43.5	420	Eutectic	7NaF·6ZrF ₄ , 3NaF·3RbF·4ZrF ₄ , and NaF·RbF·2ZrF ₄
28.5	21.5	50	443	Eutectic	3NaF·4ZrF ₄ , 7NaF·6ZrF ₄ , and NaF·RbF·2ZrF ₄
28	21.5	50.5	446	Peritectic	3NaF·4ZrF ₄ , ZrF ₄ , and NaF·RbF·2ZrF ₄
23.5	39.5	37	470	Peritectic	NaF·RbF·ZrF ₄ , 2RbF·ZrF ₄ , and 3RbF·ZrF ₄
21	40	39	438	Peritectic	NaF·RbF·ZrF ₄ , 2RbF·ZrF ₄ , and 3NaF·3RbF·4ZrF ₄
8	50	42	400	Eutectic	3NaF·3RbF·4ZrF ₄ , 2RbF·ZrF ₄ , and 5RbF·4ZrF ₄
8.5	47	44.5	395	Eutectic	3NaF·3RbF·4ZrF ₄ , 5RbF·ZrF ₄ , and NaF·RbF·2ZrF ₄
6.2	45.8	48	380	Eutectic	NaF·RbF·2ZrF ₄ , 5RbF·4ZrF ₄ , and RbF·ZrF ₄
5	42	53	398	Eutectic	NaF·RbF·2ZrF ₄ , RbF·ZrF ₄ , and RbF·2ZrF ₄
6.5	39	54.5	423	Peritectic	ZrF ₄ , NaF·RbF·2ZrF ₄ , and RbF·2ZrF ₄
33.3	33.3	33.3	642	Congruent melting point	NaF·RbF·ZrF ₄
25	25	50	462	Congruent melting point	NaF·RbF·2ZrF ₄

Minimum Temperatures on Alkemade Lines

Composition of Liquid (mole %)			Temperature (°C)	Alkemade Line
NaF	RbF	ZrF ₄		
24.75	24.75	51.5	455	NaF·RbF·2ZrF ₄ –ZrF ₄
6	44	50	402	NaF·RbF·2ZrF ₄ –RbF·ZrF ₄
7.5	46.5	46	405	NaF·RbF·2ZrF ₄ –5RbF·4ZrF ₄
8.5	48.5	43	405	3NaF·RbF·4ZrF ₄ –5RbF·4ZrF ₄
28	28	44	435	3NaF·RbF·4ZrF ₄ –NaF·RbF·2ZrF ₄
22	40	38	442	3NaF·3RbF·4ZrF ₄ –2RbF·ZrF ₄
29	41.5	49.5	450	NaF·RbF·2ZrF ₄ –7NaF·6ZrF ₄
37	24.5	38.5	610	NaF·RbF·ZrF ₄ –3NaF·ZrF ₄
47	28	25	732	3NaF·ZrF ₄ –3RbF·ZrF ₄
36	48	16	777	NaF–3RbF·ZrF ₄
30	36.7	33.3	598	NaF·RbF·ZrF ₄ –3RbF·ZrF ₄

Fig. 3.39. The System NaF–RbF–ZrF₄.

3.40. The System $\text{NaF}-\text{CrF}_2-\text{ZrF}_4$: The Section $\text{NaF}\cdot\text{CrF}_2 - \text{ZrF}_4$

R. E. Thoma, H. A. Friedman, B. S. Landau, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956-57.

Preliminary diagram. No invariant equilibria occur for compositions lying on the section $\text{NaF}\cdot\text{CrF}_2-\text{ZrF}_4$.

Fig. 3.40. The Section $\text{NaF}\cdot\text{CrF}_2-\text{ZrF}_4$.

3.41. The System NaF-CeF₃-ZrF₄

W. T. Ward, R. A. Strehlow, W. R. Grimes, and G. M. Watson, "Solubility Relationships of Rare Earth Fluorides and Yttrium Fluoride in Various Molten NaF-ZrF₄ and NaF-ZrF₄-UF₄ Solvents," *J. Chem. Eng. Data* (in press).

UNCLASSIFIED
ORNL-LR-DWG 29164R

Fig. 3.41a. The System NaF-CeF₃-ZrF₄.

UNCLASSIFIED
ORNL-LR-DWG 29165R

PROBABLE NaF·ZrF₄ PRIMARY PHASE FIELDS

A ZrF ₄	D 2NaF·ZrF ₄
B 3NaF·4ZrF ₄	E 5NaF·2ZrF ₄
C 7NaF·6ZrF ₄	F 3NaF·ZrF ₄

Fig. 3.41b. The System NaF-CeF₃-ZrF₄ in the Region 30-70 Mole % NaF, 0-20 Mole % CeF₃, 30-70 Mole % ZrF₄.

3.42. The System $\text{LiF}-\text{CeF}_3$

C. J. Barton, L. M. Bratcher, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956-57.

Preliminary diagram. The system $\text{LiF}-\text{CeF}_3$ contains a single eutectic at 81 LiF -19 CeF_3 (mole %), m.p. $755 \pm 5^\circ\text{C}$.

Fig. 3.42. The System $\text{LiF}-\text{CeF}_3$.

3.43. The System NaF–HfF₄

R. E. Thoma, C. F. Weaver, T. N. McVay, H. A. Friedman, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956–58.

Preliminary diagram.

Invariant Equilibria

Mole % HfF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
18.5	695	Eutectic	$L \rightleftharpoons \text{NaF} + 3\text{NaF} \cdot \text{HfF}_4$
25	863	Congruent melting point	$L \rightleftharpoons 3\text{NaF} \cdot \text{HfF}_4^*$
34	606	Peritectic	$L + 3\text{NaF} \cdot \text{HfF}_4 \rightleftharpoons \alpha\text{-}5\text{NaF} \cdot 2\text{HfF}_4$
–	535	Inversion	$\alpha\text{-}5\text{NaF} \cdot 2\text{HfF}_4 \rightleftharpoons \beta\text{-}5\text{NaF} \cdot 2\text{HfF}_4$
35	586	Peritectic	$L + \alpha\text{-}5\text{NaF} \cdot 2\text{HfF}_4 \rightleftharpoons \beta\text{-}2\text{NaF} \cdot \text{HfF}_4$
–	Below 350	Inversion	$\beta\text{-}2\text{NaF} \cdot \text{HfF}_4 \rightleftharpoons \gamma\text{-}2\text{NaF} \cdot \text{HfF}_4$
–	Below 350	Inversion	$\gamma\text{-}2\text{NaF} \cdot \text{HfF}_4 \rightleftharpoons \delta\text{-}2\text{NaF} \cdot \text{HfF}_4$
43	510	Eutectic	$L \rightleftharpoons \beta\text{-}2\text{NaF} \cdot \text{HfF}_4 + 7\text{NaF} \cdot 6\text{HfF}_4$
45	531	Peritectic	$L + \text{NaF} \cdot \text{HfF}_4 \rightleftharpoons 7\text{NaF} \cdot 6\text{HfF}_4$
50	557	Congruent melting point	$L \rightleftharpoons \text{NaF} \cdot \text{HfF}_4$
53	535	Eutectic	$L \rightleftharpoons \text{NaF} \cdot \text{HfF}_4 + \text{HfF}_4$

*A determination of the crystal structure of 3NaF·HfF₄ has been reported by L. A. Harris, "The Crystal Structures of Na₃ZrF₇ and Na₃HfF₇," *Acta Cryst.* 12, 172 (1959).

Fig. 3.43. The System NaF-HfF₄.

3.44. The System LiF–ThF₄

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, "Phase Equilibria in the Fused Salt Systems LiF–ThF₄ and NaF–ThF₄," *J. Phys. Chem.* **63**, 1266–74 (1959).

Invariant Equilibria

Mole % ThF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
23	568	Eutectic	$L \rightleftharpoons \text{LiF} + 3\text{LiF} \cdot \text{ThF}_4$
25	573	Congruent melting point	$L \rightleftharpoons 3\text{LiF} \cdot \text{ThF}_4$
29	565	Eutectic	$L \rightleftharpoons 3\text{LiF} \cdot \text{ThF}_4 + 7\text{LiF} \cdot 6\text{ThF}_4$
30.5	597	Peritectic	$\text{LiF} \cdot 2\text{ThF}_4 + L \rightleftharpoons 7\text{LiF} \cdot 6\text{ThF}_4$
42	762	Peritectic	$\text{LiF} \cdot 4\text{ThF}_4 + L \rightleftharpoons \text{LiF} \cdot 2\text{ThF}_4$
62	897	Peritectic	$\text{ThF}_4 + L \rightleftharpoons \text{LiF} \cdot 4\text{ThF}_4$

Fig. 3.44. The System LiF–ThF₄.

3.45. The System NaF–ThF₄

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, "Phase Equilibria in the Fused Salt Systems LiF–ThF₄ and NaF–ThF₄," *J. Phys. Chem.* **63**, 1266–74 (1959).

Invariant Equilibria

Mole % ThF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
21.5	645	Peritectic	$\text{NaF} + L \rightleftharpoons \alpha\text{-4NaF}\cdot\text{ThF}_4$
22.5	618	Eutectic	$L \rightleftharpoons \alpha\text{-4NaF}\cdot\text{ThF}_4 + 2\text{NaF}\cdot\text{ThF}_4$
–	604	Inversion	$\alpha\text{-4NaF}\cdot\text{ThF}_4 \rightleftharpoons \beta\text{-4NaF}\cdot\text{ThF}_4$
–	558	Decomposition	$\beta\text{-4NaF}\cdot\text{ThF}_4 \rightleftharpoons \text{NaF} + 2\text{NaF}\cdot\text{ThF}_4$
33.3	705	Congruent melting point	$L \rightleftharpoons 2\text{NaF}\cdot\text{ThF}_4$
37	690	Eutectic	$L \rightleftharpoons 2\text{NaF}\cdot\text{ThF}_4 + 3\text{NaF}\cdot 2\text{ThF}_4$
40	712	Congruent melting point	$L \rightleftharpoons 3\text{NaF}\cdot 2\text{ThF}_4$
41	705	Eutectic	$L \rightleftharpoons 3\text{NaF}\cdot 2\text{ThF}_4 + \alpha\text{-NaF}\cdot\text{ThF}_4$
–	683	Decomposition	$3\text{NaF}\cdot 2\text{ThF}_4 \rightleftharpoons 2\text{NaF}\cdot\text{ThF}_4 + \alpha\text{-NaF}\cdot\text{ThF}_4$
45.5	730	Peritectic	$\text{NaF}\cdot 2\text{ThF}_4 + L \rightleftharpoons \alpha\text{-NaF}\cdot\text{ThF}_4$
58	831	Peritectic	$\text{ThF}_4 + L \rightleftharpoons \text{NaF}\cdot 2\text{ThF}_4$

Fig. 3.45. The System NaF–ThF₄.

3.46. The System KF-ThF₄

W. J. Asker, E. R. Segnit, and A. W. Wylie, "The Potassium Thorium Fluorides," *J. Chem. Soc.* **1952**, 4470-79.

This system has also been reported by A. G. Bergman and E. P. Dergunov, *Doklady Akad. Nauk S.S.S.R.* **53**, 753 (1941) and by V. S. Emelyanov and A. J. Evstyukhin, "An Investigation of Fused-Salt Systems Based on Thorium Fluoride - II. NaF-KF-ThF₄, NaF-ThF₄, and KF-ThF₄," *J. Nuclear Energy* **5**, 108-14 (1957).

Invariant Equilibria

Mole % ThF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
14	694	Eutectic	$L \rightleftharpoons \text{KF} + \beta\text{-5KF}\cdot\text{ThF}_4$
-	635	Inversion	$\alpha\text{-5KF}\cdot\text{ThF}_4 \rightleftharpoons \beta\text{-5KF}\cdot\text{ThF}_4$
-	712	Peritectic	$L + 3\text{KF}\cdot\text{ThF}_4 \rightleftharpoons \alpha\text{-5KF}\cdot\text{ThF}_4$
25	865	Congruent melting point	$L \rightleftharpoons 3\text{KF}\cdot\text{ThF}_4$
-	570	Decomposition	$3\text{KF}\cdot\text{ThF}_4 \rightleftharpoons \beta\text{-5KF}\cdot\text{ThF}_4 + \beta\text{-2KF}\cdot\text{ThF}_4$
31	691	Eutectic	$L \rightleftharpoons 3\text{KF}\cdot\text{ThF}_4 + \alpha\text{-2KF}\cdot\text{ThF}_4$
-	747	Peritectic	$L + \text{KF}\cdot\text{ThF}_4 \rightleftharpoons \alpha\text{-2KF}\cdot\text{ThF}_4$
-	645	Inversion	$\alpha\text{-2KF}\cdot\text{ThF}_4 \rightleftharpoons \beta\text{-2KF}\cdot\text{ThF}_4$
50*	905	Congruent melting point	$L \rightleftharpoons \text{KF}\cdot\text{ThF}_4$
56	875	Eutectic	$L \rightleftharpoons \text{KF}\cdot\text{ThF}_4 + \text{KF}\cdot 2\text{ThF}_4$
66	930	Peritectic	$L + \text{KF}\cdot 3\text{ThF}_4 \rightleftharpoons \text{KF}\cdot 2\text{ThF}_4$
75	990	Congruent melting point	$L \rightleftharpoons \text{KF}\cdot 3\text{ThF}_4$
78	980	Eutectic	$L \rightleftharpoons \text{KF}\cdot 3\text{ThF}_4 + \text{ThF}_{4ss}$

*It has been shown that the compound labeled by Asker *et al.* has the actual formula 7KF·6ThF₄, cf. R. E. Thoma, *Crystal Structures of Some Compounds of UF₄ and ThF₄ with Alkali Fluorides*, ORNL CF-58-12-40 (December 11, 1958).

Fig. 3.46. The System KF-ThF₄.

3.47. The System RbF–ThF₄

E. P. Dergunov and A. G. Bergman, "Complex Formation Between Alkali Metal Fluorides and Fluorides of Metals of the Fourth Group," *Doklady Akad. Nauk S.S.S.R.* **60**, 391–94 (1948).

Invariant Equilibria

Mole % ThF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
15	664	Eutectic	$L \rightleftharpoons \text{RbF} + 3\text{RbF} \cdot \text{ThF}_4$
25	974	Congruent melting point	$L \rightleftharpoons 3\text{RbF} \cdot \text{ThF}_4$
37	762	Eutectic	$L \rightleftharpoons 3\text{RbF} \cdot \text{ThF}_4 + \text{RbF} \cdot \text{ThF}_4$
50	852	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot \text{ThF}_4$
54	848	Eutectic	$L \rightleftharpoons \text{RbF} \cdot \text{ThF}_4 + \text{RbF} \cdot 3\text{ThF}_4$
75	1004	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot 3\text{ThF}_4$
80	1000	Eutectic	$L \rightleftharpoons \text{RbF} \cdot 3\text{ThF}_4 + \text{ThF}_4$

Fig. 3.47. The System RbF–ThF₄.

3.48. The System $\text{BeF}_2\text{--ThF}_4$

R. E. Thoma, H. Insley, H. A. Friedman, and C. F. Weaver, "Phase Equilibria in the Systems $\text{BeF}_2\text{--ThF}_4$ and $\text{LiF--BeF}_2\text{--ThF}_4$," paper to be presented at the 136th National Meeting of the American Chemical Society, Atlantic City, N. J., Sept. 13–18, 1959.

The system $\text{BeF}_2\text{--ThF}_4$ contains a single eutectic at 98.5 $\text{BeF}_2\text{--}1.5 \text{ ThF}_4$ (mole %), m.p. $526 \pm 3^\circ\text{C}$.

Fig. 3.48. The System $\text{BeF}_2\text{--ThF}_4$.

3.49. The System $\text{BeF}_2\text{--UF}_4$

T. B. Rhinehammer, P. A. Tucker, and E. F. Joy, *Phase Equilibria in the System $\text{BeF}_2\text{--UF}_4$* , MLM-1082 (to be published).

Preliminary diagram. The system $\text{BeF}_2\text{--UF}_4$ contains a single eutectic at 99.5 $\text{BeF}_2\text{--}0.5 \text{UF}_4$ (mole %), m.p. $535 \pm 2^\circ\text{C}$.

Fig. 3.49. The System $\text{BeF}_2\text{--UF}_4$.

3.50. The System $\text{MgF}_2\text{--ThF}_4$

J. O. Blomeke, *An Investigation of the $\text{ThF}_4\text{--Fused Salt Solutions for Homogeneous Reactors}$* , ORNL-1030 (June 19, 1951) (declassified with deletions).

Invariant Equilibria			
Mole % ThF_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
25	915	Eutectic	$L \rightleftharpoons \text{ThF}_4 + \text{MgF}_2 \cdot 2\text{ThF}_4$
33.3	937	Congruent melting point	$L \rightleftharpoons \text{MgF}_2 \cdot 2\text{ThF}_4$
40	925	Eutectic	$L \rightleftharpoons \text{MgF}_2 \cdot 2\text{ThF}_4 + \text{MgF}_2$

Fig. 3.50. The System $\text{MgF}_2\text{--ThF}_4$.

3.51. The System $\text{LiF}-\text{BeF}_2-\text{ThF}_4$

R. E. Thoma, H. Insley, H. A. Friedman, and C. F. Weaver, "Phase Equilibria in the Systems $\text{BeF}_2-\text{ThF}_4$ and $\text{LiF}-\text{BeF}_2-\text{ThF}_4$," paper to be presented at the 136th National Meeting of the American Chemical Society, Atlantic City, N. J., Sept. 13-18, 1959.

Invariant Equilibria

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Invariant	Solids Present at Invariant Point
LiF	BeF_2	ThF_4			
15	83	2	497 ± 4	Peritectic	ThF_4 , $\text{LiF} \cdot 4\text{ThF}_4$, and BeF_2
33.5	64	2.5	455 ± 4	Peritectic	$\text{LiF} \cdot 4\text{ThF}_4$, $\text{LiF} \cdot 2\text{ThF}_4$, and BeF_2
47	51.5	1.5	356 ± 6	Eutectic	$2\text{LiF} \cdot \text{BeF}_2$, $\text{LiF} \cdot 2\text{ThF}_4$, and BeF_2
60.5	36.5	3	433 ± 5	Peritectic	$\text{LiF} \cdot 2\text{ThF}_4$, $3\text{LiF} \cdot \text{ThF}_4$ ss, and $2\text{LiF} \cdot \text{BeF}_2$
65.5	30.5	4	444 ± 4	Peritectic	LiF , $2\text{LiF} \cdot \text{BeF}_2$, and $3\text{LiF} \cdot \text{ThF}_4$ ss
63	30.5	6.5	448 ± 5	Peritectic	$3\text{LiF} \cdot \text{ThF}_4$ ss, $7\text{LiF} \cdot 6\text{ThF}_4$, and $\text{LiF} \cdot 2\text{ThF}_4$

Fig. 3.51a. The System $\text{LiF}-\text{BeF}_2-\text{ThF}_4$.

Limits of Single-Phase $3\text{LiF}\cdot\text{ThF}_4$ Solid Solution

Composition (mole %)		
LiF	BeF_2	ThF_4
75	0	25
58	16	26
59	20	21

A three-dimensional model of the system $\text{LiF}-\text{BeF}_2-\text{ThF}_4$ is shown in Fig. 3.51b.

Fig. 3.51b. Model of the System $\text{LiF}-\text{BeF}_2-\text{ThF}_4$.

3.52. The System NaF–ZrF₄–ThF₄

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956–57.

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
NaF	ZrF ₄	ThF ₄			
77	3	20	645	Peritectic	NaF, α -4NaF·ThF ₄ , and 3NaF·ZrF ₄
75.5	2.5	22	618	Eutectic	α -4NaF·ThF ₄ , 2NaF·ThF ₄ , and 3NaF·ZrF ₄
63	6	31*	683	Decomposition point of 3NaF·2ThF ₄ in ternary system	3NaF·2ThF ₄ , 2NaF·ThF ₄ , and NaF·ThF ₄
65.5	21.5	13	622	Peritectic	2NaF·ThF ₄ , 3NaF·ZrF ₄ , and NaF·ThF ₄
65.5	24.5	10	615	Peritectic	3NaF·ZrF ₄ , NaF·ThF ₄ , and α -5NaF·2ZrF ₄
64.5	25.5	10	593	Peritectic	α -5NaF·2ZrF ₄ , NaF·ThF ₄ , and NaF·2ThF ₄
58	38	4	538	Peritectic	α -5NaF·2ZrF ₄ , α -2NaF·ZrF ₄ , and NaF·2ThF ₄
58	40	2	495	Eutectic	β -2NaF·ZrF ₄ , NaF·2ThF ₄ , and 7NaF·6ZrF ₄ –3NaF·2ZrF ₄ ^{ss}
53	42	4	510	Peritectic	NaF·2ThF ₄ , Th(Zr)F ₄ ^{ss} , and 7NaF·6ZrF ₄ –3NaF·2ThF ₄ ^{ss}
49.5	48	2.5	505	Eutectic	7NaF·6ZrF ₄ , 3NaF·4ZrF ₄ , and Zr(Th)F ₄ ^{ss}

*Approximate composition.

Note: The boundary curve maximum on the Alkemade line 7NaF·6ZrF₄–ThF₄ is unlikely to fall exactly on this line, because pure ThF₄ does not occur along this line. Since ThF₄ forms solid solutions with both ZrF₄ and NaF·2ThF₄, it is not readily determined on which side of the Alkemade line this maximum occurs.

Fig. 3.52. The System NaF-ZrF₄-ThF₄.

3.53. The System LiF-UF₃

C. J. Barton, V. S. Coleman, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953-54.

Preliminary diagram. The system LiF-UF₃ contains a single eutectic at 73 LiF-27 UF₃ (mole %), m.p. 770°C.

Fig. 3.53. The System LiF-UF₃.

3.54. The System LiF-UF₄

C. J. Barton, H. A. Friedman, W. R. Grimes, H. Insley, R. E. Moore, and R. E. Thoma, "Phase Equilibria in the Alkali Fluoride-Uranium Tetrafluoride Fused Salt Systems: I. The Systems LiF-UF₄ and NaF-UF₄," *J. Am. Ceram. Soc.* **41**, 63-69 (1958).

Invariant Equilibria

Mole % UF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
—	470	Decomposition	$4\text{LiF} \cdot \text{UF}_4 \rightleftharpoons \text{LiF} + 7\text{LiF} \cdot 6\text{UF}_4$
26	500	Peritectic	$\text{LiF} + L \rightleftharpoons 4\text{LiF} \cdot \text{UF}_4$
27	490	Eutectic	$L \rightleftharpoons 4\text{LiF} \cdot \text{UF}_4 + 7\text{LiF} \cdot 6\text{UF}_4$
40	610	Peritectic	$L + \text{LiF} \cdot 4\text{UF}_4 \rightleftharpoons 7\text{LiF} \cdot 6\text{UF}_4$
57	775	Peritectic	$L + \text{UF}_4 \rightleftharpoons \text{LiF} \cdot 4\text{UF}_4$

Fig. 3.54. The System LiF-UF₄.

3.55. The System NaF-UF₃

C. J. Barton, V. S. Coleman, T. N. McVay, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953-54.

Preliminary diagram.

Invariant Equilibria			
Mole % UF ₃ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
27	715	Eutectic	$L \rightleftharpoons \text{NaF} + \alpha\text{-NaF}\cdot\text{UF}_3$
35	775	Peritectic	$L + \text{UF}_3 \rightleftharpoons \alpha\text{-NaF}\cdot\text{UF}_3$
—	595	Inversion	$\alpha\text{-NaF}\cdot\text{UF}_3 \rightleftharpoons \beta\text{-NaF}\cdot\text{UF}_3$

Fig. 3.55. The System NaF-UF₃.

3.56. The System NaF-UF₄

C. J. Barton, H. A. Friedman, W. R. Grimes, H. Insley, R. E. Moore, and R. E. Thoma, "Phase Equilibria in the Alkali Fluoride-Uranium Tetrafluoride Fused Salt Systems: 1. The Systems LiF-UF₄ and NaF-UF₄," *J. Am. Ceram. Soc.* **41**, 63-69 (1958).

Invariant Equilibria

Mole % UF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
21.5	618	Eutectic	$L \rightleftharpoons \alpha\text{-}3\text{NaF}\cdot\text{UF}_4 + \text{NaF}$
25	629	Congruent melting point	$L \rightleftharpoons \alpha\text{-}3\text{NaF}\cdot\text{UF}_4$
—	528	Inversion	$\alpha\text{-}3\text{NaF}\cdot\text{UF}_4 \rightleftharpoons \beta\text{-}3\text{NaF}\cdot\text{UF}_4$
—	497	Decomposition	$\beta\text{-}3\text{NaF}\cdot\text{UF}_4 \rightleftharpoons \text{NaF} + 2\text{NaF}\cdot\text{UF}_4$
28	623	Eutectic	$L \rightleftharpoons \alpha\text{-}3\text{NaF}\cdot\text{UF}_4 + 2\text{NaF}\cdot\text{UF}_4$
32.5	648	Peritectic	$L + 5\text{NaF}\cdot 3\text{UF}_4 \rightleftharpoons 2\text{NaF}\cdot\text{UF}_4$
37	673	Peritectic	$L + 7\text{NaF}\cdot 6\text{UF}_4 \rightleftharpoons 5\text{NaF}\cdot 3\text{UF}_4$
—	630	Decomposition	$5\text{NaF}\cdot 3\text{UF}_4 \rightleftharpoons 2\text{NaF}\cdot\text{UF}_4 + 7\text{NaF}\cdot 6\text{UF}_4$
46.2	718	Congruent melting point	$L \rightleftharpoons 7\text{NaF}\cdot 6\text{UF}_4$
56	680	Eutectic	$L \rightleftharpoons 7\text{NaF}\cdot 6\text{UF}_4 + \text{UF}_4$
—	660	Upper stability limit of NaF·2UF ₄	$7\text{NaF}\cdot 6\text{UF}_4 + \text{UF}_4 \rightleftharpoons \text{NaF}\cdot 2\text{UF}_4$

Fig. 3.56. The System NaF-UF₄.

3.57. The System $\text{KF}-\text{UF}_4$

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, "Phase Equilibria in the Alkali Fluoride-Uranium Tetrafluoride Fused Salt Systems: II. The Systems $\text{KF}-\text{UF}_4$ and $\text{RbF}-\text{UF}_4$," *J. Am. Ceram. Soc.* **41**, 538-44 (1958).

Invariant Equilibria

Mole % UF_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
15	735	Eutectic	$L \rightleftharpoons \text{KF} + 3\text{KF}\cdot\text{UF}_4$
25	957	Congruent melting point	$L \rightleftharpoons 3\text{KF}\cdot\text{UF}_4$
—	608	Decomposition	$2\text{KF}\cdot\text{UF}_4 \rightleftharpoons 3\text{KF}\cdot\text{UF}_4 + 7\text{KF}\cdot 6\text{UF}_4$
35	755	Peritectic	$3\text{KF}\cdot\text{UF}_4 + L \rightleftharpoons 2\text{KF}\cdot\text{UF}_4$
38.5	740	Eutectic	$L \rightleftharpoons 2\text{KF}\cdot\text{UF}_4 + 7\text{KF}\cdot 6\text{UF}_4$
46.2	789	Congruent melting point	$L \rightleftharpoons 7\text{KF}\cdot 6\text{UF}_4$
54.5	735	Eutectic	$L \rightleftharpoons 7\text{KF}\cdot 6\text{UF}_4 + \text{KF}\cdot 2\text{UF}_4$
65	765	Peritectic	$\text{UF}_4 + L \rightleftharpoons \text{KF}\cdot 2\text{UF}_4$

Fig. 3.57. The System $\text{KF}-\text{UF}_4$.

3.58. The System RbF-UF₄

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, "Phase Equilibria in the Alkali Fluoride-Uranium Tetrafluoride Fused Salt Systems: II. The Systems KF-UF₄ and RbF-UF₄," *J. Am. Ceram. Soc.* 41, 538-44 (1958).

Invariant Equilibria

Mole % UF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
10	710	Eutectic	$L \rightleftharpoons \text{RbF} + 3\text{RbF} \cdot \text{UF}_4$
25	995	Congruent melting point	$L \rightleftharpoons 3\text{RbF} \cdot \text{UF}_4$
38	818	Peritectic	$L + 3\text{RbF} \cdot \text{UF}_4 \rightleftharpoons 2\text{RbF} \cdot \text{UF}_4$
43.5	675	Eutectic	$L \rightleftharpoons 2\text{RbF} \cdot \text{UF}_4 + 7\text{RbF} \cdot 6\text{UF}_4$
44	693	Peritectic	$\text{RbF} \cdot \text{UF}_4 + L \rightleftharpoons 7\text{RbF} \cdot 6\text{UF}_4$
50	735	Congruent melting point	$L \rightleftharpoons \text{RbF} \cdot \text{UF}_4$
55	714	Eutectic	$L \rightleftharpoons \text{RbF} \cdot \text{UF}_4 + 2\text{RbF} \cdot 3\text{UF}_4$
56.5	722	Peritectic	$\text{RbF} \cdot 3\text{UF}_4 + L \rightleftharpoons 2\text{RbF} \cdot 3\text{UF}_4$
57	730	Peritectic	$\text{RbF} \cdot 6\text{UF}_4 + L \rightleftharpoons \text{RbF} \cdot 3\text{UF}_4$
70.5	832	Peritectic	$\text{UF}_4 + L \rightleftharpoons \text{RbF} \cdot 6\text{UF}_4$

Fig. 3.58. The System RbF-UF₄.

3.59. The System CsF-UF₄

C. J. Barton, L. M. Bratcher, J. P. Blakely, G. J. Nettle, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951-53.

Preliminary diagram.

Invariant Equilibria

Mole % UF ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
7.5	650	Eutectic	$L \rightleftharpoons \text{CsF} + 3\text{CsF} \cdot \text{UF}_4$
25	970	Congruent melting point	$L \rightleftharpoons 3\text{CsF} \cdot \text{UF}_4$
34	800	Peritectic	$L + 3\text{CsF} \cdot \text{UF}_4 \rightleftharpoons 2\text{CsF} \cdot \text{UF}_4$
41	695	Eutectic	$L \rightleftharpoons 2\text{CsF} \cdot \text{UF}_4 + \alpha\text{-CsF} \cdot \text{UF}_4$
50	735	Congruent melting point	$L \rightleftharpoons \alpha\text{-CsF} \cdot \text{UF}_4$
-	550	Inversion	$\alpha\text{-CsF} \cdot \text{UF}_4 \rightleftharpoons \beta\text{-CsF} \cdot \text{UF}_4$
53	725	Eutectic	$L \rightleftharpoons \alpha\text{-CsF} \cdot \text{UF}_4 + \text{UF}_4$

Fig. 3.59. The System CsF-UF₄.

3.60. The System $\text{ZrF}_4\text{--UF}_4$

C. J. Barton, W. R. Grimes, H. Insley, R. E. Moore, and R. E. Thoma, "Phase Equilibria in the Systems NaF--ZrF_4 , $\text{UF}_4\text{--ZrF}_4$, and $\text{NaF--ZrF}_4\text{--UF}_4$," *J. Phys. Chem.* **62**, 665–76 (1958).

The system $\text{ZrF}_4\text{--UF}_4$ forms a continuous series of solid solutions having a minimum melting temperature of 765°C at 77 $\text{ZrF}_4\text{--}23 \text{ UF}_4$ (mole %).

Fig. 3.60. The System $\text{ZrF}_4\text{--UF}_4$.

3.61. The System $\text{SnF}_2\text{-UF}_4$

B. J. Thamer and G. E. Meadows, *The Systems $\text{UF}_4\text{-SnF}_2$ and $\text{PuF}_3\text{-SnF}_2$* , LA-2286 (July 1959).

Invariant Equilibria

Mole % UF_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
0.5	212	Eutectic	$L \rightleftharpoons \text{SnF}_2 + 2\text{SnF}_2 \cdot \text{UF}_4$
4.5	340	Peritectic	$L + \text{SnF}_2 \cdot \text{UF}_4 \rightleftharpoons 2\text{SnF}_2 \cdot \text{UF}_4$
7.8	371	Peritectic	$L + \text{UF}_4 \rightleftharpoons \text{SnF}_2 \cdot \text{UF}_4$

Fig. 3.61. The System $\text{SnF}_2\text{-UF}_4$.

3.62. The System $\text{PbF}_2\text{--UF}_4$

C. J. Barton, L. M. Bratcher, J. P. Blakely, G. J. Nessle, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1950–51.

Preliminary diagram.

Invariant Equilibria			
Mole % UF_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
14.3	920	Congruent melting point	$L \rightleftharpoons 6\text{PbF}_2 \cdot \text{UF}_4$
35	835	Eutectic	$L \rightleftharpoons 6\text{PbF}_2 \cdot \text{UF}_4 + 3\text{PbF}_2 \cdot 2\text{UF}_4$
40	840	Congruent melting point	$L \rightleftharpoons 3\text{PbF}_2 \cdot 2\text{UF}_4$
62	762 ± 10	Eutectic	$L \rightleftharpoons 3\text{PbF}_2 \cdot 2\text{UF}_4 + \text{UF}_4$

The eutectic at quite high PbF_2 concentrations is not shown on this diagram. The authors examined thermal data but found no evidence of a eutectic thermal effect.

Fig. 3.62. The System $\text{PbF}_2\text{--UF}_4$.

3.63. The System $\text{ThF}_4\text{--UF}_4$

C. F. Weaver, R. E. Thoma, H. A. Friedman, and H. Insley, "Phase Equilibria in the Systems $\text{UF}_4\text{--ThF}_4$ and $\text{LiF--ThF}_4\text{--UF}_4$," paper presented at the 61st National Meeting of the American Ceramic Society, Chicago, Ill., May 17–21, 1959.

The system $\text{ThF}_4\text{--UF}_4$ forms a continuous series of solid solutions without a minimum.

Fig. 3.63. The System ThF₄-UF₄.

3.64. The System LiF–NaF–UF₄

R. E. Thoma, H. Insley, B. S. Landau, H. A. Friedman, and W. R. Grimes, "Phase Equilibria in the Alkali Fluoride–Uranium Tetrafluoride Fused Salt Systems: III. The System NaF–LiF–UF₄," *J. Am. Ceram. Soc.* **42**, 21–26 (1959).

Invariant Equilibria

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solid Phases Present at Invariant Point
NaF	LiF	UF ₄			
60	21	19	480	Eutectic	2NaF·UF ₄ , NaF, and LiF
65	13	22	497	Peritectic	β -3NaF·UF ₄ , 2NaF·UF ₄ , and NaF
7	65.5	27.5	470	Peritectic	7LiF·6UF _{4ss} ,* 4LiF·UF ₄ , and LiF
35	37	28	480	Eutectic	2NaF·UF ₄ , 7NaF·6UF _{4ss} , and LiF
57	13	30	630	Peritectic	2NaF·UF ₄ , 5NaF·3UF ₄ , and 7NaF·6UF _{4ss}
24.3	43.5	32.2	445	Eutectic	7NaF·6UF _{4ss} , LiF, and 7LiF·6UF _{4ss}
24.5	29	46.5	602	Eutectic	NaF·2UF ₄ , 7LiF·6UF _{4ss} , and 7NaF·6UF _{4ss}
24.3	28.7	47	605	Peritectic	NaF·2UF ₄ , LiF·4UF ₄ , and 7LiF·6UF _{4ss}
23.5	28	48.5	640	Peritectic	UF ₄ , NaF·2UF ₄ , and LiF·4UF ₄
37.5	10.5	52	660	Peritectic	UF ₄ , NaF·2UF ₄ , and 7NaF·6UF _{4ss}

*The compounds 7LiF·6UF₄ and 7NaF·6UF₄ as they occur in the ternary system are members of the solid solution 7LiF·6UF₄–7NaF·6UF₄.

Fig. 3.64a. The System LiF-NaF-UF₄.

Fig. 3.64b. The Join $7\text{NaF} \cdot 6\text{UF}_4$ - $7\text{LiF} \cdot 6\text{UF}_4$.

3.65. The System LiF-KF-UF_4

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1950-51.

Preliminary diagram. The system LiF-KF-UF_4 has not yet been defined at the Oak Ridge National Laboratory with sufficient precision to permit the temperatures and compositions of the invariant equilibria to be listed.

Fig. 3.65. The System LiF-KF-UF_4 .

3.66. The System LiF-RbF-UF₄

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1950-51.

Preliminary diagram. Phase relationships within the system LiF-RbF-UF₄ have not yet become so well defined at the Oak Ridge National Laboratory that the compositions and temperatures of the invariant points may be listed.

UNCLASSIFIED
ORNL-LR-DWG 28634R

Fig. 3.66. The System LiF-RbF-UF₄.

3.67. The System NaF-KF-UF₄

R. E. Thoma, C. J. Barton, J. P. Blakely, R. E. Moore, G. J. Nessel, H. Insley, and H. A. Friedman, unpublished work performed at the Oak Ridge National Laboratory, 1950-58.

Preliminary diagram. Phase relationships within the system NaF-KF-UF₄ have not yet become so well defined at the Oak Ridge National Laboratory that the compositions and temperatures of the invariant points may be listed.

3.68. The System NaF-RbF-UF₄

R. E. Thoma, H. Insley, H. A. Friedman, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1955-56.

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solid Phases Present at Invariant Point
NaF	RbF	UF ₄			
18	73	9	670	Eutectic	RbF, NaF, and 3RbF·UF ₄
46	33	21	470	Eutectic	NaF, NaF·RbF·UF ₄ , and 3RbF·UF ₄
44	33	23	500	Peritectic	NaF·RbF·UF ₄ , 2RbF·UF ₄ , and 3RbF·UF ₄
45	30	25	485	Peritectic	NaF, 2NaF·UF ₄ , and NaF·RbF·UF ₄
52	23	25	500	Decomposition	3NaF·UF ₄ , NaF, and 2NaF·UF ₄
41	26	33	555	Peritectic	2RbF·UF ₄ , 2NaF·UF ₄ , and NaF·RbF·UF ₄
57	8	35	630	Decomposition	2NaF·UF ₄ , 5NaF·3UF ₄ , and 7NaF·6UF ₄
33	30	37	535	Eutectic	2RbF·UF ₄ , 2NaF·UF ₄ , and 7RbF·6UF ₄
32	29	39	540	Peritectic	7RbF·6UF ₄ , 2NaF·UF ₄ , and 7NaF·6UF ₄
26	27	47	620	Eutectic	7RbF·6UF ₄ , 7NaF·6UF ₄ , and RbF·UF ₄
25	25	50	630	Eutectic	7NaF·6UF ₄ , RbF·UF ₄ , and 2RbF·3UF ₄
27	22	51	633	Peritectic	7NaF·6UF ₄ , 2RbF·3UF ₄ , and RbF·3UF ₄
33	14	53	655	Peritectic	7NaF·6UF ₄ , RbF·3UF ₄ , and RbF·6UF ₄
42	3	55	678	Peritectic	7NaF·6UF ₄ , RbF·6UF ₄ , and UF ₄

Fig. 3.68a. The System NaF-RbF-UF₄.

Fig. 3.68b. The Section 2NaF·UF₄-2RbF·UF₄.

Fig. 3.68c. The Section NaF-NaF·RbF·UF₄.

Fig. 3.68d. The Section $3\text{RbF}\cdot\text{UF}_4\text{--NaF}\cdot\text{RbF}\cdot\text{UF}_4$.

3.69. The System $\text{LiF}-\text{BeF}_2-\text{UF}_4$

L. V. Jones, D. E. Etter, C. R. Hudgens, A. A. Huffman, T. B. Rhinehammer, N. E. Rogers, P. A. Tucker, and L. J. Wittenberg, *Phase Equilibria in the $\text{LiF}-\text{BeF}_2-\text{UF}_4$ Ternary Fused Salt System*, MLM-1080 (Aug. 24, 1959).

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solid Phases Present at Invariant Temperature
LiF	BeF ₂	UF ₄			
72	6	22	480	Peritectic (decomposition of 4LiF·UF ₄ in the ternary system)	4LiF·UF ₄ , LiF, and 7LiF·6UF ₄
69	23	8	426	Eutectic	LiF, 2LiF·BeF ₂ , and 7LiF·6UF ₄
48	51.5	0.5	350	Eutectic	7LiF·6UF ₄ , 2LiF·BeF ₂ , and BeF ₂
45.5	54	0.5	381	Peritectic	LiF·4UF ₄ , 7LiF·6UF ₄ , and BeF ₂
29.5	70	0.5	483	Peritectic	UF ₄ , LiF·4UF ₄ , and BeF ₂

The minimum temperature in the quasi-binary system 2LiF·BeF₂–7LiF·6UF₄ occurs at 65 LiF–29 BeF₂–6 UF₄ (mole %) at 438°C.

A three-dimensional model of the system LiF–BeF₂–UF₄ is shown in Fig. 3.69b.

Fig. 3.69b. Model of the System LiF–BeF₂–UF₄.

3.70. The System NaF-BeF₂-UF₄

J. F. Eichelberger, C. R. Hudgens, L. V. Jones, G. Pish, T. B. Rhinehammer, P. A. Tucker, and L. J. Wittenberg, unpublished work performed at the Mound Laboratory, 1956-57.

Preliminary diagram.

Invariant Equilibria

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solid Phases Present at Invariant Temperature
NaF	BeF ₂	UF ₄			
74	12	14	500	Peritectic (decomposition of 3NaF·UF ₄ in ternary system)	3NaF·UF ₄ , NaF, and 2NaF·UF ₄
72.5	17	19.5	486	Eutectic	NaF, 2NaF·UF ₄ , and 2NaF·BeF ₂
64.5	9	26.5	630	Peritectic (decomposition of 5NaF·3UF ₄ in the ternary system)	5NaF·3UF ₄ , 2NaF·UF ₄ , and 7NaF·6UF ₄
57	42	1	378	Peritectic	2NaF·BeF ₂ , 2NaF·UF ₄ , and 7NaF·6UF ₄
56	43.5	0.5	339	Eutectic	2NaF·BeF ₂ , NaF·BeF ₂ , and 7NaF·6UF ₄
43.5	55.5	1	357	Eutectic	7NaF·6UF ₄ , BeF ₂ , and NaF·BeF ₂
41	58	1	375	Peritectic	7NaF·6UF ₄ , NaF·2UF ₄ , and BeF ₂
26	63	1	409	Peritectic	NaF·2UF ₄ , NaF·4UF ₄ , and BeF ₂
44	18	38	665	Peritectic	7NaF·6UF ₄ , UF ₄ , and NaF·2UF ₄
40	47	13	548	Peritectic	UF ₄ , NaF·2UF ₄ , and NaF·4UF ₄
27	72	1	498	Peritectic	UF ₄ , BeF ₂ , and NaF·4UF ₄

The minimum temperature in the quasi-binary system 2NaF·BeF₂-2NaF·UF₄ occurs at 66.7 LiF-25 BeF₂-8.3 UF₄ (mole %) at 528°C. The minimum temperature in the quasi-binary system NaF·BeF₂-7NaF·6UF₄ occurs at 50.5 NaF-48.5 BeF₂-1.0 UF₄ (mole %) at 367°C.

A three-dimensional model of the system NaF-BeF₂-UF₄ has been constructed by the authors and is shown in Fig. 3.70b.

Fig. 3.70a. The System $NaF-BeF_2-UF_4$.

UNCLASSIFIED
PHOTO 47483

Fig. 3.70*b*. Model of the System $\text{NaF}-\text{BeF}_2-\text{UF}_4$.

3.71. The System $\text{NaF}-\text{PbF}_2-\text{UF}_4$

C. J. Barton, J. P. Blakely, G. J. Nessel, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1950-51.

Preliminary diagram. No study has been made at the Oak Ridge National Laboratory of the phase relationships within the system $\text{NaF}-\text{PbF}_2-\text{UF}_4$.

Fig. 3.71. The System $\text{NaF}-\text{PbF}_2-\text{UF}_4$.

3.72. The system $\text{KF-PbF}_2\text{-UF}_4$

C. J. Barton, J. P. Blakely, G. J. Nettle, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1950-51.

Preliminary diagram. No study has been made at the Oak Ridge National Laboratory of the phase relationships within the system $\text{KF-PbF}_2\text{-UF}_4$.

Fig. 3.72. The System $\text{KF}-\text{PbF}_2-\text{UF}_4$.

3.73. The System NaF-ZrF₄-UF₄

C. J. Barton, W. R. Grimes, H. Insley, R. E. Moore, and R. E. Thoma, "Phase Equilibria in the Systems NaF-ZrF₄, UF₄-ZrF₄, and NaF-ZrF₄-UF₄," *J. Phys. Chem.* **62**, 665-76 (1958).

UNCLASSIFIED
ORNL-LR-DWG 19886R

Fig. 3.73a. The System NaF-ZrF₄-UF₄

Invariant Equilibria and Singular Points

Composition of Liquid (mole %)			Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
NaF	ZrF ₄	UF ₄			
69.5	4.0	26.5	646	Maximum temperature of boundary curve	3NaF·(U,Zr)F ₄ and 2NaF·UF ₄
68.5	5.5	26.0	640	Peritectic	3NaF·(U,Zr)F ₄ , 2NaF·UF ₄ , and 5NaF·3UF ₄ ^{ss}
65.5	12.0	22.5	613	Peritectic or decomposition	3NaF·(U,Zr)F ₄ , 5NaF·3UF ₄ ^{ss} , and 7NaF·6(U,Zr)F ₄
64.0	27.0	9.0	592	Peritectic	3NaF·(U,Zr)F ₄ , 5NaF·2ZrF ₄ ^{ss} , and 7NaF·6(U,Zr)F ₄
61.5	34.5	4.0	540	Peritectic	5NaF·2ZrF ₄ ^{ss} , 2NaF·ZrF ₄ , and 7NaF·6(U,Zr)F ₄
50.5	47	2.5	513	Peritectic	7NaF·6(U,Zr)F ₄ , (U,Zr)F ₄ , and 3NaF·4ZrF ₄

Fig. 3.73b. The Subsystem 3NaF·UF₄-3NaF·ZrF₄.

Fig. 3.73c. The Subsystem 7NaF·6ZrF₄-7NaF·6UF₄.

3.74. The System $\text{LiF}-\text{ThF}_4-\text{UF}_4$

C. F. Weaver, R. E. Thoma, H. Insley, and H. A. Friedman, "Phase Equilibria in the Systems UF_4-ThF_4 and $\text{LiF}-\text{ThF}_4-\text{UF}_4$," paper presented at the 61st National Meeting of the American Ceramic Society, Chicago, Ill., May 17-21, 1959.

Invariant Equilibria

Composition of Liquid (mole %)			Invariant Temperature (°C)	Type of Equilibrium	Solids Present at Invariant Point
LiF	ThF_4	UF_4			
72.5	7.0	20.5	500	Peritectic	LiF , $3\text{LiF} \cdot (\text{Th}, \text{U})\text{F}_4$, and $7\text{LiF} \cdot 6(\text{U}, \text{Th})\text{F}_4$
72.0	1.5	26.5	488	Eutectic	LiF , $4\text{LiF} \cdot \text{UF}_4$, and $7\text{LiF} \cdot 6(\text{U}, \text{Th})\text{F}_4$
53	18	19	609	Peritectic	$7\text{LiF} \cdot 6(\text{U}, \text{Th})\text{F}_4$, $\text{LiF} \cdot 2(\text{Th}, \text{U})\text{F}_4$, and $\text{LiF} \cdot 4(\text{U}, \text{Th})\text{F}_4$

A three-dimensional model of the system $\text{LiF}-\text{ThF}_4-\text{UF}_4$ is shown in Fig. 3.74e.

- (a) LIQUID + $3\text{LiF} \cdot \text{ThF}_4$ ss
- (b) LiF + LIQUID
- (c) LiF + $3\text{LiF} \cdot \text{ThF}_4$ ss + LIQUID
- (d) LiF + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss + LIQUID
- (e) $4\text{LiF} \cdot \text{UF}_4$ + LIQUID + LiF
- (f) $4\text{LiF} \cdot \text{UF}_4$ + LIQUID
- (g) $4\text{LiF} \cdot \text{UF}_4$ + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss + LIQUID
- (h) $3\text{LiF} \cdot \text{ThF}_4$ ss
- (i) $3\text{LiF} \cdot \text{ThF}_4$ ss + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss + LiF
- (j) LiF + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss
- (k) LiF + $4\text{LiF} \cdot \text{UF}_4$ + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss
- (l) $4\text{LiF} \cdot \text{UF}_4$ + $7\text{LiF} \cdot 6\text{ThF}_4 - 7\text{LiF} \cdot 6\text{UF}_4$ ss

Fig. 3.74b. The System $\text{LiF}-\text{ThF}_4-\text{UF}_4$: The Section at 75 Mole % LiF .

Fig. 3.74c. The System $\text{LiF}-\text{ThF}_4-\text{UF}_4$: The Section at 53.8 Mole % LiF .

- (a) $\text{UF}_4 - \text{ThF}_4$ ss + LIQUID
 (b) $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ + LIQUID
 (c) $\text{UF}_4 - \text{ThF}_4$ ss + LIQUID + $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ ss
 (d) $\text{LiF} \cdot 2\text{ThF}_4$ ss + LIQUID + $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ ss
 (e) $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ ss + LIQUID + $7\text{LiF} \cdot 6\text{UF}_4 - 7\text{LiF} \cdot 6\text{ThF}_4$ ss
 (f) $\text{LiF} \cdot 2\text{ThF}_4$ ss
 (g) $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ ss + $7\text{LiF} \cdot 6\text{UF}_4 - 7\text{LiF} \cdot 6\text{ThF}_4$ ss
 (h) $\text{LiF} \cdot 4\text{UF}_4 - \text{LiF} \cdot 4\text{ThF}_4$ ss + $7\text{LiF} \cdot 6\text{UF}_4 - 7\text{LiF} \cdot 6\text{ThF}_4$ ss + $\text{LiF} \cdot 2\text{ThF}_4$ ss

Fig. 3.74d. The System $\text{LiF}-\text{ThF}_4-\text{UF}_4$: The Section at 33.3 Mole % LiF .

Fig. 3.74_e. Model of the System $\text{LiF}-\text{ThF}_4-\text{UF}_4$

3.75. The System $\text{NaF}-\text{ThF}_4-\text{UF}_4$: The Section $2\text{NaF}\cdot\text{ThF}_4-2\text{NaF}\cdot\text{UF}_4$

R. E. Thoma, H. Insley, H. A. Friedman, and C. F. Weaver, unpublished work performed at the Oak Ridge National Laboratory, 1958-59.

Preliminary diagram.

Fig. 3.75. The Section $2\text{NaF}\cdot\text{ThF}_4-2\text{NaF}\cdot\text{UF}_4$.

3.76. The System $\text{LiF}-\text{PuF}_3$

C. J. Barton and R. A. Strehlow, "Phase Relationships in the System Lithium Fluoride-Plutonium Fluoride," paper presented at the 135th National Meeting of the American Chemical Society, Boston, Mass., Apr. 5-10, 1959.

The system $\text{LiF}-\text{PuF}_3$ contains a single eutectic at 80.5 LiF -19.5 PuF_3 (mole %), m.p. 743°C .

Fig. 3.76. The System $\text{LiF}-\text{PuF}_3$.

3.77. The System $\text{LiCl}-\text{FeCl}_2$

C. Beusman, *Activities in the $\text{KCl}-\text{FeCl}_2$ and $\text{LiCl}-\text{FeCl}_2$ Systems*, ORNL-2323 (May 15, 1957).

The system $\text{LiCl}-\text{FeCl}_2$ forms a continuous series of solid solutions with a minimum at 60 $\text{LiCl}-40 \text{ FeCl}_2$ (mole %), m.p. 540°C .

Fig. 3.77. The System $\text{LiCl}-\text{FeCl}_2$

3.78. The System KCl-FeCl₂

C. Beusman, *Activities in the KCl-FeCl₂ and LiCl-FeCl₂ Systems*, ORNL-2323 (May 15, 1957). A nearly identical diagram to this one has been reported by H. L. Pinch and J. M. Hirshon, "Thermal Analysis of the Ferrous Chloride-Potassium Chloride System," *J. Am. Chem. Soc.* **79**, 6149-50 (1957).

Invariant Equilibria			
Mole % FeCl ₂ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
35	385	Peritectic	$L + \text{KCl} \rightleftharpoons \alpha\text{-}2\text{KCl}\cdot\text{FeCl}_2$
-	255	Inversion	$\alpha\text{-}2\text{KCl}\cdot\text{FeCl}_2 \rightleftharpoons \beta\text{-}2\text{KCl}\cdot\text{FeCl}_2$
39	355	Eutectic	$L \rightleftharpoons \alpha\text{-}2\text{KCl}\cdot\text{FeCl}_2 + \alpha\text{-KCl}\cdot\text{FeCl}_2$
50	400	Congruent melting point	$L \rightleftharpoons \alpha\text{-KCl}\cdot\text{FeCl}_2$
-	300	Inversion	$\alpha\text{-KCl}\cdot\text{FeCl}_2 \rightleftharpoons \beta\text{-KCl}\cdot\text{FeCl}_2$
53	390	Eutectic	$L \rightleftharpoons \alpha\text{-KCl}\cdot\text{FeCl}_2 + \text{FeCl}_2$

Fig. 3.78. The System KCl-FeCl₂

3.79. The System NaCl-ZrCl₄

C. J. Barton, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1956.

Preliminary diagram.

Invariant Equilibria

Mole % ZrCl ₄ in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
28.0	535 ± 5	Peritectic	$L + \text{NaCl} \rightleftharpoons 3\text{NaCl} \cdot \text{ZrCl}_4$
—	445 ± 5	Decomposition	$3\text{NaCl} \cdot \text{ZrCl}_4 \rightleftharpoons \text{NaCl} + \alpha\text{-}2\text{NaCl} \cdot \text{ZrCl}_4$
28.5	525 ± 5	Eutectic	$L \rightleftharpoons 3\text{NaCl} \cdot \text{ZrCl}_4 + \alpha\text{-}2\text{NaCl} \cdot \text{ZrCl}_4$
33.3	626 ± 5	Congruent melting point	$L \rightleftharpoons \alpha\text{-}2\text{NaCl} \cdot \text{ZrCl}_4$
—	370 ± 5	Inversion	$\alpha\text{-}2\text{NaCl} \cdot \text{ZrCl}_4 \rightleftharpoons \beta\text{-}2\text{NaCl} \cdot \text{ZrCl}_4$
58	352 ± 5	Peritectic	$L + \beta\text{-}2\text{NaCl} \cdot \text{ZrCl}_4 \rightleftharpoons \alpha\text{-}3\text{NaCl} \cdot 4\text{ZrCl}_4$
—	312 ± 5	Inversion	$\alpha\text{-}3\text{NaCl} \cdot 4\text{ZrCl}_4 \rightleftharpoons \beta\text{-}3\text{NaCl} \cdot 4\text{ZrCl}_4$
63	312 ± 5	Eutectic	$L \rightleftharpoons \beta\text{-}3\text{NaCl} \cdot 4\text{ZrCl}_4 + \text{ZrCl}_4$

Previous reports on the system NaCl-ZrCl₄ have been made by N. A. Berlozerskts and O. A. Kucherenko, *Zhur. Priklad. Khim.* **13**, 1552 (1940), I. S. Morozov and B. G. Korshunov, *Zhur. Neorg. Khim.* **1**, 145(1956), and H. H. Kellogg, L. J. Howell, and R. C. Sommer, *Physical Chemical Properties of the Systems NaCl-ZrCl₄, KCl-ZrCl₄, and NaCl-KCl-ZrCl₄*, Summary Report, NYO-3108 (April 7, 1955).

Fig. 3.79. The System NaCl-ZrCl₄.

3.80. The System KCl-ZrCl_4

C. J. Barton, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1955.

Preliminary diagram.

Invariant Equilibria

Mole % ZrCl_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
23	600 ± 5	Eutectic	$L \rightleftharpoons \text{KCl} + 2\text{KCl} \cdot \text{ZrCl}_4$
33.3	790 ± 5	Congruent melting point	$L \rightleftharpoons 2\text{KCl} \cdot \text{ZrCl}_4$
48	565 ± 5	Peritectic	$L + 2\text{KCl} \cdot \text{ZrCl}_4 \rightleftharpoons \alpha\text{-}7\text{KCl} \cdot 6\text{ZrCl}_4$
—	222 ± 4	Inversion	$\alpha\text{-}7\text{KCl} \cdot 6\text{ZrCl}_4 \rightleftharpoons \beta\text{-}7\text{KCl} \cdot 6\text{ZrCl}_4$
65	225 ± 4	Eutectic	$L \rightleftharpoons \alpha\text{-}7\text{KCl} \cdot 6\text{ZrCl}_4 + \text{ZrCl}_4$

Some phase work on this system has been reported by H. H. Kellogg, L. J. Howell, and R. C. Sommer, *Physical Chemical Properties of the Systems NaCl-ZrCl_4 , KCl-ZrCl_4 , and NaCl-KCl-ZrCl_4* , Summary Report, NYO-3108 (April 7, 1955).

Fig. 3.80. The System KCl-ZrCl_4 .

3.81. The System $\text{LiCl}-\text{UCl}_3$

C. J. Barton, A. B. Wilkerson, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953.

Preliminary diagram. The system $\text{LiCl}-\text{UCl}_3$ contains a single eutectic at 75 LiCl -25 UCl_3 (mole %), m.p. $495 \pm 5^\circ\text{C}$.

Fig. 3.81. The System $\text{LiCl}-\text{UCl}_3$

3.82. The System $\text{LiCl}-\text{UCl}_4$

C. J. Barton, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953.

Preliminary diagram.

Invariant Equilibria

Mole % UCl_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
29	415	Eutectic	$L \rightleftharpoons 2\text{LiCl} \cdot \text{UCl}_4 + \text{LiCl}$
33.3	430 ± 10	Congruent melting point	$L \rightleftharpoons 2\text{LiCl} \cdot \text{UCl}_4$
48	405	Eutectic	$L \rightleftharpoons 2\text{LiCl} \cdot \text{UCl}_4 + \text{UCl}_4$

Some data on this system were reported by C. A. Kraus in *Phase Diagrams of Some Complex Salts of Uranium with Halides of the Alkali and Alkaline Earth Metals*, M-251 (July 1, 1943).

Fig. 3.82. The System $\text{LiCl}-\text{UCl}_4$

3.83. The System NaCl- UCl_3

C. A. Kraus, unpublished work.

Preliminary diagram constructed with the author's permission from data reported in *Phase Diagrams of Some Complex Salts of Uranium with Halides of the Alkali and Alkaline Earth Metals*, M-251 (July 1, 1943).

The system NaCl- UCl_3 contains a single eutectic at 67 NaCl-33 UCl_3 (mole %), m.p. 525°C.

Fig. 3.83. The System NaCl- UCl_3 .

3.84. The System NaCl- UCl_4

C. J. Barton, R. J. Sheil, A. B. Wilkerson, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953.

Preliminary diagram.

Invariant Equilibria

Mole % UCl_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
30	430 ± 5	Eutectic	$L \rightleftharpoons \text{NaCl} + 2\text{NaCl} \cdot \text{UCl}_4$
33.3	440 ± 5	Congruent melting point	$L \rightleftharpoons 2\text{NaCl} \cdot \text{UCl}_4$
47	370 ± 5	Eutectic	$L \rightleftharpoons 2\text{NaCl} \cdot \text{UCl}_4 + \text{unidentified compound}$
57	415 ± 5	Peritectic	$L + \text{UCl}_4 \rightleftharpoons \text{unidentified compound}$

A phase diagram of this system has been reported by C. A. Kraus in *Phase Diagrams of Some Complex Salts of Uranium with Halides of the Alkali and Alkaline Earth Metals*, M-251 (July 1, 1943).

Fig. 3.84. The System NaCl- UCl_4 .

3.85. The System KCl- UCl_4

C. J. Barton, A. B. Wilkerson, T. N. McVay, R. J. Sheil, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1954.

Preliminary diagram.

Invariant Equilibria

Mole % UCl_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
25	550	Eutectic	$L \rightleftharpoons \text{KCl} + 2\text{KCl} \cdot \text{UCl}_4$
33.3	630	Congruent melting point	$L \rightleftharpoons 2\text{KCl} \cdot \text{UCl}_4$
44	330	Eutectic	$L \rightleftharpoons 2\text{KCl} \cdot \text{UCl}_4 + \text{KCl} \cdot \text{UCl}_4$
49	345	Peritectic	$L + \text{KCl} \cdot 3\text{UCl}_4 \rightleftharpoons \text{KCl} \cdot \text{UCl}_4$
61	395	Peritectic	$L + \text{UCl}_4 \rightleftharpoons \text{KCl} \cdot 3\text{UCl}_4$

A phase diagram of this system has been reported by C. A. Kraus in *Phase Diagrams of Some Complex Salts of Uranium with Halides of the Alkali and Alkaline Earth Metals*, M-251 (July 1, 1943).

Fig. 3.85. The System KCl- UCl_4 .

3.86. The System $\text{RbCl}-\text{UCl}_3$

C. J. Barton, R. J. Sheil, A. B. Wilkerson, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1954.

Preliminary diagram.

Invariant Equilibria

Mole % UCl_3 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
15	610	Eutectic	$L \rightleftharpoons \text{RbCl} + 3\text{RbCl}\cdot\text{UCl}_3$
25	745 ± 10	Congruent melting point	$L \rightleftharpoons 3\text{RbCl}\cdot\text{UCl}_3$
40	560 ± 10	Peritectic	$L + 3\text{RbCl}\cdot\text{UCl}_3 \rightleftharpoons 2\text{RbCl}\cdot\text{UCl}_3$
45.5	513 ± 5	Eutectic	$L \rightleftharpoons 2\text{RbCl}\cdot\text{UCl}_3 + \text{RbCl}\cdot\text{UCl}_3$
49	550 ± 10	Peritectic	$L + \text{UCl}_3 \rightleftharpoons \text{RbCl}\cdot\text{UCl}_3$

Fig. 3.86. The System $\text{RbCl}-\text{UCl}_3$

3.87. The System CsCl- UCl_4

C. J. Barton, A. B. Wilkerson, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1953.

Preliminary diagram.

Invariant Equilibria			
Mole % UCl_4 in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
20	505 ± 5	Eutectic	$L \rightleftharpoons \text{CsCl} + 2\text{CsCl} \cdot \text{UCl}_4$
33.3	657 ± 5	Congruent melting point	$L \rightleftharpoons 2\text{CsCl} \cdot \text{UCl}_4$
58	370 ± 5	Eutectic	$L \rightleftharpoons 2\text{CsCl} \cdot \text{UCl}_4 + \text{CsCl} \cdot 2\text{UCl}_4$
63	382 ± 5	Peritectic	$L + \text{UCl}_4 \rightleftharpoons \text{CsCl} \cdot 2\text{UCl}_4$

Fig. 3.87. The System CsCl- UCl_4

3.88. The System K_3CrF_6 – Na_3CrF_6 – Li_3CrF_6

B. J. Sturm, L. G. Overholser, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951–52.

Preliminary diagram.

Fig. 3.88. The System K_3CrF_6 – Na_3CrF_6 – Li_3CrF_6 .

4. OXIDE AND HYDROXIDE SYSTEMS

4.1. The System $\text{SiO}_2\text{--ThO}_2$

L. A. Harris, "A Preliminary Study of the Phase Equilibria Diagram of $\text{ThO}_2\text{--SiO}_2$," *J. Am. Ceram. Soc.* **42**, 74–77 (1959).

Fig. 4.1. The System $\text{SiO}_2\text{--ThO}_2$.

4.2. The System LiOH–NaOH

C. J. Barton, J. P. Blakely, K. A. Allen, W. C. Davis, and B. S. Weaver, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. A phase diagram of the system LiOH–NaOH has been reported more recently by N. A. Reshetnikov and G. M. Oonzhakov, *Zhur. Neorg. Khim.* **3**, 1433 (1958).

Invariant Equilibria

Mole % LiOH in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
27	219	Eutectic	$L \rightleftharpoons \text{NaOH} + \alpha\text{-NaOH}\cdot\text{LiOH}$
40	250	Peritectic	$L + \text{LiOH} \rightleftharpoons \alpha\text{-LiOH}\cdot\text{NaOH}$
—	180	Inversion	$\alpha\text{-LiOH}\cdot\text{NaOH} \rightleftharpoons \beta\text{-LiOH}\cdot\text{NaOH}$

Fig. 4.2. The System LiOH–NaOH.

4.3. The System LiOH-KOH

C. J. Barton, J. P. Blakely, L. M. Bratcher, and W. R. Grimes, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram.

Invariant Equilibria			
Mole % KOH in Liquid	Invariant Temperature (°C)	Type of Equilibrium	Phase Reaction at Invariant Temperature
40	315	Incongruent melting point of $3\text{LiOH}\cdot 2\text{KOH}$	$L + \text{LiOH} \rightleftharpoons 3\text{LiOH}\cdot 2\text{KOH}$
70	245	Eutectic	$L \rightleftharpoons 3\text{LiOH}\cdot 2\text{KOH} + \beta\text{-KOH}$

Fig. 4.3. The System LiOH-KOH.

4.4. The System NaOH-KOH

G. von Hevesy, "The Binary Systems NaOH-KOH, KOH-RbOH, and RbOH-NaOH," *Z. physik. Chem.* **73**, 667-84 (1910).

Fig. 4.4. The System NaOH-KOH.

4.5. The System NaOH-RbOH

G. von Hevesy, "The Binary Systems NaOH-KOH, KOH-RbOH, and RbOH-NaOH," *Z. physik. Chem.* **73**, 667-84 (1910).

Fig. 4.5. The System NaOH-RbOH.

4.6. The System $\text{Ba(OH)}_2\text{--Sr(OH)}_2$

K. A. Allen, W. C. Davis, and B. S. Weaver, unpublished work performed at the Oak Ridge National Laboratory, 1951.

Preliminary diagram. The system $\text{Ba(OH)}_2\text{--Sr(OH)}_2$ contains a single eutectic at 37 $\text{Sr(OH)}_2\text{--}63 \text{ Ba(OH)}_2$ (mole %), m.p. 360°C .

Fig. 4.6. The System $\text{Ba(OH)}_2\text{--Sr(OH)}_2$

5. AQUEOUS SYSTEMS

5.1. The System $\text{UO}_2\text{-P}_2\text{O}_5\text{-H}_2\text{O}$, 25°C Isotherm

J. M. Schreyer, "The Solubility of Uranium(IV) Orthophosphates in Phosphoric Acid Solutions," *J. Am. Chem. Soc.* **77**, 2972 (1955).

The transition point between $\text{U}(\text{HPO}_4)_2 \cdot 6\text{H}_2\text{O}$ and $\text{U}(\text{HPO}_4)_2 \cdot \text{H}_3\text{PO}_4 \cdot \text{H}_2\text{O}$ as stable solids was at $0.62 \pm 0.02 \text{ M U}$ (10.3 wt % UO_2) and $9.8 \pm 0.1 \text{ M PO}_4$ (42.6 wt % P_2O_5), density $1.63 \pm 0.03 \text{ g/ml}$. The compound $\text{U}(\text{HPO}_4)_2 \cdot \text{H}_3\text{PO}_4 \cdot \text{H}_2\text{O}$ is incongruently soluble in water.

Fig. 5.1. The System $\text{UO}_2\text{-P}_2\text{O}_5\text{-H}_2\text{O}$, 25°C Isotherm.

5.2. The System $\text{UO}_3\text{--H}_3\text{PO}_4\text{--H}_2\text{O}$, 25°C Isotherm

J. M. Schreyer and C. F. Baes, Jr., "The Solubility of Uranium(VI) Orthophosphates in Phosphoric Acid Solution," *J. Am. Chem. Soc.* **76**, 354 (1954).

The normal phosphate $(\text{UO}_2)_3(\text{PO}_4)_2 \cdot 6\text{H}_2\text{O}$ is stable below 0.025 mole % H_3PO_4 . The stability ranges are as follows:

Compound	Total Phosphate Molarity
$(\text{UO}_2)_3(\text{PO}_4)_2 \cdot 6\text{H}_2\text{O}$	<0.014
$\text{UO}_2\text{HPO}_4 \cdot 4\text{H}_2\text{O}$	0.014–6.1
$\text{UO}_2(\text{H}_2\text{PO}_4)_2 \cdot 3\text{H}_2\text{O}$	>6.1

UNCLASSIFIED
DWG 20233

Fig. 5.2 The System $\text{UO}_3\text{--H}_3\text{PO}_4\text{--H}_2\text{O}$, 25°C Isotherm. 0–60 mole % UO_3 , 0–60 mole % H_3PO_4 .

5.3. The System $\text{U}(\text{HPO}_4)_2\text{--Cl}_2\text{O}_7\text{--H}_2\text{O}$, 25°C Isotherm

J. M. Schreyer and L. R. Phillips, "The Solubility of Uranium(VI) Orthophosphates in Perchloric Acid Solutions," *J. Phys. Chem.* **60**, 588 (1956).

The stable solids in this system are $\text{U}(\text{HPO}_4)_2 \cdot 2\text{H}_2\text{O}$, $\text{U}(\text{H}_2\text{PO}_4)_2(\text{ClO}_4)_2 \cdot 4\text{H}_2\text{O}$, and $\text{U}(\text{H}_2\text{PO}_4)_2(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$. Metastable anhydrous $\text{U}(\text{HPO}_4)_2$ formed readily in all solution compositions tested. X-ray diffraction patterns indicated the existence of three polymorphic forms of $\text{U}(\text{H}_2\text{PO}_4)_2(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$.

Fig. 5.3. Schreinemaker's Projection of the System $\text{UO}_2\text{--P}_2\text{O}_5\text{--Cl}_2\text{O}_7\text{--H}_2\text{O}$ at 25°C. A projection on the $\text{UO}_2\text{--Cl}_2\text{O}_7\text{--H}_2\text{O}$ face, plotted in rectangular coordinates.

5.4. The System $\text{UO}_3\text{--Na}_2\text{O--CO}_2\text{--H}_2\text{O}$, 26°C Isotherm

C. A. Blake, C. F. Coleman, K. B. Brown, D. G. Hill, R. S. Lowrie, and J. M. Schmitt, "Studies in the Carbonate-Uranium System," *J. Am. Chem. Soc.* **78**, 5978 (1956).

Fig. 5.4a. The System $\text{UO}_3\text{--Na}_2\text{O--CO}_2\text{--H}_2\text{O}$ at 26°C. Compositions are plotted in weight per cent. I, solutions in equilibrium with sodium uranates; II, solutions in equilibrium with sodium uranyl tricarbonate; III, solutions in equilibrium with uranyl carbonate. A, Na_2CO_3 ; B, $\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$; C, $\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$; D, $\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$; E, $\text{Na}_2\text{CO}_3\cdot\text{NaHCO}_3\cdot 2\text{H}_2\text{O}$ (trona); F, NaHCO_3 ; G, UO_2CO_3 ; H, $\text{Na}_4\text{UO}_2(\text{CO}_3)_3$. The entire upper face of the triangular prism represents pure water.

The stable uranium-carbonate solids found were UO_2CO_3 and $\text{Na}_4\text{UO}_2(\text{CO}_3)_3$. The solution composition at the transition point (not determined exactly) was at least 26 wt % UO_3 , 5.6 wt % Na_2O , and 7.2 wt % CO_2 (mole ratio $\text{UO}_3:\text{Na}_2\text{O}:\text{CO}_2 = 1:1.0:1.8$).

Further data on this system are available from the American Documentation Institute (document No. 5079). These include compositions, pH's, and densities at points in and near the planes $\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, $\text{UO}_3-\text{Na}_2\text{O}-\text{H}_2\text{O}$, $\text{UO}_2\text{CO}_3-\text{Na}_2\text{CO}_3-\text{H}_2\text{O}$, $\text{Na}_4\text{UO}_2(\text{CO}_3)_3-\text{NaHCO}_3-\text{H}_2\text{O}$, $\text{UO}_3-\text{Na}_2\text{CO}_3-\text{H}_2\text{O}$, $\text{UO}_3-\text{NaHCO}_3-\text{H}_2\text{O}$, and $\text{UO}_2\text{CO}_3-\text{Na}_2\text{O}-\text{H}_2\text{O}$.

Related phase equilibria in the system $\text{Na}_2\text{O}-\text{UO}_3-\text{H}_2\text{O}$ at 50 and 75°C are reported by J. E. Ricci and F. J. Loprest, *J. Am. Chem. Soc.* **77**, 2119 (1955).

Fig. 5.4b. The Section $\text{UO}_2\text{CO}_3-\text{Na}_2\text{CO}_3-\text{H}_2\text{O}$ at 26°C.

5.5. Portions of the System $\text{ThO}_2\text{--Na}_2\text{O--CO}_2\text{--H}_2\text{O}$, 25°C Isotherm

F. A. Schimmel, unpublished work performed at the Oak Ridge National Laboratory, 1955.

The stable thorium-carbonate solids found were $\text{ThOCO}_3 \cdot 8\text{H}_2\text{O}$ and $\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$.

Ternary Points

Solid Phases Present	Composition of Solution (wt %)		
	ThO_2	Na_2O	CO_2
$\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$, NaHCO_3 , and $\text{Na}_2\text{CO}_3 \cdot \text{NaHCO}_3 \cdot 2\text{H}_2\text{O}$ (trona)	0.62	12.0	9.96
$\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$, $\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$, and trona	0.48	14.1	10.86

Fig. 5.5a. Perspective Representation of the System $\text{ThO}_2\text{--Na}_2\text{O--CO}_2\text{--H}_2\text{O}$ as a Tetrahedron.

The highest thorium solubility found was 3.35 wt % ThO_2 (3.06 wt % Na_2O , 3.60 wt % CO_2) in equilibrium with solid $\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$ and $\text{ThOCO}_3 \cdot 8\text{H}_2\text{O}$. (The terminal points of this binary line were not established.) The maximum thorium solubility in equilibrium with $\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$ was 1.25 wt % ThO_2 (12.5 wt % Na_2O , 9.4 wt % CO_2); with $\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$ and trona, the highest solubility was 0.7 wt % ThO_2 (12.6 wt % Na_2O , 10.4 wt % CO_2). The highest found with $\text{Na}_6\text{Th}(\text{CO}_3)_5 \cdot 12\text{H}_2\text{O}$ and NaHCO_3 was 1.2 wt % ThO_2 (5.0 wt % Na_2O , 6.0 wt % CO_2), which was the most dilute NaHCO_3 solution tested.

Fig. 5.5b. The System ThO_2 - Na_2O - CO_2 - H_2O at 25°C . Projection to the ThO_2 - Na_2O - CO_2 face along radii from the H_2O vertex.

Fig. 5.5c. The System ThO₂-Na₂O-CO₂-H₂O at 25°C. Projection to the ThO₂·CO₂-3Na₂O·4CO₂-H₂O internal plane, represented as an equivalent triangle, along radii from the CO₂ vertex. Portion from 70 to 100 wt % H₂O.

5.6. The System $\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, 25°C Isotherm

F. A. Schimmel, unpublished work performed at the Oak Ridge National Laboratory, 1955.

Significant changes from the data of Freeth [*Phil. Trans. Roy. Soc. London* **233**, 35 (1922)] were found throughout the range; there was closer agreement in the bicarbonate range with the data of Hill and Bacon [*J. Am. Chem. Soc.* **49**, 2487 (1927)].

Transition Points

Solid Phases Present	Composition of Solution (wt %)	
	Na_2O	CO_2
$\text{NaOH}\cdot\text{H}_2\text{O}$ and Na_2CO_3	40.5	0.85
Na_2CO_3 and $\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$	31.4	0.72
$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$	18.7	6.9
$\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$	17.85	7.8
$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3\cdot\text{NaHCO}_3\cdot 2\text{H}_2\text{O}$ (trona)	13.7	10.35
Trona and NaHCO_3	11.9	9.5

The System $\text{Na}_2\text{O}-\text{CO}_2-\text{H}_2\text{O}$, 25°C Isotherm

Composition of Saturated Solution (wt %)		Equilibrium Solid Phase
Na_2O	CO_2	
41.3	0	$\text{NaOH}\cdot\text{H}_2\text{O}$
40.5	0.85	Na_2CO_3
40.45	0.8	Na_2CO_3
31.6	0.67	Na_2CO_3
31.4	0.72	Na_2CO_3 and $\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
31.1	0.78	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
29.9	0.5	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
26.6	0.4	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
25.2	0.55	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
23.2	0.9	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
23.15	1.0	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
21.15	2.3	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
19.3	4.3	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
18.95	6.0	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$
18.57	7.5	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$ (metastable)
18.6	7.8	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$ (metastable)
18.7	6.9	$\text{Na}_2\text{CO}_3\cdot\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$
18.5	6.75	$\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$ (metastable)
17.7	7.95	$\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$ (metastable)
18.3	7.3	$\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$
17.85	7.8	$\text{Na}_2\text{CO}_3\cdot 7\text{H}_2\text{O}$ and $\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
17.3	7.88	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
16.1	7.9	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
15.6	7.92	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
15.1	8.1	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
13.7	8.85	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
13.28	9.4	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
13.55	10.15	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$
14.1	11.0	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$ (metastable)
13.7	10.35	$\text{Na}_2\text{CO}_3\cdot 10\text{H}_2\text{O}$ and trona*
17.0	13.0	Trona (metastable)
16.7	12.7	Trona (metastable)
13.8	10.35	Trona
11.9	9.5	Trona and NaHCO_3
11.55	9.3	NaHCO_3
10.5	8.8	NaHCO_3
9.4	8.0	NaHCO_3
8.1	7.5	NaHCO_3
6.4	6.5	NaHCO_3
5.4	5.75	NaHCO_3
3.9	5.0	NaHCO_3
3.56	4.95	NaHCO_3
3.47	4.93	NaHCO_3

*The formula of trona is $\text{Na}_2\text{CO}_3\cdot\text{NaHCO}_3\cdot 2\text{H}_2\text{O}$.

Fig. 5.6. The System Na₂O-CO₂-H₂O, 25°C Isotherm. 0-50 wt % Na₂O, 0-50 wt % CO₂.

5.7. Solubility of Uranyl Sulfate in Water

1. C. H. Secoy, *J. Am. Chem. Soc.* **72**, 3343 (1950) (data above 300°C).
2. C. H. Secoy, *J. Am. Chem. Soc.* **70**, 3450 (1948) (data up to 300°C).
3. L. Helmholtz and G. Friedlander, *Physical Properties of Uranyl Sulfate Solutions*, MDDC-808 (1943) (room temperature data).
4. C. Dittrich, *Z. physik. Chem.* **29**, 449 (1899).
5. E. V. Jones and W. L. Marshall, *HRP Quar. Prog. Rep. March 15, 1952*, ORNL-1280, p 180-83.
6. E. V. Jones and W. L. Marshall, *HRP Quar. Prog. Rep. Jan. 31, 1959*, ORNL-2696, p 210-11.

The two-liquid-phase region of this system in actuality must be represented by the three components UO_3 , SO_3 , and H_2O . The curve representing the boundary of liquid-liquid immiscibility is correct for stoichiometric solutions. However, the tie lines within the region of immiscibility do not connect two solutions containing stoichiometric UO_2SO_4 but rather connect solutions containing varying concentrations and ratios of UO_3 and SO_3 (Fig. 5.10). At low concentrations and high temperatures, that is, below 0.02 *m* UO_2SO_4 and above 250°C, hydrolysis occurs to produce a nonstoichiometric solution phase. The point of hydrolysis is dependent on concentration and temperature (see Sec 5.13).

Note: Revised data (ref 5) are used to construct the immiscibility boundary in the figure. Earlier data (ref 1) showed higher temperatures in this region and a possible likelihood of acid impurity in the solutions.

Fig. 5.7. The System $\text{UO}_2\text{SO}_4\text{-H}_2\text{O}$.

5.8. Two-Liquid-Phase Region of UO_2SO_4 in Ordinary and Heavy Water

E. V. Jones and W. L. Marshall, *HRP Quar. Prog. Rep. March 15, 1952*, ORNL-1280, p 180-83.

The two curves shown are boundary curves for liquid-liquid immiscibility obtained by observing the phase-transitional behavior of known $\text{UO}_2\text{SO}_4\text{-H}_2\text{O}$ (D_2O) solutions in sealed tubes as a function of temperature. For later reference, this procedure is designated the visual-synthetic method. As was described in Sec 5.7, the concentrations and compositions of the two phases within the boundary region at equilibrium cannot be obtained from this figure. Mixtures which are solutions at lower temperatures separate into two liquid phases as the temperature is raised, according to the boundary curves.

Fig. 5.8. Two-Liquid-Phase Region of UO_2SO_4 in Ordinary and Heavy Water.

5.9. The System $\text{UO}_3\text{--SO}_3\text{--H}_2\text{O}$

1. A. Colani, *Bull. soc. chim. France* [4] 43, 754–62 (1928).
2. J. S. Gill, E. V. Jones, and C. H. Secoy, *HRP Quar. Prog. Rep. Oct. 31, 1953*, ORNL-1658, p 87.

The 25° data on the SO_3 -rich side are those of Colani. The higher temperature data are those of Gill, Jones, and Secoy, obtained by filtration techniques at temperature. The number and identity of the solid phases in the UO_3 -rich regions at 25, 100, and 175°C are somewhat uncertain. Subsequent unpublished work of F. E. Clark and C. H. Secoy has disclosed that the K solid is identical (x-ray diffraction) with the mineral uranopilite, $6\text{UO}_3 \cdot \text{SO}_3 \cdot 18(?)\text{H}_2\text{O}$, that the G solid is not identifiable as any known mineral, and that at least one additional unidentified solid phase occurs at 25 and 100°. The liquidus curves for the 25 and 100° isotherms are essentially correct as shown except that the transition points (two solid phases) occur at somewhat lower concentrations than those indicated.

Fig. 5.9a. The System $\text{UO}_3\text{--SO}_3\text{--H}_2\text{O}$ at 25°C.

Fig. 5.9b. The System $\text{UO}_3\text{--SO}_3\text{--H}_2\text{O}$ at 100°C .

UNCLASSIFIED
DWG 21936

Fig. 5.9c. The System $\text{UO}_3\text{--SO}_3\text{--H}_2\text{O}$ at 175°C .

Fig. 5.9d. The System $\text{UO}_3\text{-SO}_3\text{-H}_2\text{O}$ at 250°C .

5.10. Coexistence Curves for Two Liquid Phases in the System $\text{UO}_2\text{SO}_4\text{--H}_2\text{SO}_4\text{--H}_2\text{O}$

H. W. Wright, W. L. Marshall, and C. H. Secoy, *HRP Quar. Prog. Rep. Oct. 1, 1952*, ORNL-1424, p 108.

All curves in this figure must extrapolate to the critical temperature of water (374.2°C) at zero concentration of uranium. These are boundary curves which were determined by observing sealed tubes in which immiscibility occurred in $\text{UO}_3\text{--SO}_3\text{--H}_2\text{O}$ solutions upon varying the temperature. As was explained in Sec 5.7, they do not represent concentrations and compositions of the two liquid phases within the temperature-concentration region of immiscibility but are lower solution boundary curves.

Fig. 5.10. Coexistence Curves for Two Liquid Phases in the System $UO_2SO_4-H_2SO_4-H_2O$.

5.11. Two-Liquid-Phase Region of the System $\text{UO}_2\text{SO}_4\text{--H}_2\text{SO}_4\text{--H}_2\text{O}$

R. E. Leed and C. H. Secoy, *Chem. Quar. Prog. Rep. Sept. 30, 1950*, ORNL-870, p 29-30;
C. H. Secoy, *Chem. Quar. Prog. Rep. Dec. 31, 1949*, ORNL-607, p 33-38.

In contrast with the immiscibility curves shown in Fig. 5.10, each of these curves must extrapolate to a critical temperature curve for $\text{SO}_3\text{--H}_2\text{O}$ in which a small or negligible amount of UO_3 is dissolved. Since SO_3 dissolved in H_2O (neglecting any dissolved UO_3) will elevate the critical temperature, the curves obtained from solutions containing a constant amount of free H_2SO_4 will all extrapolate to critical temperatures higher than that (374.2°C) for H_2O alone.

Fig. 5.11. Two-Liquid-Phase Region of the System $\text{UO}_2\text{SO}_4\text{--H}_2\text{SO}_4\text{--H}_2\text{O}$.

5.12. Two-Liquid-Phase Coexistence Curves for UO_3 -Rich Solutions in the System UO_3 - SO_3 - H_2O With and Without HNO_3

E. V. Jones and W. L. Marshall, *HRP Quar. Prog. Rep. July 1, 1952*, ORNL-1318, p 147-48.

These curves have been obtained by the visual-synthetic method mentioned in Sec 5.8. They are therefore boundary curves and do not describe compositions and concentrations within the liquid-liquid immiscibility region.

Fig. 5.12. Two-Liquid-Phase Coexistence Curves for UO_3 -Rich Solutions in the System UO_3 - SO_3 - H_2O With and Without HNO_3 .

5.13. Solubility of UO_3 in $\text{H}_2\text{SO}_4\text{-H}_2\text{O}$ Mixtures

W. L. Marshall, *Anal. Chem.* **27**, 1923 (1955).

The data are represented in this manner in order to show the full concentration range on one figure and to best indicate the degree of precision. The solid phase is $\alpha\text{-UO}_3\cdot\text{H}_2\text{O}$ below 205°C and $\beta\text{-UO}_3\cdot\text{H}_2\text{O}$ above 205°C . These curves represent hydrolysis equilibria.

Fig. 5.13. Solubility of UO_3 in $\text{H}_2\text{SO}_4\text{-H}_2\text{O}$ Mixtures.

5.14. Effect of Excess H_2SO_4 on the Phase Equilibria in Very Dilute UO_2SO_4 Solutions

1. W. L. Marshall, *Anal. Chem.* **27**, 1923 (1955).
2. H. W. Wright, W. L. Marshall, and C. H. Secoy, *HRP Quar. Prog. Rep. Oct. 1, 1952*, ORNL-1424, p 108.

The curves were drawn by C. H. Secoy from data used for Figs. 5.10 and 5.13.

Fig. 5.14. Effect of Excess H_2SO_4 on the Phase Equilibria in Very Dilute UO_2SO_4 Solutions.

5.15. Second-Liquid-Phase Temperatures of $\text{UO}_2\text{SO}_4\text{--Li}_2\text{SO}_4$ Solutions

R. S. Greeley, S. R. Buxton, and J. C. Griess, "High Temperature Behavior of Aqueous $\text{UO}_2\text{SO}_4\text{--Li}_2\text{SO}_4$ and $\text{UO}_2\text{SO}_4\text{--BeSO}_4$," paper presented at American Nuclear Society 2nd Winter Meeting, New York City (Oct. 1957). Also R. S. Greeley and J. C. Griess, *HRP Dynamic Solution Corrosion Studies for Quarter Ending July 31, 1956*, ORNL CF-56-7-52, p 30-31.

These data were obtained by the visual-synthetic method mentioned in Sec 5.8.

Fig. 5.15. Second-Liquid-Phase Temperatures of $\text{UO}_2\text{SO}_4\text{--Li}_2\text{SO}_4$ Solutions.

5.16. Second-Liquid-Phase Temperatures for UO_2SO_4 Solutions Containing Li_2SO_4 or BeSO_4

R. S. Greeley and J. C. Griess, *HRP Dynamic Solution Corrosion Studies for Quarter Ending July 31, 1956*, ORNL CF-56-7-52, p 30.

Liquid-liquid immiscibility occurs at the boundary curves as the temperature is raised.

Fig. 5.16. Second-Liquid-Phase Temperatures for UO_2SO_4 Solutions Containing Li_2SO_4 or BeSO_4 .

5.17. Second-Liquid-Phase Temperatures for BeSO_4 Solutions Containing UO_3

R. S. Greeley and J. C. Griess, *HRP Dynamic Solution Corrosion Studies for Quarter Ending July 31, 1956*, ORNL CF-56-7-52, p 31.

Liquid-liquid immiscibility occurs at the boundary curves as the temperature is raised.

Fig. 5.17. Second-Liquid-Phase Temperatures for BeSO_4 Solutions Containing UO_3 .

5.18. The System $\text{NiSO}_4\text{--UO}_2\text{SO}_4\text{--H}_2\text{O}$ at 25°C

E. V. Jones, J. S. Gill, and C. H. Secoy, *HRP Quar. Prog. Rep. Oct. 31, 1954*, ORNL-1813, p 166-67.

Also included in the reference are solubility data for this system at 175 and 250°C . The data at high temperature are not in sufficient quantity for a diagram to be presented.

UNCLASSIFIED
ORNL-LR-DWG 3713A

Fig. 5.18. The System $\text{NiSO}_4\text{--UO}_2\text{SO}_4\text{--H}_2\text{O}$ at 25°C . Compositions are plotted in weight per cent.

5.19. Phase-Transition Temperatures in Solutions Containing CuSO_4 , UO_2SO_4 , and H_2SO_4

F. E. Clark, J. S. Gill, R. Slusher, and C. H. Secoy, *J. Chem. Eng. Data* **4**, 12 (1959).

All data were obtained by the visual-synthetic method mentioned in Sec 5.8 and represent temperatures at which immiscibility occurred upon raising the temperature. No deductions can be made regarding compositions within the two-liquid-phase region.

Fig. 5.19. Phase-Transition Temperatures in Solutions Containing CuSO_4 , UO_2SO_4 , and H_2SO_4 .

5.20. The Effect of CuSO_4 and NiSO_4 on Phase-Transition Temperatures: $0.04 \text{ } m \text{ } \text{UO}_2\text{SO}_4$; $0.01 \text{ } m \text{ } \text{H}_2\text{SO}_4$

C. H. Secoy *et al.*, *HRP Quar. Prog. Rep.* July 31, 1957, ORNL-2379, p 163. Also additional data: F. Moseley, *Core Solution Stability in the Homogeneous Aqueous Reactor; the Effect of Corrosion Product and Copper Concentrations*, HARD(C)/P-41 (May 1957).

The curves for liquid-liquid immiscibility were drawn from data obtained by the visual-synthetic method mentioned in Sec 5.8. Saturation temperatures for solid-liquid equilibria were obtained by measuring solution concentrations as a function of temperature and determining a break in the curve. The numbers in parentheses are temperatures at which the indicated solid phase appeared upon raising the temperature. Solid phases indicated in parentheses were found in small quantity.

Fig. 5.20. The Effect of CuSO_4 and NiSO_4 on Phase-Transition Temperatures: $0.04 \text{ } m \text{ } \text{UO}_2\text{SO}_4$; $0.01 \text{ } m \text{ } \text{H}_2\text{SO}_4$.

5.21. The System $\text{UO}_3\text{--CuO--NiO--SO}_3\text{--D}_2\text{O}$ at 300°C ; $0.06\ m\ \text{SO}_3$

J. S. Gill, R. Slusher, and W. L. Marshall, *HRP Quar. Prog. Rep. Jan. 31, 1959*, ORNL-2696, p 205-13.

The variation in solution composition in the presence of one, two, and three solid phases was determined at 300°C from 0.04 to $0.2\ m\ \text{SO}_3$. Solubility relationships at $0.06\ m\ \text{SO}_3$ were obtained from these separate solubility curves, and the skeletal volume figure for the five-component system was drawn. Other skeletal figures can be drawn at various SO_3 concentrations. This investigation is still in progress, and the data are subject to revision.

Fig. 5.21. The System $\text{UO}_3\text{--CuO--NiO--SO}_3\text{--D}_2\text{O}$ at 300°C ; $0.06\ m\ \text{SO}_3$.

5.22. Solubility of $\text{Nd}_2(\text{SO}_4)_3$ in 0.02 m UO_2SO_4 Solution Containing 0.005 m H_2SO_4 (180–300°C)

R. E. Leuze *et al.*, *HRP Quar. Prog. Rep.* April 30, 1954, ORNL-1753, p 176–79.

Also included in the reference are preliminary solubility data for rare earth sulfate mixtures as well as $\text{Ce}_2(\text{SO}_4)_3$ and $\text{La}_2(\text{SO}_4)_3$ in 0.02 m UO_2SO_4 , 0.005 m H_2SO_4 .

Fig. 5.22. Solubility of $\text{Nd}_2(\text{SO}_4)_3$ in 0.02 m UO_2SO_4 Solution Containing 0.005 m H_2SO_4 (180–300°C).

5.23. Solubility of $\text{La}_2(\text{SO}_4)_3$ in UO_2SO_4 Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solution*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

Experimental data for Figs. 5.23–5.27 were obtained by the visual-synthetic method described in Sec 5.8. The very large effect of UO_2SO_4 must be noted and indicates considerable solvation of other species by aqueous UO_2SO_4 .

Fig. 5.23. Solubility of $\text{La}_2(\text{SO}_4)_3$ in UO_2SO_4 Solutions.

5.24. Solubility of CdSO_4 in UO_2SO_4 Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solution*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

See comments in Sec 5.23.

Fig. 5.24. Solubility of CdSO_4 in UO_2SO_4 Solutions.

5.25. Solubility of Cs_2SO_4 in UO_2SO_4 Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solution*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

See comments in Sec 5.23.

Fig. 5.25. Solubility of Cs_2SO_4 in UO_2SO_4 Solutions.

5.26. Solubility of $Y_2(SO_4)_3$ in UO_2SO_4 Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solution*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

See comments in Sec 5.23.

Fig. 5.26. Solubility of $Y_2(SO_4)_3$ in UO_2SO_4 Solutions.

5.27. Solubility of Ag_2SO_4 in UO_2SO_4 Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solution*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

See comments in Sec 5.23.

Fig. 5.27. Solubility of Ag_2SO_4 in UO_2SO_4 Solutions.

5.28. Solubility at 250°C of BaSO_4 in $\text{UO}_2\text{SO}_4\text{-H}_2\text{O}$ Solutions

E. V. Jones, M. H. Lietzke, and W. L. Marshall, *J. Am. Chem. Soc.* **79**, 267 (1957); also *The Solubility of Several Metal Sulfates at High Temperatures and Pressures in Water and in Aqueous Uranyl Sulfate Solutions*, ORNL CF-55-7-69 (declassified Aug. 23, 1955).

Solutions were filtered from solids at 250°C to obtain the solubility of BaSO_4 in H_2O . Counting of radioactive barium was used to determine the solubility concentration. The very large effect of UO_2SO_4 must be noted and indicates considerable solvation of other species by aqueous UO_2SO_4 .

Fig. 5.28. Solubility at 250°C of BaSO_4 in $\text{UO}_2\text{SO}_4\text{-H}_2\text{O}$ Solutions.

5.29. Solubility of H_2WO_4 in 0.126 and 1.26 M UO_2SO_4

C. E. Coffey, W. L. Marshall, and G. H. Cartledge, *HRP Quar. Prog. Rep. Oct. 31, 1953*, ORNL-1658, p 96-99.

The data shown in the figures were obtained by direct sampling of equilibrated solutions, by the filter bomb method, and by the synthetic method described previously in Sec 5.8. In 0.126 M UO_2SO_4 the solubility of H_2WO_4 first shows a positive temperature coefficient of solubility and then, above 100°C, a negative coefficient of solubility. In 1.26 M UO_2SO_4 the temperature coefficient of solubility is positive up to 285°, the temperature at which two liquid phases form – a characteristic of $\text{UO}_2\text{SO}_4\text{--H}_2\text{O}$ solutions.

Fig. 5.29b. Solubility of H_2WO_4 in 1.26 M UO_2SO_4 .

5.30. Phase Stability of H_2WO_4 in 1.26 M UO_2F_2

C. E. Coffey, W. L. Marshall, and G. H. Cartledge, *HRP Quar. Prog. Rep. Oct. 31, 1953*, ORNL-1658, p 96-99.

The methods of solubility and stability determination were the same as for analogous studies of H_2WO_4 in UO_2SO_4 (Sec 5.29). The data on the figure as drawn show temperatures at which hydrolytic precipitation of an unidentified solid phase occurs from solutions stable at lower temperatures.

Fig. 5.30. Phase Stability of H_2WO_4 in 1.26 M UO_2F_2 .

5.31. The System $\text{UO}_2(\text{NO}_3)_2\text{-H}_2\text{O}$

1. Reported in full by W. L. Marshall, J. S. Gill, and C. H. Secoy, *Chem. Quar. Prog. Rep. March 31, 1951*, ORNL-1053, p 22-25.

2. Reported in part by W. L. Marshall, J. S. Gill, and C. H. Secoy, *J. Am. Chem. Soc.* **73**, 1867 (1951).

As in the case of the system $\text{UO}_2\text{SO}_4\text{-H}_2\text{O}$, hydrolysis occurs at high temperatures and low concentrations, resulting in precipitation of $\text{UO}_3\cdot\text{H}_2\text{O}$ from stoichiometric $\text{UO}_2(\text{NO}_3)_2$ solutions. In addition, decomposition of NO_3^- occurs at elevated temperature to produce an equilibrium vapor phase of nitrogen oxides. Letter *A* represents a region of complete solution. Data at low temperature up to point *F* were obtained by direct analysis of equilibrated solutions, whereas the high-temperature data were obtained by the visual-synthetic method described in Sec 5.8.

Fig. 5.31. The System UO₂(NO₃)₂-H₂O.

5.32. Phase Equilibria of UO_3 and HF in Stoichiometric Concentrations (Aqueous System)

W. L. Marshall, J. S. Gill, and C. H. Secoy, *J. Am. Chem. Soc.* **76**, 4279 (1954).

The two-liquid-phase region and the region of "basic" solid solution + saturated solution are in actuality segments of the three-component system $\text{UO}_3\text{--HF--H}_2\text{O}$, in which the equilibrium phases are nonstoichiometric with regard to UO_2F_2 . This system is somewhat analogous to the system $\text{UO}_3\text{--H}_2\text{SO}_4\text{--H}_2\text{O}$ (see Sec 5.7).

Fig. 5.32. Phase Equilibria of UO_3 and HF in Stoichiometric Concentrations (Aqueous System).

5.33. Solubility of Uranium Trioxide in Orthophosphoric Acid Solutions

W. L. Marshall, J. S. Gill, and H. W. Wright, *HRP Quar. Prog. Rep. May 15, 1951*, ORNL-1057, p 112-15.

The data were obtained for the most part by equilibrating solutions of known concentration in sealed tubes for various times at different temperatures. Phase changes were observed visually.

Fig. 5.33. Solubility of Uranium Trioxide in Orthophosphoric Acid Solutions.

5.34. Solubility of Uranium Trioxide in Phosphoric Acid at 250°C

J. S. Gill, W. L. Marshall, and H. W. Wright, *HRP Quar. Prog. Rep. Aug. 15, 1951*, ORNL-1121, p 119-21.

Solution and solid mixtures were equated at 250°C in sealed glass tubes. The tubes were cooled rapidly to room temperature, and the solution and solid phases were analyzed. The slow reversibility of equilibrium conditions made this procedure possible.

Fig. 5.34. Solubility of Uranium Trioxide in Phosphoric Acid at 250°C.

5.35. Solubility of UO_3 in H_3PO_4 Solution

B. J. Thamer *et al.*, *The Properties of Phosphoric Acid Solutions of Uranium as Fuels for Homogeneous Reactors*, LA-2043 (March 6, 1956); W. L. Marshall, J. S. Gill, and H. W. Wright, *HRP Quar. Prog. Rep.* May 15, 1951, ORNL-1057, p 112-15; and J. S. Gill, W. L. Marshall, and H. W. Wright, *HRP Quar. Prog. Rep.* Aug. 15, 1951, ORNL-1121, p 119-21.

This figure represents a compilation from the data of B. J. Thamer *et al.* of Los Alamos and W. L. Marshall *et al.* of Oak Ridge National Laboratory.

Fig. 5.35. Solubility of UO_3 in H_3PO_4 Solution.

5.36. The System $\text{UO}_2\text{CrO}_4\text{--H}_2\text{O}$

1. F. J. Loprest, W. L. Marshall, and C. H. Secoy, *J. Am. Chem. Soc.* **77**, 4705 (1955).
2. W. L. Marshall, *HRP Quar. Prog. Rep. March 31, 1953*, ORNL-1554, p 105-6.

The solid phases are $\text{UO}_2\text{CrO}_4 \cdot 5\frac{1}{2}\text{H}_2\text{O}$ (A) and $\text{UO}_2\text{CrO}_4 \cdot x\text{H}_2\text{O}$ (B). Curve *rs* represents the boundary of a region in which hydrolytic precipitation of a basic uranyl chromate occurs. In the same concentration range the dichromate, $\text{UO}_2\text{Cr}_2\text{O}_7$, is stable to the critical temperature and apparently dissolves in the supercritical fluid (ref 2).

Fig. 5.36. The System $\text{UO}_2\text{CrO}_4\text{--H}_2\text{O}$.

5.37. Variation of Li_2CO_3 Solubility with UO_2CO_3 Concentration at Constant CO_2 Pressure (250°C)

1. F. J. Loprest, W. L. Marshall, and C. H. Secoy, *HRP Quar. Prog. Rep. July 31, 1955*, ORNL-1943, p 227-35.

2. W. L. Marshall, F. J. Loprest, and C. H. Secoy, *HRP Quar. Prog. Rep. Jan. 31, 1956*, ORNL-2057, p 131-32.

The solubility relationships in the figure indicate strong complexing of UO_2CO_3 with Li_2CO_3 under CO_2 pressure. The formation of HCO_3^- species in solution is indicated.

Fig. 5.37. Variation of Li_2CO_3 Solubility with UO_2CO_3 Concentration at Constant CO_2 Pressure (250°C).

5.38. The System $\text{Li}_2\text{O}-\text{UO}_3-\text{CO}_2-\text{H}_2\text{O}$ at 250°C and 1500 psi

1. F. J. Loprest, W. L. Marshall, and C. H. Secoy, *HRP Quar. Prog. Rep. July 31, 1955*, ORNL-1943, p 227-35.

2. W. L. Marshall, F. J. Loprest, and C. H. Secoy, *HRP Quar. Prog. Rep. Jan. 31, 1956*, ORNL-2057, p 131-32.

As CO_2 pressure is increased, the area of complete solution (indicated by the shading) increases. As temperature is increased, the solution area decreases.

UNCLASSIFIED
ORNL-LR-DWG 11342B

Fig. 5.38. The System $\text{Li}_2\text{O}-\text{UO}_3-\text{CO}_2-\text{H}_2\text{O}$ at 250°C and 1500 psi.

5.39. The System $\text{Th}(\text{NO}_3)_4\text{--H}_2\text{O}$

1. Reported in full by W. L. Marshall, J. S. Gill, and C. H. Secoy, *HRP Quar. Prog. Rep.* Nov. 30, 1950, ORNL-925, p 279-90.

2. W. L. Marshall, J. S. Gill, and C. H. Secoy, *J. Am. Chem. Soc.* **73**, 4991 (1951).

Hydrolytic precipitation of ThO_2 and thermal decomposition of NO_3^- in this system occur in an analogous manner to the reactions occurring in the system $\text{UO}_2(\text{NO}_3)_2\text{--H}_2\text{O}$ (see Sec 5.31). At higher temperatures the two-component system in actuality must be considered in terms of the three components ThO_2 , HNO_3 , and H_2O .

Fig. 5.39. The System $\text{Th}(\text{NO}_3)_4\text{--H}_2\text{O}$.

5.40. Hydrolytic Stability of Thorium Nitrate–Nitric Acid and Uranyl Nitrate Solutions

1. W. L. Marshall and C. H. Secoy, *HRP Quar. Prog. Rep. Oct. 31, 1953*, ORNL-1658, p 93–96.

2. W. L. Marshall, J. S. Gill, and C. H. Secoy, *Chem. Quar. Prog. Rep. March 31, 1951*, ORNL-1053, p 22–25.

The data used to draw the individual curves were obtained by observations at various temperatures of tubes that contained different solutions of known concentration. The procedure was analogous to that used for obtaining data for Fig. 5.33. The curve for hydrolytic precipitation of $\text{UO}_2(\text{NO}_3)_2$ is taken from ref 2.

Fig. 5.40. Hydrolytic Stability of Thorium Nitrate–Nitric Acid and Uranyl Nitrate Solutions.

5.41. The System $\text{ThO}_2\text{-CrO}_3\text{-H}_2\text{O}$ at 25°C

1. H. T. S. Britton, *J. Chem. Soc.* **123**, 1429 (1923).
2. W. L. Marshall, F. J. Loprest, and C. H. Secoy, *HRP Quar. Prog. Rep. Jan. 31, 1955*, ORNL-1853, p 205-6.

The phase diagram at 25° was determined by Britton. Points 1-7 indicate compositions investigated at high temperature by Marshall, Loprest, and Secoy. These solution compositions were phase stable up to 160°C for point 7 and 320°C for point 1.

Fig. 5.41. The System $\text{ThO}_2\text{-CrO}_3\text{-H}_2\text{O}$ at 25°C . Compositions are in weight per cent.

5.42. Phase Stability of $\text{ThO}_2\text{-H}_3\text{PO}_4\text{-H}_2\text{O}$ Solutions at High Concentrations of ThO_2

W. L. Marshall, experimental data in ORNL research notebook 2881, p 35 (May 1953).

Solutions of appropriate concentrations were prepared and analyzed. These solutions were sealed in glass tubes and shaken at various temperatures for times varying from 1 hr to several days. In this manner the phase stability boundaries with respect to concentration and temperature were established. At first inspection of the figure it is surprising that more ThO_2 can be dissolved per liter at the lower $\text{H}_3\text{PO}_4/\text{ThO}_2$ ratio than at the higher ratio. Actually, there is a physical volume limitation due to the high fractional volumes occupied by ThO_2 and H_3PO_4 at these concentrations. As any solution is diluted with H_2O , hydrolysis of thorium in solution occurs. At room temperature, solutions in which the $\text{H}_3\text{PO}_4/\text{ThO}_2$ ratio is 5:1 are gels, whereas 10:1 ratio solutions are still fluid.

Fig. 5.42. Phase Stability of $\text{ThO}_2\text{-H}_3\text{PO}_4\text{-H}_2\text{O}$ Solutions at High Concentrations of ThO_2

ACKNOWLEDGMENT

Acknowledgment is made to the following for permission to reproduce copyrighted material:

American Ceramic Society

American Institute of Mining, Metallurgical, and Petroleum Engineers

Analytical Chemistry

Journal of Chemical and Engineering Data

Journal of Physical Chemistry

Journal of the American Chemical Society

U.S. Atomic Energy Commission

-

-

.

INTERNAL DISTRIBUTION

1. R. D. Ackley	55. J. S. Culver
2. R. E. Adams	56. D. R. Cuneo
3. G. M. Adamson	57. J. E. Cunningham
4. L. G. Alexander	58. D. G. Davis
5. K. A. Allen	59. R. J. Davis
6. C. F. Baes	60. W. C. Davis
7-11. C. J. Barton	61. J. H. DeVan
12. S. E. Beall	62. R. R. Dickison
13. P. R. Bell	63. L. M. Doney
14. J. O. Betterton	64. F. A. Doss
15. A. M. Billings	65. J. S. Drury
16. D. S. Billington	66. A. S. Dworkin
17. C. A. Blake	67. L. B. Emler (K-25)
18. J. P. Blakely	68. J. L. English
19. R. E. Blanco	69. J. E. Eorgan
20. F. F. Blankenship	70. R. B. Evans
21. M. Blander	71. D. E. Ferguson
22. E. P. Blizzard	72. J. L. Fowler
23. C. M. Blood	73. A. P. Fraas
24. A. L. Boch	74. H. A. Friedman
25. E. G. Bohlman	75. J. H. Frye, Jr.
26. S. E. Bolt	76. C. H. Gabbard
27. G. E. Boyd	77. W. R. Gall
28. M. A. Bredig	78. J. S. Gill
29. J. C. Bresee	79. J. H. Gillette
30. R. B. Briggs	80. L. O. Gilpatrick
31. H. R. Bronstein	81. J. M. Googin (Y-12)
32. J. R. Brown	82. R. S. Greeley
33. K. B. Brown	83. A. T. Gresky
34. W. E. Browning	84. J. C. Griess
35. F. R. Bruce	85-89. W. R. Grimes
36. W. D. Burch	90. E. Guth
37. S. R. Buxton	91. P. A. Haas
38. S. Cantor	92. P. H. Harley
39. W. H. Carr	93. C. S. Harrill
40. W. L. Carter	94. L. A. Harris
41. G. H. Cartledge	95. P. N. Haubenreich
42. G. I. Cathers	96. G. M. Hebert
43. C. E. Center	97. H. L. Hemphill
44. R. H. Chapman	98. J. W. Hill
45. R. A. Charpie	99. E. C. Hise
46. R. D. Cheverton	100. E. E. Hoffman
47. H. C. Claiborne	101. H. W. Hoffman
48. G. W. Clark	102. A. Hollaender
49. C. F. Coleman	103. R. W. Horton
50. E. L. Compere	104. A. S. Householder
51. J. A. Cox	105. H. Insley
52. J. H. Crawford	106. G. H. Jenks
53. C. R. Croft	107. D. T. Jones
54. F. L. Culler	108. E. V. Jones

109. W. H. Jordan
110. P. R. Kasten
111. G. W. Keilholtz
112. C. P. Keim
113. M. T. Kelley
114. M. J. Kelley
115. B. W. Kinyon
116. N. A. Krohn
117. J. A. Lane
118. S. Langer
119. C. G. Lawson
120. J. E. Lee, Jr.
121. R. E. Leed
122. R. E. Leuze
123. W. H. Lewis
124. M. H. Lietzke
125. T. A. Lincoln
126. S. C. Lind
127. R. B. Lindauer
128. R. S. Livingston
129. J. T. Long
130. R. A. Lorenz
131. R. E. Lowrie
132. M. I. Lundin
133. R. N. Lyon
134. J. P. McBride
135. W. B. McDonald
136. H. F. McDuffie
137. H. A. McLain
138. R. A. McNees
139. H. G. MacPherson
140. T. N. McVay
141. J. R. McWherter
142. W. D. Manly
- 143-147. W. L. Marshall
148. R. E. Meadows
149. R. P. Metcalf
150. R. P. Milford
151. A. J. Miller
152. J. W. Miller
153. C. S. Morgan
154. K. Z. Morgan
155. R. E. Moore
156. R. L. Moore
157. J. P. Murray (Y-12)
158. M. L. Nelson
159. G. J. Nessel
160. R. F. Newton
161. L. G. Overholser
162. L. F. Parsly, Jr.
163. R. L. Pearson
164. F. N. Peebles
165. D. Phillips
166. L. R. Phillips (Y-12)
167. M. L. Picklesimer
168. W. T. Rainey
169. A. D. Redman
170. S. A. Reed
171. P. M. Reyling
172. D. M. Richardson
173. R. C. Robertson
174. A. D. Ryon
175. H. C. Savage
176. H. W. Savage
177. F. A. Schimmel (Y-12)
178. J. M. Schmitt
179. J. M. Schreyer (Y-12)
180. H. E. Seagren
181. C. H. Secoy
182. C. L. Segaser
183. J. H. Shaffer
184. E. M. Shank
185. R. J. Sheil
186. R. P. Shields
187. E. D. Shipley
188. M. D. Silverman
189. M. J. Skinner
190. R. Slusher
191. N. V. Smith
192. W. T. Smith, Jr.
193. A. H. Snell
194. B. A. Soldano
195. I. Spiewak
196. H. H. Stone
197. R. A. Strehlow
198. B. J. Sturm
199. C. D. Susano
200. J. E. Sutherland
201. J. A. Swartout
202. A. Taboada
203. E. H. Taylor
- 204-213. R. E. Thoma
214. D. G. Thomas
215. M. Tobias
216. D. S. Toomb
217. D. B. Trauger
218. J. Truitt
219. W. E. Unger
220. R. Van Winkle
221. F. C. VonderLage
222. W. T. Ward
223. G. M. Watson
224. B. S. Weaver
225. C. F. Weaver

226. A. M. Weinberg	238. W. O. Milligan (consultant)
227. K. W. West	239. J. E. Ricci (consultant)
228. M. E. Whatley	240. G. T. Seaborg (consultant)
229. G. C. Williams	241. E. P. Wigner (consultant)
230. C. E. Winters	242. Biology Library
231. H. W. Wright	243-244. Central Research Library
232. J. P. Young	245. Health Physics Library
233. F. C. Zapp	246-265. Laboratory Records Department
234. P. H. Emmett (consultant)	266. Laboratory Records, ORNL R.C.
235. H. Eyring (consultant)	267. Reactor Experimental Engineering Library
236. D. G. Hill (consultant)	268. ORNL - Y-12 Technical Library,
237. C. E. Larson (consultant)	Document Reference Section

EXTERNAL DISTRIBUTION

- 269-270. Stanford Research Institute, Menlo Park, Calif. (1 copy ea. to D. Cubicciotti and J. W. Johnson)
- 271-272. National Lead Co., P. O. Box 158, Cincinnati 39, Ohio (1 copy ea. to R. E. DeMarco and K. Notz)
273. Geophysical Laboratory, Carnegie Institution of Washington, 2801 Upton St., Washington 8, D.C. (J. W. Grieg)
274. Mound Laboratory, Miamisburg, Ohio (E. F. Eichelberger)
- 275-276. Minnesota Mining and Manufacturing Co., Central Research Dept., 2301 Hudson Rd., St. Paul 9, Minn. (1 copy ea. to J. R. Johnson and G. D. White)
277. Chemical Separations and Development Branch, Nuclear Technology, Division of Reactor Development, AEC, Washington 25, D.C. (F. Kerze, Jr.)
278. Dept. of Chemistry, Brown University, Providence 12, R.I. (C. A. Kraus)
279. Mallinckrodt Chemical Works, Uranium Division, P.O. Box 472, St. Charles, Mo. (C. W. Kuhlman)
280. Portland Cement Association Fellowship, National Bureau of Standards, Washington 25, D. C. (E. M. Levin)
- 281-282. National Carbon Research Laboratories, P.O. Box 6116, Cleveland, Ohio (1 copy ea. to G. W. Mellors and S. Senderoff)
283. College of Mineral Industries, Pennsylvania State University, University Park, Pa. (E. F. Osborn)
284. Los Alamos Scientific Laboratory, P.O. Box 1663, Los Alamos, N. M. (B. J. Thamer)
285. Dept. of Physics, University of Chicago, Chicago, Ill. (W. H. Zachariasen)
286. Reaction Motors Division, Thiokol Chemical Corp., Denville, N.J. (F. L. Loprest)
287. National Bureau of Standards, Washington 25, D. C. (H. F. McMurdie)
288. U.S. Geological Survey, Washington 25, D. C. (G. W. Morey)
289. National Carbon Co., Union Carbide Corporation, Parma, Ohio (F. E. Clark)
290. Division of Research and Development, AEC, Washington
291. Division of Research and Development, AEC, ORO
292. Division of Reactor Development, AEC, Washington
293. Division of Reactor Development, AEC, ORO
294. Oak Ridge Institute of Nuclear Studies, Oak Ridge
- 295-851. Given distribution as shown in TID-4500 (15th ed.) under Chemistry-General category (75 copies - OTS)