### IONS OF MIXTURES

ne a copy of ref. (3) in advance et alculations of the excess function. Dr. I. R. McKinnon in his thesis, wing my attention to the work of m similar to eqn (13) with (19) responding term from the Percus

. Phys., 1967, 11, 117. ns. Faraday Soc., 1968, 64, 1447. J. Watson, Trans. Faraday Soc., 1969, 65.

1967). California, Los Angeles, 1968). Holland, Amsterdam, 1957), (a) chap. 8

1952, 20, 1538; 1953, 21, 2169.

d, J. Chem. Phys., 1957, 26, 1533; P. J 1, 1957, 27, 505. cc. Roy. Soc. A, 1957, 240, 651.

1964, 8, 549. 2 und Flüssigen Zustandes (Barth, Leipzig

vorths, London, 1969), 2nd ed., chap. 8, 9 y. Soc. A, 1951, 206, 448. Phys. Chem., 1967, 71, 3477. ysical and Chemical Constants, originally ans, London, 1966), 13th ed., p. 150. ci, Chem. Rev., 1968, 68, 659. , 670, 1441; I. M. Croll and R. L. Scott

1967, 47, 2248.

# Solid-Liquid Phase Equilibria in the Sodium-Rubidium Alloy System

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#### Received 27th June, 1969

thermal methods were used to determine with high precision, the solid-liquid phase equilibria for the sodium-rubidium system. The results differ greatly from those of earlier workers, scally near the eutectic composition where differences in melting points as large as 25 K occur. We can do not solve the eutectic composition where differences in melting points as large as 25 K occur. We can do not solve the eutectic temperature of the eutectic temperature produced any evidence for compound formation. X and for solid compound formation. Again, no evidence was the for solid compound formation.

I reezing point measurements have been made by several workers <sup>1-3</sup> on solutions badium and rubidium in an attempt to determine the solid-liquid phase diagram t this system. The measurements were made without the advantage of high purity stals, platinum resistance thermometry, and modern inert atmosphere facilities. As tesult, the data are in poor agreement, especially on the rubidium-rich side of the feetic, where few data points were obtained. Comparison of the data available beated differences in temperature of as much as 15° between the data of Rinck <sup>1</sup> and at of Goria.<sup>2</sup> No evidence was found in any of the previous work for intermetallic "pound formation. This seemed surprising since a peritectically melting Na<sub>2</sub>K termetallic compound had been well established in the sodium-potassium system,<sup>4,5</sup> d a sluggishly forming peritectically melting Na<sub>2</sub>Cs compound had been reported the sodium-caesium system.<sup>6,7</sup>

As a part of our investigation of alkali metal mixtures, we sought to obtain a studed and accurate solid-liquid phase diagram for this system. Of special interest stars the possibility for the formation of intermetallic compounds such as  $Na_2Rb$ .

#### EXPERIMENTAL

#### CHEMICALS

If the purity (99.8 % minimum) rubidium was obtained from the Kawecki Chemical rany. Batch analysis of the material by Kawecki Chemical indicated 0.034 mol % Cs, and % K, 0.018 mol % Na, 0.015 mol % Si with negligible amounts of other tattes. Oxygen analysis was not available. However, calculations from the change in g point with fraction melted indicated less than 0.02 mol % oxygen. The rubidium readered to be better than 99.9 % pure.

tor Grade sodium was obtained from the U.S. Industrial Chemical Company. stions for Reactor Grade sodium limit the impurities to <200 p.p.m. K, <100 p.p.m. <50 p.p.m. Rb and smaller amounts of other impurities. Comparison of the point of this sodium with our measurements of an ultra-pure sample as described therature <sup>5</sup> limits the impurities to <0.05 mol % total.