Thermal Properties of Aqueous Uni-univalent Electrolytes

Vivian Barfield Parker



NSRDS-NBS 2 National Standard Reference Data Series National Bureau of Standards 2

(Category 5—Thermodynamic and Transport Properties)

Issued April 1, 1965

Foreword

In 1963 the President's Office of Science and Technology established the National Standard Reference Data System to ensure that the technical community of the United States has maximum access to compilations of critically evaluated data in the physical sciences. The National Bureau of Standards was assigned administrative responsibility for this Government-wide activity.

The National Bureau of Standards, as well as other groups both in this country and abroad, has been active in the compilation of standard reference data for many years. However, in view of the great accumulation of unevaluated data, the accelerated production of new data, the known time lag in incorporating these data into critically evaluated compilations, and the urgent needs of the scientific and technical community, it was decided by the Federal Council for Science and Technology that a substantially greater effort, planned and coordinated on a national basis, was needed. The National Standard Reference Data System (NSRDS) is the outgrowth of that decision.

The NSRDS is conducted as a decentralized operation across the country, with central coordination by the NBS. It comprises a complex of data centers and other activities, carried on in government agencies, academic institutions, and non-government laboratories. The independent operational status of existing critical data projects is maintained and encouraged. The activities of data centers which are components of the NSRDS include the preparation of compilations of critically evaluated data, the preparation of critical reviews of the quantitative state of knowledge in a specialized area, and computations of useful functions derived from standard reference data.

The present publication is considered to be a product of the National Standard Reference Data System activities at the NBS. It consists of a compilation of critically evaluated data on the thermodynamic properties of aqueous uni-univalent electrolytes, prepared as one of the continuing projects of the NBS Thermochemistry Section.

A. V. ASTIN, Director.

Preface

Since 1940 the National Bureau of Standards has maintained in its Thermochemistry Section a program on the collection, analysis, and evaluation of chemical thermodynamic data on chemical substances. One of the important areas of consideration has been the thermochemistry of aqueous electrolyte solutions. Because of the comparatively regular behavior of dilute univalent electrolyte solutions in water, as predicted by classical theoretical considerations, one can establish consistent values for the heats and free energies of formation and entropies of the hypothetical single-ion species, once a value for one particular ion is assigned.

As part of the planned total review of the available chemical thermodynamic data, a systematic review of the heat-capacity, heat-of-solution, and heat-of-dilution data on simple 1-1 electrolytes has been made. From the analysis of these data, tables of selected "best" values of "apparent" heat capacities, and heats of dilution are given, as well as selected values of the heats of solution to infinite dilution. Also included is a review of data on the heats of neutralization of monobasic acids which has led to a selected "best" value for the heat of ionization of water.

The author wishes to acknowledge the many stimulating and fruitful discussions with D. D. Wagman and W. H. Evans of the Thermochemistry Section. The continued financial assistance by the Division of Research of the U.S. Atomic Energy Commission is also gratefully acknowledged.

VIVIAN B. PARKER.

Contents

	ord	
Prefac	e	
I. Introd	uction	
II. Units	of energy and constants	
III. Uncertainties		
IV. The apparent molal heat capacity		
V. Heats of dilution VI. Heat of neutralization		
	w papers	
	ences	
X. Tables	5	
	List of Figures	
Figure I.	The apparent molal heat capacity of KCl at 25 °C	
Figure II.	The apparent molal heat capacity: acids and lithium compounds	
Figure III.	The apparent molal heat capacity: ammonium and sodium compounds	
Figure IV.	The apparent molal heat capacity: potassium, rubidium, and cesium com-	
	pounds	
Figure V.	The apparent molal heat capacity: the oxyanions of hydrogen, sodium and	
Ü	potassium	
Figure VI.	The apparent molal heat capacity: the chlorides	
Figure VII.	The apparent molal heat capacity: carboxylates	
Figure VIII.		
Figure IX.	The chord area plot for NaCl at 25 °C	
Figure X.	The relative apparent molal enthalpy: acids and ammonium and lithium	
	compounds	
Figure XI.	The relative apparent molal enthalpy: sodium compounds	
Figure XII.	The relative apparent molal enthalpy: potassium, rubidium and cesium	
-	compounds	
Figure XIII.	The relative apparent molal enthalpy: the hydroxides and chlorides	
	The relative apparent molal enthalpy: concentrated solutions	

Thermal Properties of Aqueous Uni-univalent Electrolytes

Vivian Barfield Parker

The available specific heat, heat of dilution, and heat of solution data for aqueous solutions of uni-univalent electrolytes have been critically reviewed and tables of selected "best" values at 25 °C prepared. In addition, the neutralization data have been critically reviewed in order to obtain the "best" value, 13,345 cal/mole, for the heat of ionization of water at 25 °C.

The electrolytes reviewed include the hydroxides, halides and oxyhalides, nitrates and nitrites,

formates and acetates, cyanides, cyanates and thiocyanates, and permanganates of hydrogen, ammonium and the methyl ammonium derivatives, silver, and the alkali metals.

I. Introduction

The data on the thermal properties of aqueous uni-univalent electrolytes have been critically reviewed by Rossini [1, 2, 3, 4, 5], Bichowsky and Rossini [6], Rossini et al. [7], Harned and Owen [8, 9], Pitzer and Brewer [10], Britske et al. [11], Lange and Robinson [12], and Lange and Monheim [13]. In view of the current program of the Thermochemistry Section on the revision of the tables of Selected Values of Chemical Thermodynamic Properties of chemical substances, it was felt advisable to re-examine all the available data on the thermal properties of the aqueous uni-univalent electrolytes and select systematically the most consistent set of values. It is hoped that this review, in addition to making available the experimental data, will serve to indicate areas where the data are inadequate or completely nonexistent.

The terminology and conventions used are those of Lewis and Randall [14], Lewis, Randall, Pitzer, and Brewer [10], Rossini [2, 3, 4], Harned and Owen [9], and Glasstone [15]. A summary of the terms

used and their definitions follows:

 Φ_c is the apparent molal heat capacity of the solute, equal to $[(1000 + mM_2)C - 1000 C^{\circ}]/m$ where C and C° are the specific heats (per unit mass) of the solution and pure solvent, respectively, m is the molality, and M_2 is the molecular weight of the solute.

 ΔH_{diln} , the integral heat of dilution, is the change in enthalpy, per mole of solute, when a solution of concentration m1 is diluted to a final finite concen-

If the dilution is carried out by the addition of an infinite amount of solvent, so that the final solution is infinitely dilute, the enthalpy change is the integral heat of dilution to infinite dilution. Since Φ_L , the relative apparent molal enthalpy, is equal to and of opposite sign to this, only Φ_L is referred to here when discussing dilution to infinite dilution.

 \bar{L}_2 is the relative partial molal enthalpy of the solute in the given solution, equal to $\Phi_L + \frac{1}{2}m^{1/2}$

 $(d\Phi_L/dm^{1/2}).$

 L_1 is the relative partial molal enthalpy of the solvent in the given solution, equal to $-(M_1m^{3/2}/2000)(d\Phi_L/dm^{1/2})$, where M_1 is the molecular weight of the solvent. It is also the partial or differential heat of dilution and may be defined as the change

in enthalpy per mole of solvent when it is added to a large volume of the solution at the given concentration.

 $\overline{J_2}$ is the relative partial molal heat capacity of the solute, equal to $\overline{C}_2 - \overline{C}_2^\circ$, which is equal to $(\partial \overline{L}_2/\partial T)_p$, where \overline{C}_2 is equal to $\Phi_C + \frac{1}{2} m^{1/2} (d\Phi_C/dm^{1/2})$ and \overline{C}_2° is equal to Φ_C° .

 Φ_C° is the value of the apparent molal heat capacity of the solute at infinite dilution. It is not to be confused with the heat capacity of the pure solvent.

The limiting slopes of the apparent molal properties were taken as two thirds the limiting slopes for the partial molal properties given by Harned and Owen [9].

 ΔH_{soln} is the total or integral heat of solution of one mole of solute to form a solution of concentration m. It is equal to $(H_m-55.506H_1-mH_2)/m$, where H_m is the enthalpy of the solution, and H_1 and H2 are the molar enthalpies of the pure solvent

and solute, respectively.

The partial or differential heat of solution is the increase in enthalpy per mole of solute when it is dissolved in a large volume of solution at a concentration m, so that there is no appreciable change in the latter. It is equal to $\overline{L_2} - L_2$ where L_2 is the relative partial molal enthalpy of the pure solute.

 $\Delta H_{\infty}^{\circ}$ is the heat of solution at infinite dilution,

equal to $\Delta H_{\text{soin}}(m) - \Phi_L(m)$.

 ΔH_N° is the heat of neutralization at infinite dilution. In a reaction between a strong acid and a strong base at finite and equal concentrations, given by $H\bar{X} \cdot nH_2O + MOH \cdot nH_2\bar{O} = MX \cdot (2n+1)H_2O$, ΔH_N^o is equal to $\Delta H_N - [\Phi_L \mathbf{M} \mathbf{X} \cdot (2n+1) \mathbf{H}_2 \mathbf{O} - \Phi_L \mathbf{M} \mathbf{O} \mathbf{H} \cdot n \mathbf{H}_2 \mathbf{O} - \Phi_L \mathbf{H} \mathbf{X} \cdot n \mathbf{H}_2 \mathbf{O}]$.

 ΔH_i is the heat of ionization or dissociation of a weak acid or base into its ionic species, such as, $HA + H_2O = H_3O^+ + A^-$ and $RNH_2 + H_2O - RNH_3^+$

+ OH-.

 K_i refers to the equilibrium constant of the above type reactions, with K_a and K_b referring specifically to the equilibrium constants of the acid and base

reactions, respectively.

The properties were calculated, wherever possible, over the entire range from infinite dilution to approximate saturation or to a 1:1 mole ratio. Use was made of Seidell's and Linke's compilations [615, 616, 617] to obtain the saturation concentrations. Values for Φ_L and Φ_C were estimated for some electrolytes when there was enough information available on the other members of the family.

¹ Figures in brackets indicats the literature references on pages 34-41.

II. Units of Energy and Constants

The unit of energy used is the thermochemical calorie, defined by the relation 1 thermochemical calorie = 4.1840 J. The 1961 atomic weights are used [16]. The values for the specific heat of water are those recommended by the International Committee of Weights and Measures as given by Stimson [17]. The conversion factors for correction to the thermochemical calorie are taken from [18] and [618]. Unless otherwise mentioned the specific heats of the pure solutes used to correct $\Delta H_{\rm soln}$ to 25 °C have been obtained from Kelley [19] and Kelley and King [20]. The values of the limiting slopes appearing in the Debye-Hückel theory of strong electrolytes are taken from Harned and Owen [9]. The emf data, on the whole, were used as summarized by Harned and Owen. The heats of vaporization of H₂O used in the calculations on the dissociation pressures of the hydrates were obtained from Osborne, Stimson, and Ginnings [649]. Whenever sufficient data appeared in the original article, the results were completely recalculated, making appropriate corrections for the specific heat of water, atomic weights, calorie used, etc. When the details given were insufficient to make exact corrections, the experimental results were corrected on the basis of reasonable estimates of the quantities employed by the authors.

III. Uncertainties

The uncertainties assigned to the various values have been based on many factors - the experimental technique used, the details given, the number of measurements, the standard deviation, the magnitude of the corrections to 25 °C (and to infinite dilution, for ΔH_{ω}^{0} , the inherent error in the methods used, the reliability of previous work of the investigators, etc. A strictly mathematical evaluation can therefore not be made. The personal appraisal is an important part of the overall evaluation. For example, the standard deviation of Wüst and Lange's [21] work on KCl is ±1 cal, but the assigned uncertainty is larger; the value selected as the "best" value is 40 cal lower than theirs. As in the above case, the standard deviation and also the assigned uncertainty can be misleading. In other cases it is not possible to assign an uncertainty because of the lack of information.

IV. The Apparent Molal Heat Capacity

A great many investigators have measured the specific heats of aqueous electrolyte solutions although only a few have done so with sufficient precision to give reliable results for the apparent molal heat capacity of the solute. Harned and Owen [8] have pointed out the difficulty in obtaining satisfactory results in very dilute solutions. They show, for example, that at a concentration of 0.01 molal an error of 0.01 percent in the specific heat measurements results in an error of about 10 cal/deg mole

in the apparent molal heat capacity of the solute, whereas an error of 1 percent in the determination of the concentration results in an error of only 0.5 cal/deg mole. For this reason a great deal of reliance has been placed on the Debye-Hückel theory in the very dilute region.

The measurements of specific heat were converted to Φ_C . The values for Φ_C were plotted against $m^{1/2}$ on large scale graphs. The first three systems treated were NaCl, HCl, and NaOH, at the various experimental temperatures. This was done to determine the temperature coefficients, which in general were applied to other systems when necessary to correct to 25 °C. In general the curve was drawn down to $m^{1/2} = 0.4$ following the data available as closely as possible. The curve was continued so that as $m^{1/2} \rightarrow 0$, the slope, $d_{\Phi_C}/dm^{1/2}$, approached the limiting value (6.13 at 25 °C).

Preliminary values for $\Phi_{\mathcal{C}}^{\circ}$ at 25 °C for the hydroxides, chlorides, bromides, iodides, and nitrates of H, Li, Na, and K were obtained. It was assumed that the heat capacities of the individual ionic species are additive at infinite dilution. Equations of the following sort were set up:

```
\begin{split} &\Phi_{\mathcal{C}}^{\circ}(H^{+}) + \Phi_{\mathcal{C}}^{\circ}(Cl^{-}) = \Phi_{\mathcal{C}}^{\circ}(HCl) \\ &\Phi_{\mathcal{C}}^{\circ}(Na^{+}) + \Phi_{\mathcal{C}}^{\circ}(Cl^{-}) = \Phi_{\mathcal{C}}^{\circ}(NaCl) \\ &\Phi_{\mathcal{C}}^{\circ}(K^{+}) + \Phi_{\mathcal{C}}^{\circ}(Cl^{-}) = \Phi_{\mathcal{C}}^{\circ}(KCl) \\ &\Phi_{\mathcal{C}}^{\circ}(H^{+}) + \Phi_{\mathcal{C}}^{\circ}(Br^{-}) = \Phi_{\mathcal{C}}^{\circ}(HBr) \\ &\Phi_{\mathcal{C}}^{\circ}(Na^{+}) + \Phi_{\mathcal{C}}^{\circ}(Br^{-}) = \Phi_{\mathcal{C}}^{\circ}(NaBr) \\ &\Phi_{\mathcal{C}}^{\circ}(K^{+}) + \Phi_{\mathcal{C}}^{\circ}(NO_{3}^{-}) = \Phi_{\mathcal{C}}^{\circ}(HNO_{3}) \\ &\Phi_{\mathcal{C}}^{\circ}(Na^{+}) + \Phi_{\mathcal{C}}^{\circ}(NO_{3}^{-}) = \Phi_{\mathcal{C}}^{\circ}(NaNO_{3}), \end{split}
```

 $\Phi_C^{\circ}(H^+)$ was set equal to 0. The values of Φ_C° for each electrolyte were weighted proportionally to their estimated reliability. By cross substituting values, values for the individual ions were obtained. The various values for each ion were weighted, depending on the source, and averaged. The Φ_C° values of the electrolytes were then adjusted to be self-consistent.

The values obtained were checked by the method of Guggenheim and Prue [22] as adapted by Pitzer and Brewer [10]. The results agreed well within the uncertainties. Thus, Pitzer and Brewer's values of Φ_c° for NaCl (-22.1 cal/deg mole) and KCl (-27.7) compare favorably with -21.5 and -27.4, respectively, obtained here by the above simple approach. If their method is used with the experimental data used here, values of -21.9 and -27.8 are obtained. The difference between $\Phi_c^{\circ}(Na^+)$ and $\Phi_c^{\circ}(K^+)$ in this work is 5.9 as compared to 5.6; for Cl⁻ and Br⁻, 1.3 as compared to 1.2. It was therefore felt that there was no appreciable difference in results due to the method of extrapolation, but rather that it was which data were used and how the experimental data were treated that was more important. It was also felt advantageous

to treat each system individually. Very few adjustments were found to be necessary to make the results consistent, not only for Φ_C° , but also for Φ_L , $\Delta H_{\infty}^{\circ}$, and ΔH_{N}° , at 25 °C.

Once the values for the above mentioned ions were determined, they were subtracted from the Φ_C° values obtained later for other compounds to obtain the values for the other ions in table I.

The values of Φ_C° of HCl, NaOH, and NaCI (table II) at temperatures other than 25 °C were not adjusted, since there were insufficient data available on the other systems to give an adequate number of values of Φ_C° for smoothing.

The values of $\Phi_{\mathcal{C}}^{\circ}$ for all compounds considered are given in tables III to IX. Graphic representation of the various families of curves, as well as a detailed curve for KCl, are given in figures I through VIII.

The available data for each compound are summarized in the following section. For each substance are listed the various investigations, with the temperature and range of concentrations measured. The investigations given the greatest weight in the evaluation of Φ_C are marked with an asterisk. The substances are discussed in the order: acids, ammonium and amine salts, silver salts, and salts of the alkali metals.

HF

The specific heat has been measured by Baud [23] (15 °C, m=11.72), Pranschke and Schwiete [24]* (18 °C, m=34.30 to 5.44), Kolesov, Skuratov, and Uvarov [25]* (25 °C, m=2.01 to 0.01), Thorvaldson and Bailey [26]* (19 °C, m=46.37 to 0.28), Roth and Chall [27] (50 °C, and between 20 and 50 °C, m=13.26), Roth, Pahlke, Bertram, and Börger [28]* (20 °C, m=2.22 to 0.11), and Mulert [29] (17–18 °C, m=17.5 to 2.63).

HCl

The specific heat has been measured by Thomsen [30]* (18 °C, m=5.55 to 0.56), Marignae [31, 32]* (22 °C, m = 2.22 to 0.56), Richards and Rowe [33, 34] (18 °C, m = 0.28), [35, 36] (18 °C, m = 0.56 and 0.28), and [37]* (18 °C, m = 5.55 and 2.77), Richards, Rowe, and Burgess [38] (20 °C, m=6.30 and 2.78), Roth and Chall [27] (50 °C, and between 22.5 and 50 °C, m=2.25), Ackermann [39]* (10, 20, 40 to 120 °C, m=2.00 to 0.50), Vrevskii and Kaigorodov $[40, 41]^*$ (3.3, 20.5, 40.4, and 60.5 °C, m = 16.60 to 0.17), Tucker [42] (10 °C, m = 12.67 to 2.57), Richards, Mair, and Hall [43]* (16, 18, and 20 °C, m = 2.22to 0.56), Thorvaldson, Brown, and Peaker [44]* (18 and 19 °C, m = 2.78), Drucker [45]* (18 °C, m = 2.65 and 1.25), Radulescu and Jula [46]* (15 °C, m = 2.08), Wicke, Eigen, and Ackermann [47]* (20, 40, 60 to 130 °C, m = 1.91 to 0.59), Randall and Ramage [48]* (25 °C, m=1.05 to 0.08), Gucker and Schminke [49]* (25 °C, m = 2.01 to 0.01), and Sturtevant [50]* (25 °C, m = 0.87 to 0.02). The \overline{J}_2 values obtained here agree well with Bates and Bower's [51] values of \overline{J}_2 from emf measurements below m=0.1 and

Harned and Owen's [9] tabulation of \overline{J}_2 obtained from the extended Debye-Hückel equation. For $0.1 \le m \le 4.0$ the \overline{J}_2 values are below and within 2 cal/deg mole of the values tabulated by Harned and Owen from Harned and Ehlers' emf data. As $m \to 4.0$ the difference becomes 1 cal/deg mole.

Other emf measurements since 1955 have been reported by Hayes and Lietzke [52] (25 to 50 °C, m=0.1 to 0.001), Greeley [53] (25 to 200 °C, m=1.0 to 0.005), Covington and Prue [54] (25 °C, m=0.07 to 0.005) and [55] (25 °C, m=0.1 to 0.005), Oiwa [56] (25 °C, m=0.1 to 0.001), Taylor and Smith [57] (10 to 45 °C, m=1.5 to 0.9) and Aston and Gittler [58] (25 °C, m=4.0 to 1.2).

HClO₄

The specific heat has been measured by Berthelot [59, 60] (between 15 and 40 °C, m=9.0 to 0.05), and Richards and Rowe [36] (18 °C, m=0.56).

The data were not considered reliable for a Φ_C curve.

\mathbf{HBr}

The specific heat has been measured by Tolloczko and Meyer [61] (13.25 °C, m=4.1), Richards and Rowe [35, 36]* (18 °C, m=0.56), Roozeboom [62]* (between 12 and 30 °C, m=30.17 to 0.28), and Johnson, Gilliland, and Prosen [63]* (25 °C, m=11.10)

 J_2 values were compared with Harned and Owen's [9], and Biermann and Yamasaki's [64] emf data. The agreement is good.

HI

The specific heat has been measured by Richards and Rowe [36]* (18 °C, m=0.56), and Johnson, Gilliland, and Prosen [63]* (25 °C, m=7.60).

HIO.

The specific heat has been measured by Randall and Taylor [65]* (25 °C, m = 3.60 to 0.07).

HNO

The specific heat has been measured by Thomsen [30]* (18 °C, m=5.55 to 0.23), Marignac [66]* (between 21 and 52 °C, m=1.10 to 0.56), Mishchenko and Ponomareva [67]* (21 °C, extremely concentrated solutions to 1.24), Forsythe and Giauque [68]* (25 °C, m=35.55 and 18.50), Roth [69]* (20 °C, m=0.27 to 0.03), Drucker [45]* (18 °C, m=2.41 to 1.06), Richards and Rowe [36] (18 °C, m=0.56) and [70]* (18 °C, m=5.55 to 0.14), Lloyd and Wyatt [71] (0 °C, extremely concentrated solutions), Bump et al. [72] (0 to 26.7 °C, extremely concentrated solutions), and Booman, Elverum, and Mason [73] (0 °C, extremely concentrated solutions). $\mathbf{CH_2O_2}$

The specific heat of aqueous formic acid has been measured by Bury and Davies [74]* (15 °C, m=10.06 to 1.33), Ackermann and Schreiner [75]* (0, 10, 20, 30 to 130 °C, m=2.84 to 0.49), and Glagoleva and Cherbov [76]* (25, 60, and 80 °C, 100% to m=4.185).

The specific heat of CH₂O₂(1) has been measured by von Reis [77], Radulescu and Jula [46], Berthelot and Ogier [78], Pettersson [79], Glagoleva and Cherbov [76], Gibson, Latimer, and Parks [80]*, and Stout and Fisher [81]*.

$C_2H_4O_2$

The specific heat of aqueous acetic acid has been measured by Marignac [32, 66]* (between 21 and 52 °C, m=22.2 to 2.22), Neiman [82, 83]* (23.9, 42.0, 59.5, and 80.5 °C, 100% to m=0.96), Richards and Gucker [84]* (16, 18, and 20 °C, m=2.22), von Reis [85]* (18 °C, m=2.22), Bury and Davies [74]* (15 °C, m=8.66 to 0.91), and Ackermann and Schreiner [75]* (10, 20, 30 to 130 °C, m=3.00 to 0.50).

Richards and Gucker [84]* calculated the specific heat (18 °C, m=1.11 to 0.07) from $\Delta H_{\rm diln}$ and the value of the specific heat at m=2.22.

The specific heat of C₂H₄O₂(1) has been measured by Radulescu and Jula [46]*, von Reis [77], Parks and Kelley [86]*, Swietoslawski and Zielenkiewicz [87], Neiman [82, 83]*, Marignac [32], and Berthelot [88].

$C_3H_6O_2$

The specific heat of aqueous propionic acid has been measured by Bury and Davies [74]* (15 °C, m=5.84 to m=0.49) and Ackermann and Schreiner [75]* (0, 10, 20, 30 to 130 °C, m=2.00 to 0.50).

NH_3

The specific heat has been measured by Rühlemann [89] (between 22 and 99 °C, m=1.35), Vrevskiĭ and Kaĭgorodov [40, 41]* (2.4, 20.6, 41, and 60.9 °C, m=28.01 to 0.84), and Hildenbrand and Giauque [90]* (25 °C, m=55.51, and -73 to -18 °C, more concentrated solutions).

NH₄F

The specific heat has been measured by Labowitz and Westrum [91]* (25 °C, m = 0.50 to 0.005).

NH₄Cl

The specific heat has been measured by Thomsen [30]* (18 °C, m=7.40 to 0.28), Marignac [32]* (hetween 20 and 52 °C, m=4.44 to 0.56), Drucker [45]* (18 °C, m=4.73 to 1.10), Mishchenko and Ponomareva [67, 92]* (25 °C, m=7.03 to 0.31), D'Ans and Tollert [93]* (21, 41, 70, and 86 °C, m=3.28 and 1.67), Winkelmann [94]* (15 to 28.6 °C, m=6.23 to 0.58), Faasch [95]* (18 °C, m=3.96 to 0.47), and Urban [96]* (20, 25, 30, 35, and 40 °C, m=2.00).

The value for $\Phi_c^\circ NH_4^+$ was obtained by subtracting $\Phi_c^\circ Cl^-$ from $\Phi_c^\circ NH_4 Cl$.

NH₄Br

The specific heat has been measured by Thomsen [30] (19 °C, m=0.28) and Faasch [95] (18 °C, m=4.02 to 0.50). The Φ_C line, however, was estimated. $\Phi_C^\circ NH_4 Br$ was obtained from $\Phi_C^\circ NH_4^+$ and $\Phi_C^\circ Br^-$

NH₄I

Thomsen [30] measured the specific heat (18 °C,

m = 0.28). The Φ_C curve was essentially estimated, using $\Phi_C^{\circ} NH_4I$ from the ions.

NH₄NO₃

The specific heat has been measured by Thomsen [30]* (18 °C, m=11.10 to 0. 56), Marignac [32, 66]* (between 20 and 50 °C, m=11.10 to 0.56), Tollinger [97]* (23.5 °C, m=0.56). Zwicky [98] (20 °C, m=28.61 to 1.39), Lerner-Steinberg [99]* (21.6 °C, m=0.60 and 0.44), Chauvenet [100] (15 °C, m=1.33), Drucker [45]* (20 °C, m=4.99 and 1.05), Rutskov [101, 102]* (25, 50, and 75 °C, m=18.50 to 0.56), Winkelmann [94]* (16 to 38 °C, m=8.33 to 0.39), Cohen, Helderman, and Moesveld [103, 104] (32.3 °C, m=28.41 to 1.39), and Gucker, Ayres, and Rubin [105, 106]* (25 °C, m=24.00 to 0.10).

CH₃NH₃Cl

The specific heat has been measured by Whitlow and Felsing [107]* (25 °C, m=0.96 to 0.09). Φ_C is tabulated only in the range from m=1.11 to 0.09 since the data available are not considered reliable for extrapolation to infinite dilution. A rough extrapolation leads to a value for Φ_C° of -1 cal/deg mole.

(CH₃)₂NH₂Cl

The specific heat has been measured by Whitlow and Felsing [107]* (25 °C, m=0.98 to 0.11). Φ_C is tabulated only in the range from m=1.11 to 0.09 since the data available are not considered reliable for extrapolation to infinite dilution. A rough extrapolation leads to a value for Φ_C° of +2 cal/deg mole.

(CH₃)₃NHCl

The specific heat has been measured by Whitlow and Felsing [107]* (25 °C, m=1.0 to 0.09). Φ_C is tabulated only in the range from m=1.11 to 0.09 since the data available are not considered reliable for extrapolation to infinite dilution. A rough extrapolation leads to a value for Φ_C° of +4 cal/deg mole.

AgF

The specific heat has been measured by Jahn-Held and Jellinek [110]* (25 °C, m=14 to 1). Values for Φ_C are given at infinite dilution and above m=2.

AgNO₃

The specific heat has been measured by Marignac [66]* (between 25 and 52 °C, m=2.22 to 0.56) and Kapustinskii, Yakushevskii, and Drakin [111]* (25 °C, m=1.80 to 0.28). Roth [69]* gives an equation for the range m=0.22 to 0.08 at 18.5 °C.

LiOH

The specific heat has been measured by Richards and Rowe [70]* (18 °C, m=2.22), and [36] (18 °C, m=0.56), and Gucker and Schminke [112]* (25 °C, m=2.27 to 0.04).

Richards and Rowe [70]* calculated the specific heat (18 °C, $m\!=\!1.11$ to 0.14) from $\Delta H_{\rm diln}$ at 16 and 20 °C and the specific heat at $m\!=\!2.22$.

The Φ_C curve was based primarily upon Gucker and Schminke's measurements.

LiC

The specific heat has been measured by Tucker [42] (14 to 22 °C, m=16.34 to 2.52), Chernyak [113] (25 °C, m=1.0), Bennewitz and Kratz [114]* (20 °C, m=1.0 to 0.05), White [115]* (25 °C, m=0.05), Jauch [116] (18 °C, m=4.06 to 0.50) Hess [117]* (25 °C, m=0.15), Drucker [45]* (18 °C, m=9.8 to 1.3), Gucker and Schminke [49]* (25 °C, m=2.46 to 0.04), Richards and Rowe [70]* (18 °C, m=2.22), and Lange and Dürr [118]* (25 to 28 °C, m=19.89 to 0.56).

Richards and Rowe [70]* calculated the specific heat (18 °C, m=1.11 to 0.14) from $\Delta H_{\rm diln}$ and the specific heat at m=2.22.

The Φ_C lines of Gucker and Schminke, and Lange and Dürr are parallel to one another. There seems to be a systematic error in the latter's work. The final Φ_C line is based primarily on Gucker and Schminke's work.

LiBr

The specific heat has been measured by Jauch [116]* (18 °C, m=4.45 to 0.51), Lange and Schwartz [119]* (26.5 °C, m=17.92 to 0.04), and Eigen and Wicke [120]* (10, 20, 40, 60 to 120 °C, m=0.56, 0.91, and 1.32).

Here as in LiCl, the Φ_C lines of Eigen and Wicke and Lange and Schwartz are parallel. Eigen and Wicke's line was essentially used, with Lange and Schwartz's Φ_C values lowered by a constant amount to coincide with Eigen and Wicke's.

Lil

The specific heat has been measured by Jauch [116]* (18 °C, m=3.17 to 0.51).

LiNO₃

The specific heat has been measured by Chauvenet [100] (15 °C, m=2), Drucker [45]* (18 °C, m=1.89 to 0.91), and Richards and Rowe [70]* (18 °C, m=2.22). The latter also used $\Delta H_{\rm diln}$ at 16 and 20 °C to calculate the specific heat from m=1.11 to 0.14 at 18 °C.

NaOH

The specific heat has been measured by Thomsen [30]* (19 °C, m=7.40 to 0.28), Bertetti and McCabe [121]* (3.12 to 60 °C, m=22.81 to 1.06), Richards and Rowe [36]* (18.1 °C, m=0.56 and 0.28), Ackermann [39, 122]* (10, 20, 40, 60 to 130 °C, m=2.00 to 0.50), Richards and Gucker [84]* (16, 18, and 20 °C, m=2.22), Sturtevant [50]* (25 °C, m=0.087 to 0.006), Tucker [42] (16 to 21 °C, m=22.54 to 4.92), Richards and Hall [123]* (18 °C, m=2.22 to 0.56), Roth, Wirths, and Berendt [124] (19.8 °C, m=10.11), Wilson and McCabe [125]* (29.8 to 122 °C, m=26.76 to 25.20), and Gucker and Schminke [112]* (25 °C, m=2.55 to 0.04).

The values of \bar{J}_2 obtained here agree well with those given by Harned and Owen [9] and Akerlöf and Kegeles [126] below m=2.

NaCl

The specific heat has been measured by Thomsen [30] (19 °C, m=5.55 to 0.28), Marignac [32] (18 °C and between 22 and 52 °C, m=2.22 to 0.56), Stakhanova and Vasilev [127]* (25 °C, m=4.37 to 0.20), Biron [128] (19 °C, m=0.28 to 0.07), Blaszkowska [129]* (21.4 °C, m=2.22 to 0.28), Zdanovskii [130]* (25 °C, m=2.33), Zdanovskii and Suslina [131]* (25 °C, m=1.89), Zdanovskii and Matsenok [132]* (25 °C, m=6.11 to 1.75), Eigen and Wicke [120]* (10, 20, 40 to 130 °C, m=1.12 to 0.41), D'Ans and Tollert [93] (21 and 41 °C, m=5.75 to 1.66), Drucker [45]* (20 °C, m=4.36 to 1.02), Chipman, Johnson, and Maass [133]* (18.8 and 25.4 °C, m=2.54 to 0.43), Nikolaev, Kogan, and Ogorodnikov [134]* (25 °C, m=6.15 to 0.34), Winkelmann [94] (17 to 46 °C, m=5.48 to 0.53), Leitner [135] (25.7 to 95.7 °C, m=5.97), Hess [117]* (25 °C, m=0.15 to 0.003), White [136]* (35 to 45 °C, m=0.20 to 0.01), Hess and Gramkee [137]* (15, 25, 35, 45 °C, m=1.03 to 0.01), Voskresenskaya and Yankovskaya [138] (50 °C, m=57.6 to 3.45), Lipsett, Johnson, and Maass [139]* (25 °C, m=6.01 to 0.17), Bousefield and Bousefield [140]* (7, 20, and 33 °C, m=5.70 to 0.25), Rutskov [101, 102]* (25, 50, and 75 °C, m=5.55 to 0.20), Ackermann [39]* (10, 20, 40, 60 to 130 °C, m=2.00 to 0.50), Randall and Ramage [48]* (25 °C, m=1.90 to 0.07), Sandonnini and Gerosa [141] (18.5 °C, m=6.20 and 4.04), Randall and Rossini [144]* (25 °C, m=6.20 and 4.04), Randall and Rossini [144]* (25 °C, m=6.20 and 4.04), Randall and Rossini [144]* (25 °C, m=2.30 to 0.04), and Richards and Gucker [84]* (18 °C, m=2.22).

The J_2 values were checked with those obtained from emf data as given by Harned and Owen [9]. The agreement is excellent below m=1 and within 3 cal/deg mole between m=1 and 4.

NaClO₃

The specific heat has been measured by Colomina and Nicolas [145]* (24 °C, m=15.39 to 0.20) and Ferrer [146]* (25 °C, m=3.11 to 0.19).

NaClO₄

The specific heat has been measured by Colomina and Nicolas [145]* (24.5 °C, m=12.22 to 0.13), and Ferrer [146]* (25 °C, m=2.59 to 0.24).

The values of Φ_C chosen were based on Ferrer's data up to m=2.6 and above that on Colomina's.

NaBr

The specific heat of aqueous NaBr has been measured by Marignac [32, 66]* (between 20 and 52 °C, m=2.22 to 0.56), Chipman, Johnson, and Maass [133] (25.4 and 18.8 °C, m=1.88 to 0.24), Bender and Kaiser [147]* (25 and 30 °C, m=8.37 to 0.97), Rossini [148]* (25 °C, m=0.43 to 0.08), and Randall and Rossini [144]* (25 °C, smoothed values from m=1.00 to 0.01).

Below m=2.0 the values obtained here for \overline{J}_2 were below but within 3 cal/deg mole of the emf values listed in Harned and Owen [9].

NaBrO₃

The specific heat has been measured by Ferrer [146]* (25 °C, m = 2.34 to 0.21).

NaI

The specific heat has been measured by Thomsen [30]* (18 °C, m=0.28), Marignac [32, 66]* (between 20 and 51 °C, m=2.22 to 0.56), Rossini [148]* (25 °C, m=0.85 to 0.9), and Randall and Rossini [144]* (25 °C, smoothed values from m=1.00 to 0.01).

NaNO₂

The specific heat has been measured by Ferrer [146]* (25 °C, m=2.25 to 0.10) and Perreu [149] (15 °C, m=11.87 to 1.45).

NaNO₃

The specific heat has been measured by Thomsen [30]* (18 °C, m=5.55 to 0.28), Marignac [66]* (20.5 °C and between 22 and 52 °C, m=2.22 to 0.56), Biron [128] (19 °C, m=0.28 to 0.03), Person [142] (0 to 20 °C, m=7.7 to 0.59), Magie [150] (room temperature, m=1.11 to 0.19), D'Ans and Tollert [93]* (21 °C, m=5.0), Winkelmann [94, 151] (2 to 60 °C, m=2.35 to 0.37), Gerlach [152] (between 20 and 95 °C, m=10.2), Richards and Gucker [84]* (18 °C, m=2.22), Zdanovskiĭ [130]* (25 °C, m=2.58), Rossini [148]* (25 °C, m=0.85 to 0.08), Randall and Rossini [144]* (25 °C, smoothed values at m=1.00 to 0.01), Richards and Rowe [70]* (18 °C, m=2.22), Drucker [45]* (18 °C, m=4.59 to 1.03), and Chauvenet [100] (15 °C, m=2.13).

Richards and Rowe [70]* calculated the specific heat (18 °C, m=1.11 to 0.14) from the dilution data at 16 and 20 °C and the specific heat at m=2.22.

NaCHO₂

The specific heat of aqueous sodium formate has been measured by Ackermann and Schreiner [75]* (10, 20, 30 to 120 °C, m=2.85 to 0.50).

NaC₂H₃O₂

The specific heat of aqueous sodium acetate has been measured by Thomsen [30]* (17 °C, m=2.78 to 0.28), Marignac [32, 66]* (22.5 °C, and between 19 and 52 °C, m=2.22 to 0.56), Richards and Gucker [84]* (16, 18, and 20 °C, m=2.22), Perreu [153]* (19 °C, m=4.78 to 1.36), Bindel [154] (21 °C, m=5.55 to 1.11), Ferrer [146]* (25 °C, m=2.79 to 0.23), Kapustinskii and Stakhanova [155]* (25 °C, m=17.13 to 3.14), Ackermann and Schreiner [75]* (10, 20, 30 to 120 °C, m=2.53 to 0.60). Gnesotto and Fabris [156] measured the specific heat of the liquid trihydrate at 0 to 100 °C.

Richards and Gucker [157]* calculated the specific heat (m=1.09 to 0.03) from dilution data and the specific heat at m=2.22, obtained in [84]. $NaC_0H_0O_0$

The specific heat of aqueous sodium propionate

has been measured by Ackermann and Schreiner [75]* (10, 20, 30 to 120 °C, m = 2.24 to 0.50).

KOH

The specific heat has been measured by Thomsen [30]* (18 °C, m=1.85 to 0.28), Jaquerod [158] (18 °C, m=4.92 to 0.95), Gawlik [159]* (20 °C, m=0.33 to 0.04), Voskresenskaya [160]* (20 °C, m=4.24), Richards and Hall [123]* (18 °C, m=0.56), Gucker and Schminke [49]* (25 °C, m=2.59 to 0.04), Richards and Rowe [36]* (18 °C, m=0.56) and [70]* (18 °C, m=13.38 and 5.55), Sabatier [161] (temperature uncertain, m=13.37).

Richards and Rowe [70]* calculated the specific heat (18 °C, m=2.22 to 0.14) from the dilution data and specific heats of the more concentrated solutions.

The agreement of the calculated \bar{J}_2 values with those given by Harned and Owen [9] for the emf values is excellent.

KCI

The specific heat has been measured by Thomsen [30]* (18 to 20 °C, m=3.7 to 0.28), Marignac [32, 66]* (19.5 °C and between 20 and 51 °C, m=1.11 to 0.56), Andrews [162] (18 °C, m = 0.25), Stakhanova and Vasilev [127]* (25 °C, m = 4.60 to 0.20), Voskresenskaya and Rutskov [163]* (25 °C, m=1.40), Biron [128] (19 °C, m=0.28 to 0.03), Winkelmann [94] (0 to 55 °C, m = 5.59 to 0.42), and [151] (20 °C, m=0.71), Chipman, Johnson, and Maass [133]* (18.8 and 25.4 °C, m=2.48 to 0.33), Magie [150] (room temperature, m=1.11 to 0.14), Sandonnini and Gerosa [141] (18.5 °C, m=0.75 to 0.17), Jaquerod [158] (18 °C, m=3.8 to 0.95), Cobb and Gilbert [164]* (25 °C, m=1.00 and 0.50), D'Ans and Tollert [93]* (21 and 41 °C, m = 4.40 to 1.7), Nikolaev, Kogan, and Ogorodnikov [134]* (25 °C, m=4.80 to 0.07), Zdanovskii [130]* (25 °C, m=2.56), Rutskov [101, 102]* (25 °C, m=4.63 to 0.10), Popov, Bundel, and Choller [165]* (20 °C, m = 0.56 to 0.14), Popov, Khomyakov, Feodos'ev, and Schirokich [166]* (same data as [165]), Urban [96]* (10 to 40 °C, m = 2.00 to 0.10), Ferrer [146]* (25 °C, m = 2.98 to 0.05), Hess and Gramkee [137]* (15, 25, 35, 45 °C, m = 1.03 to 0.01), Clews [167]* (15.6 to 43.2 °C, m=2.50 to 0.10), Gucker [168]* (20 °C, m = 2.22), Kapustinskiĭ, Yakushevskiĭ, and Drakin [111]* (25 °C, m = 1.71 to 0.52), Varasova, Mishchenko, and Frost [169]* (25 °C, m = 4.85 and 2.73), Randall and Bisson [143] (25 °C, m = 4.85and 2.73), Bousefield and Bousefield [140]* (20 and 33 °C, m = 3.40 to 0.25), Randall and Rossini [144]* (25 °C, m = 2.50 to 0.05), Rossini [148]* (25 °C, m = 2.25 to 0.04), Cohen and Moesvell [170]* (19 °C, m = 3.39 and 1.03), Parshikov [171]* (25 °C, m = 0.41), and Richards and Rowe [70]* $(18 \, ^{\circ}\text{C}, m = 2.22)$

Richards and Rowe [70]* calculated the specific heat (18 °C, m=1.11 to 0.14) from the directly determined specific heat at m=2.22 and their dilution data. The $\Delta H_{\rm soln}$ and $\Delta H_{\rm diin}$ experiments of Wüst and Lange [21]* were sufficiently detailed

to permit the computation of the heat capacity of the solutions. The $\Delta H_{\rm soln}$ measurements of KCl at various temperatures were also used to compute Φ_C at the mean temperature.

The calculated $\overline{J_2}$ values were compared with those obtained from the emf measurements as given by Harned and Owen [9]. The agreement was good, within 1 cal/deg mole below m=2.

KBr

The specific heat has been measured by Thomsen [30]* (18 °C, m = 0.28), Marignac [32, 66]* (between 20 and 51 °C, m = 2.22 to 0.56), Popov, Bundel, and Choller [165]* (20 °C, m = 0.19), Bender and Kaiser [147]* (25 and 35 °C, m = 5.52 to 1.00), Rossini [148]* (25 °C, m = 0.97 to 0.04), Randall and Rossini [144]* (25 °C, smoothed values from m = 1.00 to 0.01), Chipman, Johnson, and Maass [133]* (25.4 and 18.4 °C, m = 1.57 to 0.21), Urban [96]* (10, 15, 20, 25, 30, 35, and 40 °C, m = 2.00), Popov, Skuratov, and Strel'tsova [172]* (20.2 °C, m = 0.32).

KI

The specific heat has been measured by Thomsen [30]* (18 °C, m=0.28), Marignac [32] (between 20 and 51 °C, m=2.22 to 0.56), Drucker [45]* (18 °C, m=3.24 to 0.68), Popov, Skuratov, and Strel'tsova [172]* (20.2 °C, m=1.63), Rossini [148]* (25 °C, m=0.85 to 0.04), and Randall and Rossini [144] (25 °C, smoothed values from m=1.00 to 0.01).

KIO₃

The specific heat has been measured by Randall and Taylor [65]* (25 °C, m = 0.38 to 0.04).

KNO₃

The specific heat has been measured by Thomsen [30]* (18 °C, m=2.22 to 0.23), Marignac [32, 66]* (20.5 °C and between 22 and 52 °C, m=2.22 to 0.56), Person [142] (0.6 to 20.6 °C, m=2.00 to 0.50), Drucker [45]* (19 °C, m=2.18 and 1.05), Voskresenskaya [160]* (30 °C, m=2.10), Rutskov [101, 102]* (25, 50, and 75 °C, m=0.50 to 6.94), Zdanovskii [130]* (25 °C, m=3.12), Magie [150] (room temperature, m=2.22 to 0.14), Biron [128] (19 °C, m=0.28 and 0.14), D'Ans and Tollert [93] (41 °C, m=2.78), Winkelmann [94, 151] (4 to 60 °C, m=2.44 to 0.30), Chauvenet [100] (15 °C, m=2.22), Andrews [162] (16.6 °C, m=0.26), Gucker [168]* (20, 50, and 80 °C, m=2.22), Richards and Rowe [70]* (18 °C, m=2.22), Rossini [148]* (25 °C, m=1.05 to 0.05), Randall and Rossini [144]* (25 °C, smoothed values at m=1.00 to 0.01).

Lange and Monheim [173]* calculated the specific heat at 18.75 °C from the solution and dilution experiments at 12.5 and 25.0 °C at m=0.146 to 0.0004. Richards and Rowe [70]* calculated the specific heat (18 °C, m=1.11 to 0.14) from dilution data and the measured specific heat at m=2.22.

KCNS

Urban [96]* has measured the specific heat (10, 15, 20, 25, 30, 35, and 40 °C, m=2.63 to 0.50).

$KC_2H_3O_2$

The specific heat of aqueous potassium acetate has been measured by Marignac [32, 66]* (between 20 and 51 °C, m=11.11 to 2.22), Drucker [45]* (18.6 °C, m=5.62 to 1.05), Urban [96]* (10, 15, 20, 25, 30, 35, and 40 °C, m=2.36 to 0.25), and Andrews [162] (18 °C, m=0.25).

KMnO₄

The specific heat has been measured by Perreu [174]* (19 °C, m=0.40 to 0.063) and Kapustinskiĭ and Klokman [175]* (25 °C, m=0.32 to 0.03).

KClO:

The specific heat has been measured by Bindel [154] (20 °C, m=2.78 to 0.56). The data were not considered reliable for a Φ_C line.

Cs and Rb Compounds

The scarcity of concordant information on the Cs compounds makes the selection of Φ_C^* Cs⁺ difficult. The selection depends on whether the data of Richards and Rowe [70] at 18 °C on CsCl or CsNO₃, or the data of Kapustinskii and coworkers [176, 177, 178, 179] on CsI at 25 °C are utilized. It was felt that the more recent measurements on CsI are the more reliable. This choice necessitated estimating the other Φ_C lines, which were drawn to be consistent with the CsI and the K compounds.

Since there was no reliable information available on the Rb compounds to determine $\Phi_c^{\circ}Rb^+$, the value was estimated on the basis of values of Φ_c° of Cs⁺, K⁺, and Na⁺.

The data available for each compound follow:

Rbl

Jauch [116] has measured the specific heat (18 °C, m = 3.14 to 0.05).

RbNO₃

Chauvenet [100] has measured the specific heat $(15 \, ^{\circ}\text{C}, m = 1.45)$.

CsCl

The specific heat has been measured by Bennewitz and Kratz [114]* (20 °C, m=0.41 to 0.01) and Richards and Rowe [70]* (18 °C, m=1.11 to 0.14).

The specific heat has been measured by Kapustinskii, Yakushevskii, and Drakin [176]* (25 °C, m=0.73 to 0.15), and Kapustinskii, Lipilina, and Samoilov [177, 178]* (25 °C, m=1.06 to 0.22). Kapustinskii [179]* reported the same values.

CsNO;

The specific heat has been measured by Chauvenet [100] (15 °C, m=0.61) and Richards and Rowe [70]* (18 °C, m=0.93 to 0.14).

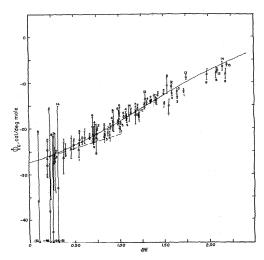


FIGURE 1. The apparent modal heat capacity, Φ_{C_0} of KCl in calldeg mole, at 25 °C as a function of mth. The vertical line indicates the assigned uncertainty of the determination. The series of the determination of the determination of the determination. The series of the determination of the determination of the determination of the determination. The series of the determination of the determinatio

(1) Ferrer [146]
(2) Rossini [148]
(3) Randall and Bisson [143]
(4) Kandall and Bisson [143]
(5) Kandall and Gramkee [137]
(6) Urban [96]
(7) Varasova, Mischenko, and Frost [169]

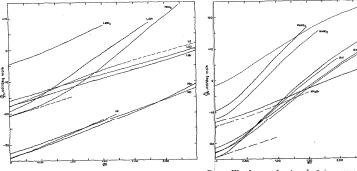


FIGURE III. Φ_{C} as a function of m^{u} in aqueous solutions NH₄Cl, NH₄Br, NH₄NO₃, NaOH, NaCl, NaBr, NaI, NaNC and NaNO₃ at 25 °C. NH₄Br is estimated. Figure II. Φ_C as a function of m^{sc} in aqueous solutions of HCl, HBr, HI, HNO₂, LiOH, LiCl, LiBr, LiI, and LiNO₂ at 25 °C.

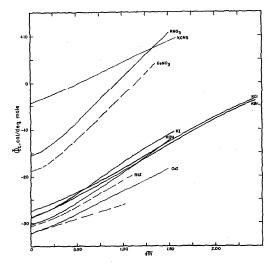
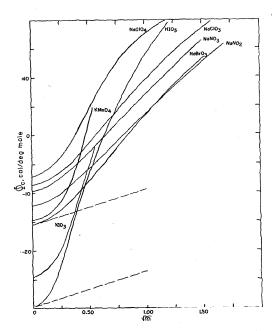


FIGURE IV. Φ_c as a function of mⁿ in aqueous solutions of KOH, KCl, KBr, KI, KNO₃, KCNS, RbI, CsI, and CsNO₃ at 25 °C.
CsNO₃ and RbI are estimated.

FIGURE VI. Φ_c as a function of m^w in aqueous solutions of HCl, NH₄Cl, NaCl, KCl, RbCl, and CsCl at 25 °C.



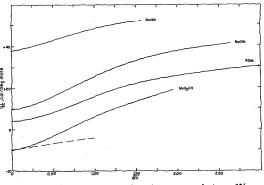


FIGURE VII. P. as a function of mⁿ in aqueous solutions of Na formate, Na acetate, Na propionate, and K acetate at 25 °C.

FIGURE V. Φ_c as a function of mth in aqueous solutions of HIO₃, NaClO₃, NaClO₄, NaBrO₃, NaNO₂, NaNO₃, KIO₃, and KMnO₄ at 25 °C.

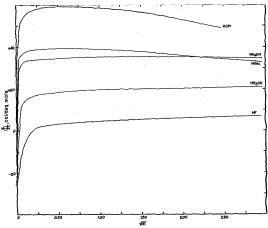


FIGURE VIII. Φ_c as a function of m^* in aqueous solutions of HF, formic acid, acetic acid, propionic acid, and NH₄OH at 25 °C.

V. Heats of Dilution

A great deal of the data from which Φ_L is obtained appears in the literature in the form of measurements of ΔH_{soln} as a function of concentration. By taking the difference between two measured $\Delta H_{\rm soln}$ at concentrations m_1 and m_2 , the $\Delta H_{\rm diln}$ between the two concentrations is obtained. The $\Delta H_{\rm soln}$ data treated in this way can be combined with the directly measured ΔH_{diln} data.

As in the case of the specific heat measurements, the original data were corrected as completely as possible to conform with the present best values of atomic weights, energy units, etc. The more important sets of measurements were completely recalculated from the experimental data whenever the necessary information was available. necessary, the data were corrected to 25 °C using

the values of Φ_C selected earlier.

The method used to obtain the values of Φ_L is the chord-area method developed by Young and coworkers [180, 181, 182, 183] (An alternative method is that of Guggenheim and Prue [184] which leads to essentially the same results.) chords $\Delta\Phi_L/\Delta(m^{1/2})$ were plotted against $m^{1/2}$ on large scale graphs. The extrapolation to infinite dilution was made using 472 cal/mole^{3/2} for the limiting value of $d\Phi_L/dm^{1/2}$. A smooth curve was drawn through all the data, giving greater weight to the more reliable data. The $d\Phi_L/dm^{1/2}$ curves for NaCl and KCl resulting from the graphical method were compared with the curves obtained from least squares treatments of the same data. results agreed well: it was therefore felt unnecessary to apply the least squares treatment to the less extensive data for the other systems. The resultant $d\Phi_L/dm^{1/2}$ curve was integrated to obtain Φ_L . In areas where Φ_L was not well defined by the chord area plot, and long chords were available, Φ_L values were obtained directly by using the experimental ΔH_{diln} and Φ_L for the defined area. In other cases where reliable ΔH_N data were available, Φ_L was obtained directly. All Φ_L values so obtained were consistent with the Φ_L values obtained by the chord area method. There was very little necessity for adjusting values, lending support to the methods used and to the corrections applied. A detailed plot of the chord area treatment of NaCl is shown in figure IX. The values of $d\Phi_L/dm^{1/2}$, Φ_L , and \overline{L}_2 for NaCl are given in table X. Representative graphs of Φ_L for the various families are given in figures X-XIV. The detailed results are tabulated in tables XI-XVI.

The data available for each compound follow:

 ΔH_{diln} has been measured by Petersen [185] (21.5 °C, one dilution from m = 0.20 to 0.10), Roth [186] (20 °C, m = 0.37 to 0.11), Guntz [187] (17 °C, m = 11.012 to 0.14), Schäfer and Kahlenberg [188]* (25 °C, one dilution from m = 34.43 to 2.82) and [189]* (25 °C, one dilution from m=0.33 to 0.03), and Roth, Pahlke, Bertram, and Börger [28]* $(20 \, ^{\circ}\text{C}, m = 24.0 \text{ to } 0.04).$

The values of ΔH_i were obtained from the variation of K_i with temperature for the reactions: (1) $HF = H^+ + F^-$ and (2) $HF + F^- = HF_{\overline{2}}$, using the data of Roth [186], Broene and DeVries [190]*,

and Wooster [191]*.

Other information on the ionization of HF can be obtained from the ΔH_N data of Thomsen [30] (with NaOH), von Steinwehr [192] (unspecified), Petersen [193] (with LiOH), Arrhenius [194] (with NaOH), Guntz [187] (with KOH and NH₃), Mulert [29] (with NaOH and KOH), Roth and Becker [195] (with NaOH), and Berthelot and Guntz [196] (HCl with KF).

The greatest weight was given to Wooster's ata. The selected values at 25 °C are: $\Delta H(1)$ =-3,000 cal/mole and $\Delta H(2)$ =+662 cal/mole.

Elmore, Hatfield, Mason, and Jones [197] reviewed the thermal properties of aqueous HF and, using Wooster's K_i 's, calculated the percentage of each ion in solution at various concentrations of HF. These percentages were used with the above determined ΔH_i 's for reactions (1) and (2) to determine their contribution to ΔH_{soln} and ΔH_{diln} . These heats were combined with the dilution data to obtain Φ_L for the complete range. The Φ_L values as tabulated are to be used only with the ΔH_∞° for complete dissociation into the ions H+ and F-.

 ΔH_{diln} has been measured by Thomsen [30]* (16.5 °C, m = 21.2 to 1.11), Berthelot [198]* (15 °C, m = 25.58 to 0.25), Naude [199]* (12 and 18 °C, m = 1.0 to 0.0001), Tucker [42]* (15 to 20 °C, m = 13.0 to 1.34), Becker and Roth [200]* (20 °C, m = 5.38 to 0.28), Richards, Mair, and Hall [43] (18 °C, m = 2.22 to 0.07), Richards and Rowe [37]* (18 and 20 °C, m = 5.55 to 0.14), von Steinwehr [192] (16 °C, m=4.7 to 0.005), Muller [201] (16, 26, and 38 °C, m=1.0 to 0.01), Thorvaldson, Brown, and Peaker [44]* (20 °C, m=2.78 to 0.28), Petersen [185] (21.5 °C, m=0.56 to 0.19), Richards, Royl, 1012 (21.5 °C, m=0.56 to 0.19), Richards, Royl, 1014 (21.5 °C) Burgess [38] (25 °C, m = 6.30 to 0.28), Roth [69]* (20 °C, m=1.11 to 0.02), Vrevskii and Zavaritskii [202]* (15 °C, m = 15.59 to 1.10), Myers [203]* and Vanderzee and Myers [204]* (25 °C, m=4.49 to 0.07), Sacconi, Paoletti, and Ciampolini [205]* (25 °C, one dilution from m = 1.70 to 0.03), Pitzer [206]* (25 °C, one dilution from m=1.19 to 0.05), Sturtevant [50, 207, 208]* (25 °C, m = 3.34 to 0.005). Ellis [209] has calculated the heat of dilution from emf measurements (25 °C, m = 4.484 to 0.10).

 \bar{L}_1 has been measured by Roth and Wienert [210] (20.7 °C, m=17.28), Schäfer, Wittig, and Jori [211]* (25 °C, m=16.9), and Payn and Perman [212]* (20 and 30 °C, m=16.38 to 1.60).

Bender and Biermann [213]* measured ΔH_N of HCl with NaOH (25 °C, $m_{HCl} = 16.39$ to 3.01).

Vrevskii and Zavaritskii [202, 214] measured ΔH_{soln} as a function of concentration (3.6, 21.5, 42, and 62.3 °C, m = 16.37 to 0.18).

The chord area plot was used to define the curve; chords were obtained from $\Delta H_{\rm diln}$ and $\Delta H_{\rm soln}$. The Φ_L curve which was obtained was adjusted in

the region from m=16.4 to 4.9 to include the Φ_L obtained from the neutralization measurements. The greatest weight below m=5 was given to Sturtevant's corrected data, and above to Bender and Biermann's. Thomsen's and Berthelot's data were used between m=16 and 27. The last two values are extrapolated ones.

The calculated values of \bar{L}_2 agree within 10 cal/mole through m=2.56 with those given by Harned and Owen [9] from emf measurements.

HClO₄

 $\Delta H_{\rm diln}$ has been measured by Berthelot [59, 215, 216, 220] (19 and 21 °C, m=55.5 to 0.28), Krivtsov, Rosolovskii, and Zinov'ev [217]* (25 °C, extremely concentrated solutions to m=0.07), Wood [218] (25 °C, one dilution from m=0.52 to 0.49), Austin and Mair [219] (25 °C, one dilution from m=0.52 to 0.02), and Vanderzee and Swanson [221]* (25 °C, m=10.76 to 0.004).

 Φ_L was based primarily upon Vanderzee and Swanson's measurements. Above m=11, the results of Krivtsov et al., were used.

HRr

 $\Delta H_{\rm diln}$ has been measured by Thomsen [30]* (19 °C, $m\!=\!25.12$ to 0.28), Berthelot [198]* (15.5 °C, $m\!=\!27.14$ to 0.10), Petersen [185] (21.5 °C, $m\!=\!0.56$ to 0.28), Roozeboom [62] (0 and 8.2 °C, $m\!=\!27.8$ to 0.25), and Louguinine and Schukareff [222] (18 °C, $m\!=\!6.49$ to dilute solutions).

Biermann and Weber [223]* measured ΔH_N of HBr with NaOH (25 °C, $m_{\rm HBr}=16$ to 3). Their results were used in this range to obtain Φ_L . Thomsen's data and Berthelot's data were used above and below this range and coordinated with the emf data, as given by Harned and Owen [9], below m=0.4 to complete the Φ_L curve.

HI

Below m=0.56 values of Φ_L were estimated on the basis of a comparison of LiCl-LiBr, NaCl-NaBr-NaI, and HCl-HBr. Above m=0.56 the $\Delta H_{\rm diln}$ measurements of Thomsen [30]* (18.5 °C, m=19.41 to 0.23) and Berthelot [198]* (14 °C, m=18.82 to 0.16) were used.

Richards and Rowe's [224]* ΔH_N (15.5 and 17°C, $m_{\rm HI} = 0.56$) with NaOH and KOH were used as corroborating evidence.

HNO_3

 $\Delta H_{\rm diln}$ has been measured by Thomsen [30]* (18.8 °C, extremely concentrated solutions to $m\!=\!0.28$), Berthelot [225] (9.7 and 26 °C, $m\!=\!2.72$ to 0.69) and [226, 227]* (9.7 °C, extremely concentrated solutions to 0.69), Petersen [185] (21 °C, $m\!=\!0.67$ to 0.17), Richards and Rowe [70]* (20 °C, $m\!=\!5.55$ to 0.14), Naude [199]* (18 °C, $m\!=\!1.03$ to 0.0003), Becker and Roth [228]* (19.8 °C, $m\!=\!1.36$ to 0.01), and Good, Scott, Lacina, and McCullough [229] (25 °C, $m\!=\!1.50$ to 1.29).

 \overline{L}_1 has been measured by Wilson and Miles [230] (20 °C, extremely concentrated solutions), Roth

[231] (20 °C, m=1.52), and Chedin and Dhers [232] (25 °C, m=151.0 to 6.64).

Forsythe and Giauque [68]* measured $\Delta H_{\rm soln}$ of the liquid monohydrate and trihydrate to final molalities of 0.13 and 0.14 at 25 °C.

The chord area plot was used up to m=5. The resultant Φ_L curve was accepted as such. In the more concentrated region, another chord area plot was made using Thomsen's, Berthelot's, and Chedin and Dher's values. A preliminary curve of Φ_L was then made which was adjusted by using the long chords of Thomsen, Berthelot, and Forsythe and Giauque and the dilute Φ_L curve. The greatest weight was given to Forsythe and Giauque's measurements on the solution (dilution) of the liquid monohydrate and trihydrate to m=0.13.

CH₂O

 $\Delta H_{\rm diln}$ of aqueous formic acid has been measured by Thomsen [30]* (19 °C, m=111.01 to 0.28), Berthelot [233] (8.5 °C, m=1.0 to 0.25), and Faucon [234] (9 °C, m=413 to 1). $\Delta H_{\rm soln}$ as a function of concentration has been measured by Glagoleva [235]* (25 °C, m=677 to 0.52).

 ΔH_i of the reaction HCOOH(aq) \rightarrow H⁺ + HCOO-, has been estimated from the K_a 's given by Harned and Owen [9]* and Everett and Wynne-Jones [236]* and the ΔH_N experiments of von Steinwehr [192] and Canady, Papée, and Laidler [237]*.

The heat contribution due to ionization has been calculated by using the K_a listed in Harned and Owen [9] and incorporated into the Φ_L values. The Φ_L values are to be used only with $\Delta H^{\circ}_{\infty}$ (complete ionization).

$C_2H_4O_2$

 $\Delta H_{\rm diln}$ of aqueous acetic acid has been measured by Thomsen [30]* (19 °C, $m\!=\!111.01$ to 0.28), Berthelot [238] (23 and 24 °C, $m\!=\!2.78$ to 0.35), and [239] (8.5 °C, $m\!=\!1.0$ to 0.25), Faucon [234] (7.4 °C, extremely concentrated solution to $m\!=\!0.88$) and Richards and Gucker [157]* (16 and 20 °C, $m\!=\!2.22$ to 0.07).

 \bar{L}_1 has been measured by Payn and Perman [212] (20, 30, 40 to 70 °C, extremely concentrated solutions to m=1.72).

 $\Delta H_{\rm soin}$ as a function of concentration has been measured by Klibanova and Neĭman [240]* (24°C, m=538 to 0.77).

The ΔH_i of C₂H₄O₂ was estimated to be -60 cal/mole from the neutralization data of Thomsen [30]* (18 °C, with NaOH, $m_{\text{C}_2\text{H}_4\text{O}_2} = 0.28$), Canady, Papée, and Laidler [237]* (25 °C, with NaOH, $m_{\text{C}_2\text{H}_4\text{O}_2} = 0.005$), Richards and Mair [241]* (20 °C, with NaOH, $m_{\text{C}_2\text{H}_4\text{O}_2} = 0.56$), Berthelot [242] (9 °C, with NaOH, $m_{\text{C}_2\text{H}_4\text{O}_2} = 0.50$), Arrhenius [194] (21.5 °C, with NaOH), Lunden [243] (25 °C), Devoto [244], Jordan and Dumbaugh [245], Graham [246], Favre [247], and Andrews [248], and the K_a 's and emf data as listed in Harned and Owen [9]*.

The heat contribution due to ionization has been calculated and incorporated into the Φ_L table, which is to be used only with the $\Delta H_{\infty}^{\circ}$ for complete ionization.

NH₃ and NH₄OH

 $\Delta H_{\rm diln}$ has been measured by Thomsen [30]* (19 °C, $m\!=\!17.34$ to 1.11), Berthelot [249] (14 °C, $m\!=\!56.6$ to 0.28), Calvet [250] (17 °C, $m\!=\!3.60$ to 0.28), Ramstetter and Hantke [251] (20 °C, $m\!=\!2.56$ to 0.14), Baud and Gay [252] (12.5 °C, $m\!=\!12.97$ to 2.22), and Vrevskii and Zavaritskii [202, 214]* (14 °C, $m\!=\!18.50$ to 0.28).

°C, m = 18.50 to 0.28). Vrevskii and Zavaritskii [202, 214]* (3, 19.9, 41, and 61 °C, m = 53.89 to 0.44) measured ΔH_{soin} as a

function of concentration.

 ΔH_i is calculated as +865 cal/mole, by utilizing the data of Pitzer [206]* (25 °C, reaction of NaOH with NH₄Cl in dilute solution), the neutralization data of Thomsen [30]* (18 °C, with HCl and HNO₃, $m_{\rm NH_4OH} = 0.28$), Berthelot [253] (20 °C with HCl, $m_{\rm NH_4OH} = 0.50$), [215, 220] (18 °C, with HClO₄, $m_{\rm NH_4OH} = 0.25$), and the K_B 's at various temperatures and emf data as tabulated by Harned and Owen [9]*.

Other neutralization measurements are those of Bruni and Levi [254] (with HNO₃ and HClO₄), Berthelot [242, 255] (with acetic acid), [216] (with HClO₄), [256, 257] (with HNO₃ and HCl), Biltz and Messerknecht [258] (with HCl), Hess [259] (with HCl and HNO₃), Bouzat [260] (with HCl), Vorob'ev, Privalova, Monaenkova, and Skuratov [261] (with HClO₄), Favre and Silbermann [262] (with HCl and HNO₃), Favre [247] (with HNO₃, HCl and acetic acid), Andrews [263] (with unspecified acids), [248] (with HCl, HNO₃, and acetic acid), Lunden [243] (unspecified acids), Guntz [187] (with HF), and Delépine [264] (with HCl). Moore and Winmill [265] calculated the ΔH_i from conductivity measurements and partition coefficients at various temperatures

The degree of ionization was calculated using the value given by Harned and Owen [9] for K_B for NH₄OH. The heat contribution of the ionization has been incorporated into the tabulated Φ_L values. For convenience, the values are given for NH₄OH · nH₂O and NH₃ · nH₂O. The Φ_L values are to be used only with the value given for $\Delta H_{\infty}^{\circ}$ (complete ionization).

NH₄Cl

 $\Delta H_{\rm diin}$ has been measured by Thomsen [30]* (19 °C, m=5.55 to 0.28), Pollok [266] (9 °C, one dilution of a saturated solution to m=4), Mondain-Monval [267] (0.5 °C, one dilution from m=5.55 to 0.19 and 18.1 °C, one dilution from m=6.89 to 0.25) and [268] (0 and 19 °C, one dilution of a saturated solution to m=0.37), Pratt [269]* (0 to 35.5 °C, m=2.22 to 0.14), Magic [270]* (3.7 and 22.3 °C, m=2.22 to 0.28), Fedorov and Sil'chenko [271]* (22 °C, m=7.12 to 0.28), and Streeck [272]* (25 °C, m=1.07 to 0.0002).

 \overline{L}_1 has been measured by Dunnington and Hoggard [273] (20 to 25 °C, m = 5.05 to 2.31) and Rümelin

[274] (13 °C, m = 2.78 and 2.22).

 $\Delta H_{\rm soin}$ as a function of concentration has been measured by Mishchenko and Ponomareva [92]* (25 °C, m=7.09 to 0.03), Fedorov and Sil'chenko [271]* (22 °C, m=0.56 to 0.14), and Winkelmann

[94] (17 to 28 °C, m=6.23 to 0.58). von Stackelberg [275] measured the differential heat of solution (18 °C, m=5.88 to 0.18).

The chord area plot was used. The greatest weight was placed upon Streeck's data in the dilute region. The resultant Φ_L curve was checked by using the long chords of Thomsen, Fedorov and Sil'chenko, Magie, Pratt, and Mishchenko and Ponomareva.

NH₄ClO₄

The Φ_L values given in the discussion of $\Delta H_{\infty}^{\circ}$ were estimated, by considering the behavior of related compounds and the ΔH_N data of Berthelot [216] (20 °C, m=0.10) and Vorob'ev, Privalova, Monaenkova, and Skuratov [261]* (25 °C, $m_{\text{NH4 ClO4}} = 0.14$).

NH₄Br

Only one measurement of the $\Delta H_{\rm diln}$ has been reported, by Fedorov and Sil'chenko [271] (18 °C, one dilution from m=7.71 to 0.28). The tabulated values of Φ_L have been estimated, by considering the behavior of NH₄Cl, HCl, HBr, NaCl, and NaBr.

NH₄I

The tabulated values of Φ_L have been estimated. NH₄IO₃

The Φ_L values given for $\Delta H^{\circ}_{\infty}$ at 25 °C are estimated.

NH₄NO₃

 $\Delta H_{\rm diln}$ has been measured by Thomsen [30]* (18 °C, $m\!=\!22.20$ to 0.28), Pratt [269]* (0 to 33.5 °C, $m\!=\!2.22$ to 0.14), Mondain-Monval [276] (19 °C, one dilution of a saturated solution to $m\!=\!0.37$) and [267] (0 °C, $m\!=\!14.6$ to 0.39), Federov and Sil'chenko [271]* (21.5 °C, one dilution from $m\!=\!24.13$ to 0.28), Tollinger [97]* (21 °C, $m\!=\!18.50$ to 0.56), Wassermann [277]* (18.2 °C, $m\!=\!22.21$ to 0.55–0.14), von Wartenberg and Lerner-Steinberg [278]* (18.2 °C, $m\!=\!22.21$ to 0.56–0.24).

 \bar{L}_1 has been measured by Dunnington and Hoggard [273] (22.5 °C, m=18.50 to 2.22) and Rümelin [274] (11 °C, m=16.8 and 12 °C, m=8.0). Fricke and Havestadt [279] (0 °C, m=15.13 to 1.63) measured the vapor pressure of aqueous NH₄NO₃ from

which \bar{L}_1 may be calculated.

The $\Delta H_{\rm soln}$ data of Fedorov and Sil'chenko [271]* (21.5 °C, m=0.56 to 0.14) have also been used for

the Φ_L determinations.

The chord area plot was used below m=1. The long chords were then used with the Φ_L values obtained to obtain Φ_L for the more concentrated solutions.

NH₄C₂H₃O₂

 $\Delta H_{\rm diin}$ of aqueous ammonium acetate has been measured by Thomsen [30]* (20 °C, m=27.75 to 0.28), and Dunnington and Hoggard [273] (22.5 °C, m=18.50 to 2.28).

CH₃NH₂

 ΔH_{diin} has been measured by Bonnefoi [280] (12 °C, one dilution from m = 0.50 to 0.10).

 $\Delta H_{\rm soin}$ as a function of concentration has been measured by Felsing and Wohlford [281]* (35 °C, $m\!=\!0.19$ to 0.05). The dilution values at 35 °C in this range are assumed to be relatively the same as at 25 °C. Values of Φ_L are given relative to $m\!=\!0.056$.

$(CH_3)_2NH$

 $\Delta H_{\rm soin}$ as a function of concentration has been measured by Patterson and Felsing [282]* (30 °C, m=0.15 to 0.04). The scatter of their values permits only approximate estimates of Φ_L , relative to m=0.056.

$(CH_3)_3N$

 $\Delta H_{\rm soln}$ as a function of concentration has been measured by Patterson and Felsing [282]* (30 °C, m=0.15 to 0.05). The scatter of their values permits only approximate estimates of Φ_L , relative to m=0.056.

CH₃NH₃Cl

 $\Delta H_{\rm diln}$ has been measured by Streeck [272]* (25 °C, m=0.52 to 0.0002) and Whitlow and Felsing [107]* (25 °C, m=1.02 to 0.04). The chord area method was used to obtain preliminary values of Φ_L . These values were adjusted to fit Streeck's long chords.

(CH₃)₂NH₂Cl

 $\Delta H_{\rm diin}$ has been measured by Streeck [272]* (25 °C, m=0.52 to 0.0002) and Whitlow and Felsing [107]* (25 °C, m=1.01 to 0.05). The chord area method was used to obtain preliminary values of Φ_L . These values were adjusted to fit Streeck's long chords.

(CH₃)₃NHCl

 $\Delta H_{\rm diin}$ has been measured by Streeck [272]* (25 °C, $m\!=\!0.53$ to 0.0002) and Whitlow and Felsing [107]* (25 °C, $m\!=\!1.03$ to 0.05). The chord area method was used to obtain preliminary values of Φ_L . These values were adjusted to fit Streeck's long chords.

N(CH₃)₄Cl

 $\Delta H_{\rm diln}$ has been measured by Streeck [272]* (25 °C, m=0.53 to 0.0004). The chord area method was used to obtain preliminary values of Φ_L . These values were adjusted to fit Streeck's long chords.

AgF

 $\Delta H_{\rm diln}$ has been measured by Jahn-Held and Jellinek [110]* (25 °C, m=13.96 to 0.97). The tabulated values of Φ_L are referred to m=0.14, since AgF is partially unionized and the association constants available are not sufficiently extensive to enable calculating the contribution of the ionization to Φ_L at greater dilutions. Equilibrium constants have been given by Connick and Paul [283], Paul [284], and Leden and Marthen [285].

AgClO₄

von Wartenberg [286] reported one measurement (temperature unknown, m=2.41 to 1.0).

$AgNO_3$

Jackson, Smith, Gatty, and Wolfenden [287]* made one measurement (25 °C, m = 0.37 to 0.17) of $\Delta H_{\rm diln}$.

Terasaki [288] measured $\Delta H_{\rm soin}$ as a function of concentration (25 °C, 20 measurements, m=1.09 to 0.02). There is an error in the way the concentrations are expressed. The concentrations were corrected to the most reasonable form. The $\Delta H_{\rm diln}$ were then used to determine Φ_L . Although there is still an error in the concentrations, it was felt that the values obtained were reasonable and in line with the Φ_L values of the other nitrates.

LiOH

 $\Delta H_{\rm din}$ has been measured by Richards and Rowe [70]* (16 and 20 °C, m=2.22 to 0.14) and Ueda [289, 290]* (25 °C, saturated solution to 0.14). Juza and Laurer [291] measured ΔH_N of LiOH with HCI (20 °C).

°C). The chord area method was used to obtain Φ_L . These values were correlated with the emf data as given by Harned and Owen [9]*.

LiF

The Φ_L values were obtained from the $\Delta H_{\rm diln}$ measurements of Leighton [292]* (25 °C, m=0.04 to 0.01) and Lange and Leighton [293]* (25 °C, m=0.04 to 0.01) by using the chord area plot.

LiCl

 $\Delta H_{\rm diln}$ has been measured by Tucker [42]* (15 °C, $m\!=\!15.15$ to 1.82), Naude [199]* (17.5 to 26 °C, $m\!=\!1.0$ to 0.0003), Nernst and Orthmann [294, 295]* (18 °C, $m\!=\!0.10$ to 0.001), Lemoine [296] (10 and 20 °C, $m\!=\!16.62$ to 0.49), Lange and Messner [297, 298]* (25 °C, one dilution from $m\!=\!0.01$ to 0.0014), Richards and Rowe [70]* (16 and 20 °C, $m\!=\!2.22$ to 0.14), Birnthaler and Lange [299] (25 °C, values on graph from $m\!=\!18.32$ to 0), Lange and Dürr [118]* (25 °C, $m\!=\!19.88$ to 0.63), and Monheim and Streeck [300]* (25 °C, $m\!=\!0.28$ to 0.05).

 \bar{L}_1 has been reported by Dunnington and Hoggard [273] (22.5 °C, m=13.88 to 2.52) and Johnson and Molstad [301] (30, 50, and 70 °C, m=21.27 to 0.77).

 $\Delta H_{\rm soln}$ as a function of concentration has been measured by Lange and Dürr [118]* (25 °C, m = 18.31 to 0.08).

LiClO₄

 $\Delta H_{\rm diln}$ has been measured by Jones [302]* (25 °C, m=3.98 to 0.007) and Austin and Mair [219]* (25 °C, one dilution from m=1.0 to 0.025). The chord area method was used.

LiBr

 $\Delta H_{\rm diln}$ has been measured by Lange and Schwartz [119]* (25 °C, m=17.92 to 0.07), Lange and Messner [297, 298]* (25 °C, m=0.03 to 0.001), Birnthaler and Lange [299]* (25 °C, m=16.65 to 2.22), Mishchenko [303]* (25 °C, m=1.44 to 0.12), Lange and

Mishchenko [304]* (25 °C, one dilution from m = 1.44to 0.13) and Pennington [305] (25 °C, equation for the dilution of concentrated solutions from m =17.3-11.5 to a dilute solution).

 $\Delta H_{\rm soln}$ as a function of concentration has been measured by Lange and Schwartz [119]* (25 °C, m = 17.12 to 0.22).

The chord area plot was used to obtain Φ_L .

The tabulated values of Φ_L were estimated.

LiNO₂

Dodé [306] measured ΔH_N of LiOH with HNO₂; however, the concentration and temperature are uncertain.

LiNO₃

Richards and Rowe [70]* measured ΔH_{diln} (20 °C, m = 2.22 to 0.14). Φ_L was obtained by the chord area method.

Richards and Rowe [224] measured ΔH_N of LiOH with HNO₃ (16 and 17 °C, $m_{\text{LiNO}_3} = 0.28$). This was used to check Φ_L obtained by the chord area method.

NaOH

 ΔH_{diln} has been measured by Thomsen [30]* (19 °C, m = 18.50 to 0.56), Berthelot [307] (11 °C, m = 21.6 to 0.12), [225] (9.5 and 24 °C, one dilution from m = 6.3 to 0.66) and [233, 239] (8.5 °C, m = 1to 0.13), Fricke [308]* (19 to 20 °C, m=26.42 to 0.18), Calvet [250] (17 °C, m=2.58 to 0.28), Sturtevant [50, 309]* (25 °C, m=4.18 to 0.004), Pratt [269] (0 to 33.5 °C, m=4.11 to 0.14), Sabatier [161, 310] (17 °C, m=11.10 to 0.14), Richards and Gucker [157]* (16 and 20 °C, m = 2.22 to 0.02), Tucker [42]* (17 to 22 °C, m = 20.91 to 5.36), Bertetti and McCabe [311]* (20 °C, m = 23.20 to 3.92), Richards and Rowe [33]* (18 °C, one dilution from m = 9.49 to 1.28) and [70] (16 and 20 °C, m=9.64 to 0.14), de Forcrand [312] (15 °C, one dilution of a saturated solution to m = 10.8) and [313] (20 °C, m = 0.5 to 0.13), and Wilson and McCabe [125] (93.3 °C, m = 100 to 21.3). Richards and Hall [314]* recalculated the dilution data of Richards and Gucker with new specific heat data.

 \bar{L}_1 , from vapor pressure measurements, has been calculated by Payn and Perman [212] (20 to 70 °C, m=10.30 to 1.22) and Hayward and Perman [315] (20 and 30 °C, m = 10.7 to 6.25).

The chord area method was used. Greater weight was given in drawing the curves to Sturtevant's data below m=2, and to Bertetti and McCabe's data above that.

The calculated values of \bar{L}_2 check reasonably well with Akerlöf and Kegeles' [126] emf values below m=12. In the very dilute region, $m \le 0.5$, the agreement with the emf results given by Harned and Owen [9] is only fair.

The tabulated Φ_L values are estimated on the

basis of a comparison of LiF, LiOH, KF, KOH and NaOH.

NaCl

 $\Delta H_{\rm diln}$ has been measured by Scholz [316] (0 °C $m=6.\overline{01}$ to 0.25), Sandonnini and Gerosa [141] (18.5 °C, m=6.07 to 0.44), Pollok [266] (9 °C, one)dilution of a saturated solution to m=6), Backlund [317] (17 °C, m=4.40 to 0.37), Magie [270] (16.3 and 26.4 °C, m=2.22 to 0.28), Pratt [269] (0 to 33.3 $^{\circ}$ C, m = 2.22 to 0.14), von Stackelberg [275] (0 and 18 °C, m=4.40 to 0.56), Leitner [135] (20 °C, m=5.5to 0.56), van Deventer and van de Stadt [318] (one dilution from m=2.78 to 0.28), Naude [199, 319] (18 °C, m=1.0 to 0.001), Richards and Rowe [70] (20 °C, m = 3.2 to 0.19), Randall and Bisson [143] (25 °C, m=6 to 0.4), Malcolm, Parton, and Watson [320]* (25 °C, one dilution from m=2.70 to 0.025) Young and Machin [181]* (25 °C, m = 5.97 to 0.19) Lange and Messner [297, 298]* (25 °C, one dilution from m = 0.01 to 0.001), Robinson [321]* (25 °C) m = 0.10 to 0.0008), Gulbransen and Robinson [322] (25 °C, m = 0.82 to 0.005) in the form shown by Young and Groenier [180], Young and Vogel [183] $(25 \, ^{\circ}\text{C}, m = 6.15 \text{ to } 0.05)$, and Monheim and Streed [300]* $(25 \, ^{\circ}\text{C}, m = 0.32 \text{ to } 0.06)$.

 L_1 has been reported by Rümelin [274] (14 °C m=2.78 and 2.22) and Dunnington and Hoggard [273] (22.5 °C, m = 6.17 to 2.22).

The ΔH_{soln} data of Lipsett, Johnson, and Maass $[323]^*$ (20 and 25 °C, m=5.92 to 0.04), Wüst and Lange [21]* $(25 \, ^{\circ}\text{C}, m = 5.62 \text{ to } 0.52)$, and Benson and Benson [324]* (25 °C, m = 1.3 to 0.05), were used in the chord area method, along with the measured $\Delta H_{\rm diln}$. The resultant Φ_L curve was checked with the other ΔH_{diln} and ΔH_{soln} data. The agreement was satisfactory.

The \bar{L}_2 values were checked with those given by Harned and Owen [9] for combined calorimetic, em and boiling point data. The agreement is good be low m=2 and fair above.

NaClO₃

 ΔH_{diln} has been measured by Colomina and Nicolas [145]* (25 °C, one dilution from m=9.05 to 0.43) and Lange and Robinson [325]* (25 °C, m=0.10 to 0.0004). Colomina and Nicolas [145]* (25 °C, m = 8.54 to 0.11) measured ΔH_{soln} .

NaClO₄

 $\Delta H_{\rm diln}$ has been measured by Jackson, Smith Gatty, and Wolfenden [287]* (20 °C, one dilution from m = 0.51 to 0.23), Colomina and Nicolas [145] (25 °C, m=15.39 to 0.54), Austin and Mair $[219]^*$ (25 °C, one dilution from m = 1.0 to 0.025), and Van derzee and Swanson [221]* (25 °C, m = 11.80 to 0.01) $\Delta H_{\rm soln}$ has been measured by Colomina and Nic

olas [145]* (25 °C, m = 12.22 to 0.13).

The Φ_L values were based primarily upon Van derzee and Swanson's measurements. Colomina and Nicolas' values above m = 12 were coordinated with them to complete the Φ_L curve.

NaBr

 $\Delta H_{\rm diln}$ has been measured by Lange and Robinson [326, 327]* (25 °C, m = 0.16 to 0.0025), van Deventer and van de Stadt [318] (16 °C, m=8.20 to 0.28), Hammerschmid and Robinson [329]* (25 °C, m = 0.10 to 0.0003), Wüst and Lange [21]* (25 °C. m = 9.16 to 1.10), and Wallace [328]* (25 °C, m = 9.00to 0.25). Dunnington and Hoggard [273] (24 °C, m=4.27 to 2.52) measured \bar{L}_1 .

 $\Delta H_{\rm soln}$ has been measured by Wüst and Lange [21]* (25 °C, m=8.63 to 1.02).

NaBrO₃

The Φ_L values were obtained from Lange and Robinson's [325]* ΔH_{diln} measurements (25 °C, m = 0.10to 0.0004).

 ΔH_{diln} has been measured by Thomsen [30]* (18) °C, m = 5.55 to 0.56), Wüst and Lange [21]* (25 °C, m=12.34 to 9.85), and Lange and Robinson [326, 327]* (25 °C, $m \le 0.1$).

Wüst and Lange [21]* measured ΔH_{soln} (25 °C, m = 9.82 to 0.63).

NaIO₃

The Φ_L values are based on Lange and Robinson's [325]* measurements of ΔH_{diln} (25 °C, m=0.1 to 0.0004).

NaNO₂

The Φ_L values given are based on Perreu's [149]* measurements of ΔH_{diln} (14 °C, m = 10.86 to 0.19). Φ_L was assumed to be +40 at m=0.19, based on a comparison of the other Na+ electrolytes. The scale, however, is subject to adjustment.

NaNO₃

 ΔH_{diln} has been measured by Thomsen [30]* (18 °C, m = 9.25 to 0.28), Pollok [266, 330] (9 °C, one dilution of a saturated solution to m=5), von Stackelberg [275] (0 and 17 °C, m=8.88 to 0.28), Bishop [331]* (25 °C, m=2.17 to 1.50), Pratt [269]* (0 to 33 °C, m=2.22 to 0.14), Richards and Rowe [70]* (16 and 20 °C, m=2.22 to 0.14), van Deventer and van de Stadt [318] (17 °C, one dilution from m=4.91 to 0.28), Stahlberg [332]* (18 °C, m=7.40 to 0.19), Mondain-Monval [276]* (16 °C, m=9.39 to 0.54), and [267, 268] (0 and 16 °C, one dilution of a saturated solution to 0.37), Naude [199, 319]* (11 and 18 °C, m = 0.33 to 0.001), and Lange and Robinson [325]* (25 °C, m = 0.10 to 0.0002). \bar{L}_1 has been measured by Rümelin [274] (13 °C, m = 8.71 to 0.55).

Below m=0.1 the greatest weight in the chord area plot was given to the measurements of Lange and Robinson. The Φ_L values obtained in this region were used in conjunction with the long chords of the other investigators to adjust the Φ_L values obtained from the integration of the complete $d\Phi_L/dm^{1/2}$ plot.

NaC₂H₃O₂

 ΔH_{diln} has been measured by Thomsen [30]*

(19 °C, m = 5.55 to 0.56), Berthelot [333] (temperature unknown, one dilution from m = 0.50 to 0.25) and [233] (8.5 °C, m = 0.50 to 0.25), Richards and Gucker [157]* (16 and 20 °C, m = 2.22 to 0.03), Perreu [153] (18 °C, m = 4.78 to 0.12), and Fricke $[308]^*$ (19.5 °C, m = 14.92 to 0.28).

Fricke and Havestadt [279] (0 and 15 °C, m = 14.03to 5.24) and Perreu [153] (18 °C, saturated solution) measured L_1 .

NaCNS

 $\Delta H_{\rm diln}$ has been measured by Partington and Soper [334]* (25 °C, m=15.17 to 10.72). They [334]* also measured $\Delta H_{\rm soln}$ (25 °C, m=15.25 to

KF

 ΔH_{diln} has been measured by de Forcrand [335] (15 °C, m = 14.23 to 0.25), Guntz [187, 336] (temperature uncertain, m=2.0 to 0.50), Birnthaler and Lange [337]* (25 °C, m=15.55 to 0.06), Lange and Eichler [338]* (25 °C, m=16.92 to 1.52), Messner [339]* and Lange and Messner [340]* (m=0.21 to)0.004), and Monheim and Streeck [300] (25 °C, m = 0.30 to 0.06).

 ΔH_{soln} has been measured by Lange and Eichler

 $[338]^*$ (25 °C, m = 3.16 to 0.21).

 ΔH_{diln} has been measured by Thomsen [30]* (20 °C, m = 18.50 to 0.56), Berthelot [307]* (11.5) $^{\circ}$ C, m = 18.14 to 0.25) and [225] (11.5 and 24 $^{\circ}$ C, one dilution from m=1 to 0.5), Sabatier [310, 161]* (17 °C, m = 13.88 to 0.21), de Forcrand [312] (15 °C, one dilution of a saturated solution to a dilute solution), Pratt [269] (0 to 33.5 °C, m=3.65 to 0.14), Richards and Rowe [70]* (16 and 20 °C, m=5.55to 0.14), and Roth and Zeumer [341]* (20 °C, one measurement m = 0.83 to 0.03).

Biermann [342]* measured ΔH_N of KOH with HCl (25 °C, $m_{\text{KOH}} = 16$ to 3).

A chord area plot of the dilution data was made to obtain preliminary values of Φ_L . These values were adjusted to agree with the Φ_L values obtained from Biermann's neutralization data and with the emf data in the dilute region as summarized by Harned and Owen [8, 9]*.

 ΔH_{diln} has been measured by Stearn and Smith [343] (25 °C, m=3.16 to 0.18), Nernst and Orthmann [294, 295, 344]* (18 °C, m=0.33 to 0.002), Naude [1°9]* (12.2 and 17.8 °C, m=0.33 to 0.007) and [319] (18 °C, m=0.33 to 0.004), Richards and Rowe [71]* (16 and 20 °C, m=2.22 to 0.14), Lange and Leighton [345]* (12.5 and 25 °C, m=0.50 to 0.002), Bishop [331] (25 °C, m=4.49 to 0.06), Mondain-Monval [268] (18 °C, one dilution of a saturated solution to m = 0.37), Wüst and Lange [21]* (25 °C, m=4.82 to 0.57), Partington and Soper [334]* (25 °C, m = 4.82 to 0.30), Pratt [269] (0 to 33.5 °C, m = 2.22 to 0.14), Magie [270] (3 and 21 °C, m=2.22 to 0.28), van Deventer and van de Stadt

[318] (17 and 18 °C, m=4.24 to 0.28), Pollok [266] (9 °C, saturated solution to m=3), von Stackelberg [275] (0 and 18 °C, m=2.22 to 0.28), Sandonnini and Gerosa [141] (18.5 °C, m=4.33 to 0.28), Scholz [316] (0 °C, one dilution from m=3.62 to 1.00), and Monheim and Streeck [300]* (25 °C, m=0.30 to 0.06).

 \overline{L}_1 has been measured by Dunnington and Hoggard [273] (22.5 °C, m=4.27 to 2.0), Rümelin [274] (13.4 and 14.4 °C, m=2.78 and 2.22), and Harrison and Perman [346] (40, 50 to 80 °C, m=4.29 to 1.21).

 $\Delta H_{\rm soln}$ has been measured by Wüst and Lange [21]* (25 °C, m=4.47 to 0.37) and Partington and Soper [334]* (25 °C, m=3.55 to 0.31).

KClO₃

The Φ_L values are based on Andauer and Lange's [347]* measurements of $\Delta H_{\rm diln}$ (15 and 25 °C, m=0.10 to 0.0002).

Consideration was also given to von Stackelberg's [275] $\Delta H_{\rm soln}$ measurements (16 °C, m=0.22 and 0.11) and differential heats of solution (16 °C, m=0.50 to 0.22).

KClO₄

The Φ_L values are based on Andauer and Lange's [347]* measurements of $\Delta H_{\rm diln}$ (15 and 25 °C, m=0.10 to 0.0002).

KBr

 $\Delta H_{\rm diin}$ has been measured by Berthelot [348] (11 °C, one dilution from m=0.50 to 0.15), Scholz [316] (0 °C, m=4.0 to 0.25), Hammerschmid and Robinson [329]* (25 °C, m=0.10 to 0.0003), Wüst and Lange [21]* (25 °C, m=5.70 to 0.42), and Monheim and Streeck [300]* (25 °C, m=0.32 to 0.06).

 \bar{L}_1 has been measured by Dunnington and Hoggard [273] (22.5 °C, m=5.55 to 2.05) and Rümelin [274] (13.7 °C, m=2.78 and 11.5 °C, m=2.2).

 ΔH_{soln} has been measured by Wüst and Lange [21]* (25 °C, m=5.25 to 0.37).

KBrO₃

The tabulated Φ_L values have been estimated.

KI

 $\Delta H_{\rm diln}$ has been measured by Scholz [316] (0 °C, m=3.99 to 0.15), Monheim and Streeck [300]* (25 °C, m=0.32 to 0.06), and Wüst and Lange [21]* (25 °C, m=7.95 to 0.28); Dunnington and Hoggard [273] measured \bar{L}_1 (22.5 °C, m=9.25 to 1.85).

 $\Delta \hat{H}_{\rm soln}$ has been measured by Wüst and Lange [21]* (25 °C, m=8.97 to 7.34).

KIO₃

The tabulated Φ_L values were estimated considering the behavior of related compounds. The $\Delta H_{\rm diln}$ measurements of Berthelot [216, 349] (13 °C, m=0.5 to 0.13) and the differential heats of solution of von Stackelberg [275] (16 °C, m=0.28 to 0.11) were considered.

$KC_2H_3O_2$

 ΔH_{diin} has been measured by Thomsen [30]* (19 °C, m = 11.10 to 0.28).

KNO₃

 $\Delta H_{\rm diln}$ has been measured by Colson [350] (13.5 to 18 °C, $m\!=\!0.25$ to 0.01 and from a saturated solution to $m\!=\!0.13$), Naude [199]* (0, 12, 18, and 31 °C, $m\!=\!0.33$ to 0.002) and [319]* (18 °C, $m\!=\!0.35$ to 0.04), Lange and Messner [297, 298]* (25 °C, $m\!=\!0.01$ to 0.009), Lange and Monheim [173]* (12.5 and 25 °C, $m\!=\!0.13$ to 0.0006), Berenger-Calvet [351]* (16 °C, one dilution from $m\!=\!0.78$ to 0.06). Mondain-Monval [268] (0 and 18 °C, one dilution of a saturated solution to $m\!=\!0.37$), Nernst and Orthmann [294, 295, 344]* (18 °C, $m\!=\!0.10$ to 0.0003), Richards and Rowe [70]* (20 °C, $m\!=\!2.25$ to 0.14), Bishop [331]* (25 °C, $m\!=\!2.38$ to 0.49). Pratt [269]* (0 to 33.5 °C, $m\!=\!2.22$ to 0.14), Federov and Sil'chenko [271]* (22 °C, one dilution from $m\!=\!3.38$ to 0.28), Scholz [316] (0 °C, $m\!=\!1.0$ to 0.13). Pollok [266] (9 °C, one dilution of a saturated solution to $m\!=\!2$), and Graham [352] (temperature un known, $m\!=\!1$ to 0.5).

 \bar{L}_1 has been measured by Hunter and Bliss [353] (30 °C, m=2.55 to 1.12), and Rümelin [274] (13 °C, m=2.29 and 1.67).

 $\Delta H_{\rm soin}$ has been measured by Berenger-Calvet [351]* (16 °C, m=2.49 to 0.09), von Stackelberg [275]* (15 °C, m=2.44 to 0.22) and Winkelmann [94] (27.5 °C, m=2.41 to 0.31).

Below m=0.1 Lange and Monheim's data were given greater weight in the chord area plot. The Φ_L values obtained were adjusted above this molality by using the long chords.

KCNO

Myers [203]* and Vanderzee and Myers [204]* reported one dilution at 25 °C from m = 0.26 to 0.01. However, this was insufficient upon which to base a Φ_L curve.

KCNS

 $\Delta H_{\rm diin}$ has been measured by Partington and Soper [334]* (25 °C, m=23.96 to 15.6 and one dilution from m=23.96 to 0.65). Hunter and Bliss [353] measured \overline{L}_1 (30 °C, m=15.4 to 2.57).

Partington and Soper [334]* measured ΔH_{soln} (25 °C, m = 2.98 to 0.25).

The chord area plot was used below m=3. The long chord of Partington and Soper was used directly with the Φ_L at m=0.65 to determine Φ_L at m=23.96. This permitted the calculation of Φ_L from $\Delta H_{\rm diin}$ down to m=15.6.

The intermediate region was estimated with the aid of \bar{L}_1 .

KMnO₄

The Φ_L values were obtained from Perreu's [174]* measurements of $\Delta H_{\rm diln}$ (18 °C, m=0.40 to 0.05).

RbF

The Φ_L values are based on Lange and Monheim's [173]* measurements of $\Delta H_{\rm diln}$ (25 °C, m=0.21 to 0.0003).

RbCl

Monheim and Streeck [300] reported two measure-

ments of $\Delta H_{\rm diln}$ (25 °C, m=0.32 to 0.08 and m=0.32 to 0.06). However, their values on NaCl, KBr, KCl, KF, and KI exhibit systematic deviations. These measurements can therefore not be accepted. The tabulated Φ_L values were estimated on the basis of the behavior of CsCl, KCl, and KBr.

RbBr, RbI

The tabulated Φ_L values were estimated on the basis of the behavior of CsCl, KCl, KBr, and KI.

RbOH

de Forcrand [312] reported one measurement of ΔH_{diln} (15 °C, saturated solution to m=0.25).

de Forcrand [354] reported one measurement of ΔH_N of RbOH with HCl at 15 °C. No concentrations were mentioned.

RbNO₃

The tabulated Φ_L values were estimated on the basis of the behavior of CsNO₃, KNO₃, and NaNO₃.

CsOH

de Forcrand [312]* reported one measurement of ΔH_{diln} (15 °C, saturated solution to m=0.25).

Beketov [593] and de Forcrand [354] measured

 ΔH_N of CsOH with HCl at 16 and 15 °C, respectively. No concentrations were given; therefore the data cannot be used for the Φ_L determinations. CsF

The tabulated Φ_L values were estimated on the basis of the behavior of KF and RbF.

CsCl

 $\Delta H_{\rm diln}$ has been measured by Richards and Rowe [70]* (16 and 20 °C, $m\!=\!1.11$ to 0.14), Lange and Kohler [355]* (25 °C, below $m\!=\!0.25$), Messner [339]* (25 °C, below $m\!=\!0.25$), and Monheim and Streeck [300] (25 °C, two dilutions from $m\!=\!0.32$ to 0.08 and from $m\!=\!0.32$ to 0.06). Monheim and Streeck's measurements appear to have a systematic error.

CsBr and CsI

The tabulated Φ_L values were estimated on the basis of the behavior of CsCl, KCl, KBr, and KI. CsNO₃

The Φ_L values are based on Richards and Rowe's [70]* measurements of $\Delta H_{\rm diln}$ (20 °C, m=0.93 to 0.14).

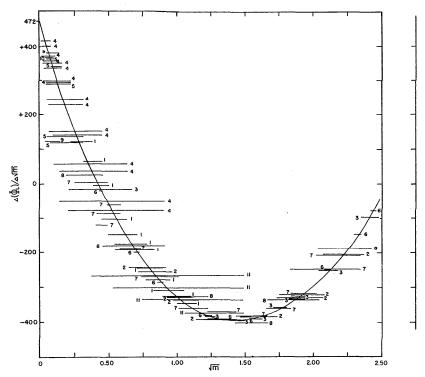


FIGURE IX. $\Delta(\Phi_L)/\Delta m^{\frac{1}{2}}$ versus $m^{\frac{1}{2}}$ for NaCl at 25 °C.

	The series of determinations are		
1.	Benson and Benson	25 °C ΔH _{soln} [324]	
2.	Wüst and Lange	25 °C ΔH _{soln} [21]	
3.	Young and Vogel	25 °C ΔH _{diin} [183]	
4.	Gulbransen and Robinson	25 °C ΔH _{diln} [322]	
5.	Robinson	25 °C ∆H _{diln} [321]	
6.	Young and Machin	25 °C ΔH _{diln} [181]	
7.	Lipsett, Johnson and Maass	25 °C ΔH _{soln} [323]	
8.	Lipsett, Johnson and Maass	20 °C ΔH soln [536]	
9.	Criss and Cobble	25 °C ∆H _{soln} [541]	
0.	Lange and Messner	20 °C ΔH _{diin} [297, 29	
1.	Richards and Rowe	20 °C ΔH _{aun} [70]	

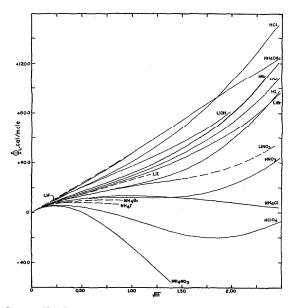


FIGURE X. The relative apparent molal enthalpy, Φ_L, in call mole, as a function of m^ω at 25 °C in aqueous solutions of IICl, HClO₄, HBr, HI, HNO₃, NH₄Cl, NH₄Br, NH₄I, NH₄NO₃, NH₄ acetate, LiOH, LiBr, LiI, and LiNO₃.

LiI, NH₄Br, and NH₄I are estimated. The broken portion of the lines for HClO₄ and LiNO₃ indicate the estimated values.

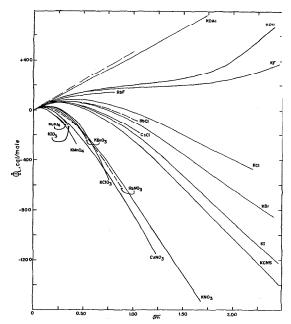


FIGURE XII. Φ_L as a function of m^{ν_L} in aqueous solutions of K, Rb, and Cs compounds at 25 °C.

The curves shown for RbNO₃ and RbCl are estimated.

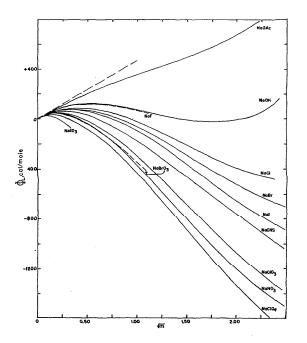


FIGURE XI. Φ_L as a function of m⁴ in aqueous solutions of NaOH, NaF, NaCl, NaBr, NaI, NaClO₃, NaClO₄, NaBrO₃, NaIO₃, NaNO₃, NaCNS, and Na acetate, at 25 °C.

NaF and NaBrO₂ are estimated.

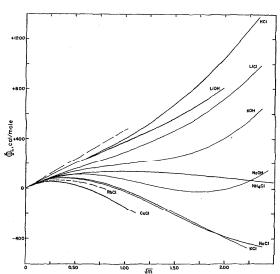


FIGURE XIII. Φ_L as a function of m^μ in aqueous solutions of the hydroxides and chlorides of H, NH4, Li, Na, K, Rb, and Cs, at 25 °C.

The curve for RbCl is estimated.

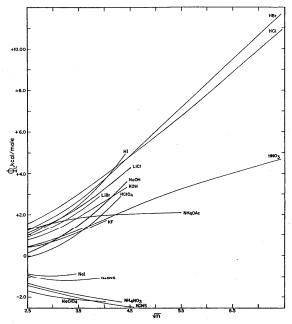


FIGURE XIV. Φ_L as a function of m¹⁶ $\geqslant 2.5$ in aqueous solutions at 25 °C.

VI. Heat of Neutralization

This discussion on ΔH_N is limited to the determination of ΔH_N° ; citations are given only for systems considered in this determination. Wherever ΔH_N° was used to determine Φ_L or ΔH_i for a compound, reference is made to it in section V.

Rossini's [3] procedure was used. All the data utilized have been recalculated, using the newly evaluated Φ_L and Φ_C values for the key components. Only the most reliable data, for systems where Φ_L for all three components was well established, were used to determine ΔH_N^o .

Table XVII lists the measurements (and their standard deviations) actually used for the final evaluation. This was limited to NaCl, KCl, NaNO₃, KNO3, NaClO4, and to the work of Richards and Rowe [224], Gillespie, Lambert, and Gibson [356], Bender and Biermann [213], Pitzer [206], and Vanderzee and Swanson [221, 357], despite the presence of reliable data, such as those of Richards and Rowe [224] on NaBr, NaI, KBr, KI, LiCl, LiBr, LiI, and LiNO₃, Biermann and Weber [223] on NaBr, and Biermann [342] on KCl, since the Φ_L values for HI, HBr, KOH, and LiOH were uncertain at the concentrations used in the neutralization experiments. Once ΔH_N^o was established, their results were used to help define these uncertain regions. There are two recent series of measurements that require particular mentionthose of Papée, Canady, and Laidler [358] and those of Bidinosti and Biermann [359]. Papée, Canady, and Laidler's measurements on the NaCl system in a microcalorimeter give a value of $-13,490 \pm 50$ cal/mole for ΔH_N° . The paper is not sufficiently detailed to determine whether precautions were taken to remove CO_2 from their dilute acid solutions, what ratio of NaOH to HCl was used, and how the extrapolation to infinite dilution was made. Bidinosti and Biermann's measurements on the NaClO₄ system give values for ΔH_N° that range from -13,207 to -13,171 cal/mole between 1 and 3 molal HClO₄ and from -13,237 to -13,300 cal/mole between 5 and 11 molal. There is no explanation for these discrepancies.

The tabulated values result in the following: Richards and Rowe, $-13,346\pm30$; Gillespie, Lambert, and Gibson, $-13,363\pm50$; Bender and Biermann, $-13,331\pm25$; Pitzer, $-13,366\pm25$; and Vanderzee and Swanson, $-13,336\pm20$ cal/mole. The "best" value for ΔH_N^o , 25 °C, has been selected as $-13,345\pm25$ cal/mole.

Using the Φ_{C}° values for HCl, NaOH, and NaCl at the various temperatures, ΔH_{N}° may be expressed, in the range 15 to 30 °C, as $(\Delta H_{N}^{\circ})_{t} = -13,345 + 53.5(t-25^{\circ}) - 0.46(t-25)^{2}$, or ΔH_{i} of H₂O as $(\Delta H_{i}^{\circ})_{t} = 13,345-53.5(t-25^{\circ}) + 0.46(t-25)^{2}$.

This equation differs from those of Rossini [3] and Pitzer [206], primarily because of $d(\Delta C_p^o)/dT$ which is now calculated to be 0.9 cal/mole deg instead of 0.3. The recent work of Ackermann [39, 122] indicates a greater variation of Φ_c^o of NaOH with temperature than previously accepted.

The value for ΔH_i° , at 20 °C = 13,710 cal/mole, given by Harned and Owen [9] from emf data, is 85 cal higher than that obtained by the use of the above equation. The equation given by Harned and Owen for the variation of ΔH_i° with temperature indicates a value of 13,510 cal/mole at 25 °C. The use of emf data for the heat of ionization involves the first and second derivatives with respect to T. The value obtained this way is generally not as reliable as that obtained calorimetrically.

Other measurements on the NaCl, NaNO₃, KCl, KNO₃, and NaClO₄ systems are as follows:

NaC

 ΔH_N has been measured by Thomsen [30], Berthelot [255, 257], Berthelot and Louguinine [360], Hess [361], von Steinwehr [192], Mathews [362], Mathews and Germann [363], Ellingson [364], Favre [247], Favre and Silbermann [262], Andrews [162, 248, 365], Roth [186], Könnecke, Steinert, and Leibnitz [366], Eley and Watts [367], Davies, Singer, and Stavely [368], Schultz [369], Keyes, Gillespie, and Mitsukuri [370], Douglas [371], Pitzer [206], Sacconi, Paoletti, and Ciampolini [205], Myers [203], Muller [372], Arrhenius [194], Lindner [373], Speakman and Stott [374], Gillespie, Lambert, and Gibson [356], Bender and Biermann [213], Olcott [375], Richards and Rowe [224], and Papée, Canady, and Laidler [358].

NaCIO

 ΔH_N has been measured by Thomsen [30], Berthelot [215, 216, 220], Vorob'ev, Privalova, Monaenkova, and Skuratov [261], and Bidinosti and Biermann [359].

NaNO₃

The ΔH_N has been measured by Thomsen [30], Berthelot [257], Arrhenius [194], Hess [259], Andrews [248], Favre [247], Mathews and Germann [363], Richards and Rowe [224], and Favre and Silbermann [262].

KCI

 ΔH_N has been measured by Thomsen [30], Muller [372, 376], Berthelot [257, 377, 378, 379], Hess [361,

380], Andrews [162, 248, 365], Favre and Silbermann [262], Favre [247], Gillespie, Lambert, and Gibson [356], Richards and Rowe [224], and Biermann [342].

KNO₃

 ΔH_N has been measured by Thomsen [30], Berthelot [257], Graham [246], Hess [259], Andrews [162, 248, 365], Favre and Silbermann [262], Favre [247], and Richards and Rowe [224].

VII. Heat of Solution at Infinite Dilution

The experimental data on $\Delta H_{\rm soln}$ were corrected to 25 °C, as explained earlier. The value of $\Delta H_{\infty}^{\circ}$ is obtained by using the relationship, $\Delta H_{\infty}^{\circ}$ = $\Delta H_{\rm soln,\,m} - \Phi_{L_m}$. A summary of the "best" values of $\Delta H_{\infty}^{\circ}$ for all the compounds can be found in table XXI.

The discussion below lists the experimental data (year reported, temperature of measurement, number of measurements, and concentration range), the values of ΔH_{∞}^{o} obtained and the "best" value for each compound. Wherever information is not reported by the investigator, it is designated by ?.

KCl(c)

 $\Delta H_{\rm soln}$ has been reported by a great many investigators. As KCl has been proposed as a standard for use in solution calorimetry and as, in many cases, the older literature must be relied upon for the values of $\Delta H_{\rm soln}$ of other compounds, all values between 15 and 30 °C reported for KCl have been tabulated, together with the estimated uncertainties, (table XVIII). The values may be separated into two distinct groups, $\Delta H_{\infty}^{\circ}=4,150$ cal/mole as shown by Wüst and Lange [21] and $\Delta H_{\infty}^{\circ}=4,115$ cal/mole.

The only apparent explanation at present, other than systematic errors in measurements, is that of variations in the treatment of the KCl sample. The "best" value of ΔH_{∞}^{o} has been selected as $4,115\pm10$ cal/mole, as indicated by the preponderance of data in this region in the past 20 years.

Other measurements on KCl besides those tabulated have been reported by Berthelot [225], Lange and Monheim [173, 613], Yatsimirskii and Grafova [437], Gire [438], Kaganovich and Mishchenko [439], Favre and Silbermann [262], Lehtonen [440], Zemczuzny and Rambach [441], Varali-Thevenet [442], Ostwald [443], Scholz [316], Samoĭlov and Buslacva [444], Kessler and Corbanev [445], Agostini [446], Richardson and Wells [447], Austin and Mair [219], and Vrzhesnevskii [448].

HF(g)

 $\Delta H_{\rm soln}$ has been measured by von Wartenberg and Fitzner [449] (1926; 32 °C, 10 measurements, m=0.14) and Roth [69] (1944; 20 °C, five measurements, m=33.32 to 5.55). The values of $\Delta H_{\infty}^{\circ}$ at 25 °C (complete ionization to H⁺ and F⁻) are -14,700 and -14,720 cal/mole, respectively. Khaĭdukov,

Linetskaya, and Bognovarov [450] have calculated $\Delta H_{\rm soln}$ at m=0.14 from vapor pressure measurements. Their value corrects to -14,817 cal/mole. Guntz [187] has also measured $\Delta H_{\rm soln}$.

von Wartenberg and Fitzner corrected their experimental values at m=0.14 on the basis of the presence of only the hexamer and monomer of HF in the gaseous state. The more recent data and calculations of Franck and Meyer [451], Smith [452, 453], Spalthoff and Franck [454], Long, Hildebrand, and Morrell [455], Hu, White, and Johnston [456], and Jarry and Davis [457] indicate the presence of other polymer species. These more recent data were used to recalculate the association factors for the conditions of von Wartenberg and Fitzner's experiments. A value of -11,600 cal/mole was obtained at m=0.14, t=25 °C, which leads to the above value of -14,700 cal/mole. This is in excellent agreement with a value independent of the polymerization of 14,690 cal/mole for the difference, ΔH_f HF(g) $-\Delta H_f$ F- (aqueous), obtained from the work of Hood and Woyski [634] on the reaction of NaCl(c) and HF(g) to form NaF(c) and HCl(g). It was not possible to recalculate Roth's value.

By using the value for ΔH_i selected in section V, we obtain for $\Delta H_{\infty}^{\circ}$ (no dissociation) -11,700 cal/mole, and $\Delta H_{\infty}^{\circ}$ (compete ionization) $=-\frac{14,700\pm200}{1000}$ cal/mole. The latter value is to be used with the values of Φ_L given in table XV, which include the effects of the partial ionization and association.

HCl(g)

 $\Delta H_{\rm soln}$ has been measured by Berthelot and Louguinine [458] (1875; 20 °C, four measurements, $m\!=\!0.40$ to 0.28), Thomsen [30] (1882; 19 °C, three measurements, $m\!=\!0.20$ to 0.14), Vrevskii and Zavaritskii [202, 214] (1923; 21.5 and 42 °C, 15 measurements, $m\!=\!4.83$ to 0.19), Roth and Richter [459] (1934; 21 °C, nine measurements, $m\!=\!0.09$ to 0.006), Roth and Bertram [460] (1937; 19 °C, seven measurements, $m\!=\!0.006$ to 0.0002), Roth, Börger, and Siemonsen [461] (1938; 20 °C, graph), Slansky [399] (1940; 25 °C, ?; ?), Gunn and Green [462] (1962; 25 °C, five measurements, $m\!=\!0.02$), and Vanderzee and Nutter [652] (1963; 25 °C, 12 measurements, $m\!=\!0.02$). The values of $\Delta H_{\infty}^{\circ}$ are: Berthelot and Louguinine, $-17,780\pm250$; Thomsen, $-17,760\pm100$; Vrevskii and Zavaritskii, $-17,837\pm50$; Roth and Richter, $-17,940\pm100$; Roth and Bertram, $-17,810\pm70$; Gunn and Green, $-17,888\pm15$; Van-

derzee and Nutter, $-17,875 \pm 12$ cal/mole. Roth. Börger, and Siemonsen's extrapolated value corrects to -17,950 cal/mole at 25 °C. Slansky gives a value of -17,820 cal/mole from his extrapolation. Vanderzee and Nutter's value, received after the value selected for $\Delta H_{\infty}^{\circ}$ was incorporated into the revision of Circular 500, confirms the selection.

Measurements have also been made by Favre and Silbermann [262], Abria [463], and Favre [464]. "Best" value, $\Delta H_{\infty}^{\circ} = -17,888 \pm 15$ cal/mole.

$HClO_4(1)$

 $\Delta H_{\rm soln}$ has been measured by Krivtsov, Rosolovskii, and Zinov'ev [217] (1960; 25 °C, one measurement, m = 0.07), Rosolovskii, Krivtsov, and Zinov'ev [465] (same measurement as first) and Berthelot

[215, 216] (1882; 19 °C, one measurement, m = 0.10). The values of $\Delta H_{\infty}^{\circ}$ are: Krivtsov et al., $-21,215 \pm 40$; Berthelot, -20,570 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -21,215 \pm 40$ cal/mole.

$HClO_4 \cdot H_2O(c)$

 $\Delta H_{\rm soln}$ has been measured by Krivtsov, Rosolovskii, and Zinov'ev [217] (1960; 25 °C, one measurement, m = 0.07); Arlman [466] (1939; one measurement, ?, ?), and Berthelot [215, 216, 220] (1882; 19 °C, one measurement, ?).

Krivtsov's value corrects to -7.875 ± 40 cal/mole for $\Delta H_{\infty}^{\circ}$. Arlman and Berthelot both report 7,700 cal/mole for $\Delta H_{\rm soln}$. "Best" value, $\Delta H_{\infty}^{\circ} = -7,875 \pm 40$ cal/mole.

HBr(g)

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, three measurements, m = 0.15 to 0.11), Roth and Bertram [460] (1935; 20.5 °C, 17 measurements, m=0.22 to 0.0005), Roth, Börger, and Siemonsen [461] (1938; 20.5 °C, results given in graph), and Vanderzee and Nutter [652] (1963; 25 °C, eight measurements, m = 0.02).

The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, $-20,290 \pm 150$; Roth and coworkers, $-20,260 \pm 100$; Vanderzee and Nutter, $-20,350 \pm 12$ cal/mole.

Measurements have also been reported by Favre and Silbermann [262], Ruff and Menzel [467], and Berthelot [198].

"Best" value, $\Delta H_{\infty}^{\circ} = -20,350 \pm 12$ cal/mole.

HI(g)

 $\Delta H_{\rm soln}$ has been measured by Berthelot and Louguinine [458] (1875; 20 °C, three measurements, m = 0.12 to 0.06), Thomsen [30] (1883; 18 °C, three measurements, m=0.11), Roth, Börger, and Siemonsen [461] (1938; 20 °C, results given in graph, m = 0.006 to 0.0004), and Roth [69] (1944; seven measurements, m = 1.11 to 0.02). The values of $\Delta H_{\infty}^{\circ}$ are: Berthelot and Louguinine, $-19,850 \pm 300$; Thomsen, $-19,390 \pm 150$; Roth, Börger, and Siemonsen, -19,680; Roth, $-19,520 \pm 50$ cal/mole.

Measurements have also been made by Favre and Silbermann [262] and Cottrell [468] (of HI(1)). "Best" value, $\Delta H_{\infty}^{\circ} = -19,520 \pm 50$ cal/mole.

HIO₃(c)

 ΔH_{soln} has been measured by Berthelot [216] (1883; 12 °C, one measurement, m = 0.13), Thomsen [30] (1883; 18 °C, two measurements, m = 0.28), and Moles and Perez-Vitoria [394] (1932; 20.5 °C, four measurements, m = 0.08). The values of ΔH_{soln} at 25 °C are: Berthelot, 2,060; Thomsen, $1,850 \pm 50$; Moles and Perez-Vitoria, 2,380 ± 50 cal/mole.

Ditte [469] also measured $\Delta H_{\rm soln}$. The specific heat of the solid used to correct to 25 °C was estimated. Since Φ_L is unknown, the extrapolation to infinite dilution cannot be made.

"Best" value, $\Delta \hat{H}_{\text{soin}} = 2,100 \pm 300 \text{ cal/mole.}$

$HNO_3(1)$

 ΔH_{soln} has been measured by Thomsen [30] (1882; 19 °C, ?, m = 0.19), Forsythe and Giauque [68] (1942; 25 °C, four measurements, m = 0.13), Karve [470], and Hess [471]. The differential heat of solution has been measured by Leclerc and Chédin [472]

and Chédin and Dhers [232]. The values of $\Delta H_{\infty}^{\circ}$ from the first two are: Thomsen, $-7,850\pm100$; Forsythe and Giauque,

 -7.954 ± 20 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -7,954 \pm 20$ cal/mole.

$CH_2O_2(1)$

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, one measurement, m=1.11), Glagoleva [235] (1936; 25 °C, 24 measurements, m=677 to 0.52), Berthelot [233,239], and André [473]. The values of $\Delta H_{\infty}^{\circ}$ (complete ionization) from the first two are -204 ± 20 cal/mole and -205 ± 10 cal/mole, respec-

By using the value for ΔH_i selected in section V we obtain for the best value, $\Delta H_{\infty}^{\circ}$ (no ionization) -175 ± 10 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ}$ (complete ionization)=-205

±20 cal/mole.

The latter value is to be used with the values of Φ_L tabulated.

$C_2H_4O_2(1)$

 $\Delta H_{\rm soin}$ has been measured by Berthelot [238] (1875; 7 to 23 °C, three measurements, m=2.78 to 0.35), Berthelot and Louguinine [458] (1875; 20 °C, three measurements, m = 0.44 to 0.19), Thomsen [30] (1883; 19 °C, two measurements, m=2.78 and 1.11), Pickering [474] (1895; 13.6 to 18.5 °C, five measurements, m=0.28 to 0.14), Klibanova and Neiman [240] (1933; 24 °C, 16 measurements, m=115 to 0.77), and Pritchard and Skinner [475] (1950; 21 °C, one measurement, m=0.08). The values of $\Delta H_{\infty}^{\circ}$ (complete ionization) are: Berthelot, -350; Berthelot and Louguinine, -446; Thomsen, -364; Pickering, -326; Klibanova and Neiman, -347; Pritchard and Skinner, -361 cal/mole.

By using the value for ΔH_i of $C_2H_4O_2$ determined

in section V we obtain:

value, $\Delta H_{\infty}^{\circ}$ (no ionization) = -300 "Best" cal/mole. "Best"

value, $\Delta H_{\infty}^{\circ}$ (complete ionization) = -360

cal/mole.

The latter value is to be used with the Φ_L values which include the ionization contribution.

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, three measurements, m=0.43 to 0.24), Vrevskii and Zavaritskii [202, 214] (1924; 19.9 and 41 °C, 29 measurements, m=34.05 to 0.61), and Ramstetter and Hantke [251] (1931; 20 °C, five measurements, m=2.57 to 0.80). The values, recomputed to $\Delta H_{\infty}^{\circ}$ (complete ionization) are: Thomsen, -7.587 ± 100 ; Vrevskii and Zavaritskii, $-7,443\pm40$; Ramstetter and Hantke, $-7,290\pm150$

Other measurements of ΔH_{soin} are those of Vrevskii and Zavaritskii [202, 214] (3 °C and 61 °C), Berthelot [249], Favre and Silbermann [262], Baud

and Gay [252], and Bouzat [260].

Vrevskii and Zavaritskii's value is preferred; however, as will be evident in the revision of Circular 500, Ramstetter and Hantke's value is more consistent with values obtained from other systems.

By using the value for ΔH_i NH₄OH (ΔH_i =+865 ±30 cal/mole) determined in section V we obtain:

"Best" value, $\Delta H_{\infty}^{\circ}$ (no ionization) = -8,160 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ}$ (complete ionization) = -7,290±150 cal/mole.

The values of Φ_L given for NH₄OH and NH₃ are to be used only with $\Delta H_{\infty}^{\circ}$ (complete ionization).

NH₄F(c)

Guntz [187] quotes a value of 1.5 kcal/mole from Favre for the $\Delta H_{\rm soln}$ at 13 °C. The concentration is not mentioned. The reference cannot be located. $\Delta H_{\rm soln} = 1,500$ cal/mole.

NH₄Cl(c)

 $\Delta H_{\rm soln}$ has been measured by Winkelmann [94] (1873; 17.2 to 28 °C, 10 measurements, m = 6.23 to 0.58), Berthelot [377, 476] (1873; 10 °C, one measurement, m=0.46), Thomsen [30] (1883; 18 °C, two measurements, m = 0.28), von Stackelberg [275] (1898; 18 °C, three measurements, m = 0.74 to 0.37), Colson [350] (1915; 14.9 and 22.3 °C, two measurements, m = 0.23), Mondain-Monval [268] (1923; 18 °C, one measurement, m=0.37), Nacken [477] (1930; 20 °C, two measurements, m = 0.27), Drucker [45] (1934; 18.1 °C, one measurement, m = 0.24), Federov and Sil'chenko [271] (1937; 22 °C, three measurements, m = 0.56 to 0.14), and Mishchenko and Ponomareva [92] (1956; 25°C, 21 measurements, m=7.09 to 0.03). The values of $\Delta H_{\infty}^{\circ}$ are: Winkelmann, 3,560 \pm 50; Berthelot, 3,400; Thomsen, 3,530 \pm 50; von Stackelberg, 3,586 \pm 50; Colson, 3,615 ±100; Mondain-Monval, 3,498; Nacken, $3,623 \pm 75$; Drucker, 3,528; Fedorov and Sil'chenko, 3,691 ±100; Mishchenko and Ponomareva, 3,533

±15 cal/mole. Measurements have also been reported by Favre and Silbermann [262], Anderson and Noyes [478], van Deventer and van de Stadt [318], Lehtonen [440], Varet [479], and André [480]. "Best" value, $\Delta H_{\infty}^{o} = 3,533 \pm 15$ cal/mole.

NH₄ClO₄(c)

 ΔH_{soln} has been measured by Birky and Hepler [481] (1960; 25 °C, six measurements, m = 0.009 to 0.001) and Vorob'ev, Privalova, Monaenkova, and Skuratov [261] (1960; 25 °C, one measurement, m=0.14). Φ_L has been estimated as 10 and 0 cal/mole, respectively, at the concentrations mentioned above. The values of $\Delta H_{\infty}^{\circ}$ are 8.020 ± 70 and $7,984 \pm 20$ cal/mole, respectively.

Berthelot [216, 220] has also measured ΔH_{soln} . "Best" value, $\Delta H_{\infty}^{\circ} = 8,000 \pm 40$ cal/mole.

NH₄Br(c)

 ΔH_{soin} has been measured by Thomsen [30, 482] (1877; 18 °C, four measurements, m = 0.28) and Federov and Sil'chenko [271] (1937; 18°C, three measurements, m=0.56 to 0.14). The values of ΔH_{∞}^{o} are 4,064±50 and 3,920±50 cal/mole, respectively.

Measurements have also been reported by Varet

[479] and André [480].

"Best" value, $\Delta H_{\infty}^{\circ} = 4{,}010 \pm 100$ cal/mole.

$NH_4I(c)$

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, four measurements, m = 0.28) and Varet [479] (1896; 18 °C, one measurement, ?). The specific heat of the crystal used to correct to 25 °C was estimated. From Thomsen's measurements, ΔH_{∞}^{o} is $3,280 \pm 60$ cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 3,280 \pm 60$ cal/mole.

$NH_4IO_3(c)$

 ΔH_{soln} has been measured by Kapustinskii, Shidlovskii, and Shidlovskaya [483] (1958; 25 °C, three measurements, m=0.07) and Spencer and Hepler [484] (1960; 25 °C, four measurements, m=0.014 to 0.007). Φ_L was assumed to be -20and +20 cal/mole, respectively. The values of $\Delta H_{\infty}^{\circ}$ are $7,590 \pm 100$ and $7,610 \pm 100$ cal/mole, respec-

tively. "Best" value, $\Delta H_{\infty}^{\circ} = 7,600 \pm 70$ cal/mole.

NH₄NO₂(c)

 ΔH_{soln} has been measured by Berthelot [485, 486] (1874; 12.5 °C, one measurement, m=0.13) and Bureau [487] (1937; 20 °C, one measurement, m=0.14). The specific heat of the solid and Φ_c used to correct to 25 °C were estimated. Φ_L has been estimated as 60 cal/mole at the above concentrations. The values of $\Delta H_{\infty}^{\circ}$ are 4,300 and 4,610 cal/mole, respectively. Favre and Silbermann [262] also report a measurement.

"Best" value, $\Delta H_{\infty}^{\circ} = 4,600 \pm 200$ cal/mole.

NH₄NO₃(IV)

 $\Delta H_{\rm soln}$ has been measured by Winklemann [94] (1873; 16 to 22 °C, six measurements, m = 3.12 to 0.39), Tollinger [97] (1876; 19 to 25 °C, three measurements) urements, m = 0.56), Thomsen [488] (1878; 18 °C two measurements, m = 0.28), Mondain-Monval [267] (1925; 28 °C, one measurement, m = 0.28), Becker and Roth [489] (1934; 20 °C, seven measurements, m = 0.05), and Fedorov and Sil'chenko [271] (1937; 21.5 °C, three measurements, m=0.56 to 0.14). The values of $\Delta H_{\infty}^{\circ}$ are: Winkelmann, 6,125 ±100; Tollinger, 6,040 ±100; Thomsen, 6,125±100; Tollinger, 6,040±100; Thomsen, 6,117±50; Mondain-Monval, 5,990; Becker and Roth, 6,093±40; Fedorov and Sil'chenko, 6,216±50 cal/mole.

Other measurements of $\Delta H_{\rm soln}$ are those of Berthelot [377], van Deventer and de Stadt [318], Favre and Silbermann [262], Cohen and Helderman [490, 491], Cohen, Moesveld, and Helderman [492], and Mondain-Monval [268, 493].

"Best" value, $\Delta H_{\infty}^{\circ} = 6{,}140 \pm 50$ cal/mole.

$NH_4C_2H_3O_2(c)$

 $\Delta H_{\rm soin}$ has been measured by Berthelot [494] (1873; 24 °C, one measurement, m=0.28). His value, recomputed to 25 °C, infinite dilution, and no hydrolysis, is -570 ±200 cal/mole.

 Φ_L as tabulated includes the hydrolysis effect.

NH₄CN(c)

 ΔH_{soln} has been measured by Berthelot [377] (1875; 18 °C, one measurement, m = 0.06) and [216, 495] (1871; ?, one measurement, m = 0.13). The ΔC_p for correction to 25 °C has been estimated. Φ_L has been estimated as -15 cal/mole at m = 0.06. ΔH_{\odot}° from Berthelot's first value is $4,200 \pm 250$ cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 4,200 \pm 250$ cal/mole.

NH₄CNS(c)

 $\Delta H_{\rm soln}$ has been measured by Joannis [496] (1882: 12 °C, four measurements, ?). We have assumed the concentration to be m=0.56, and Φ_L to be 30 cal/mole. $\Delta H_{\infty}^{\circ}$ is 5,400 ±200 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 5,400 \pm 200 \text{ cal/mole.}$

$CH_3NH_2(g)$

 ΔH_{soln} has been measured by Aston and Ziemer [108] (1946; 25 °C, five determinations, m = 0.104), Felsing and Thomas [497] (1929; 25 or 30 °C, seven determinations, m=0.16 to 0.06), Felsing and Wohlford [281] (1932; 35 °C, 12 determinations, m = 0.19 to 0.05), and Bonnefoi [280] (1901; 12 °C, two measurements, m = 0.09).

Aston and Ziemer's measurements on CH2NH2(1) were corrected to CH₃NH₂(g). Their corrected value for $\Delta H_{\text{soln, }m=0.104}$, $-11,167\pm57$ cal/mole, has been corrected to $\Delta H_{\text{soln, }m=0.0555}$. There appears to be a systematic error in Felsing's measurements.

The data are insufficient to warrant extrapolation to infinite dilution.

"Best" value, $\Delta H_{\text{soln, }m=0.0555} = -11,290 \pm 75 \text{ cal/}$ mole.

$(CH_3)_2NH(g)$

 $\Delta H_{\rm soln}$ has been measured by Patterson and Felsing [282] (1938; 30 °C, 15 measurements, m = 0.15 to 0.04). Their values were corrected to 25 °C; however, the random scatter of the values permits only an approximate value of $\Delta H_{\text{soln}, m=0.056}$, $=-12,900\pm200$ cal/mole. Extrapolation to infinite dilution is not possible.

$(CH_3)_3N(g)$

 ΔH_{soln} has been measured by Berthelot [498] (1880; 20 °C, one determination, m=0.21) and Patterson and Felsing [282] (1938; 30 °C, 15 measurements, m = 0.15 to 0.05). The values were corrected to 25 °C; however, the random scatter of the values permits only an approximate value of $\Delta H_{\text{soln}, m=0.056}$ $=-12,800 \pm 200 \text{ cal/mole}.$

CH₃NH₃Cl(e)

 ΔH_{soln} has been measured by Aston and Ziemer [108] (1946; 25 °C, four determinations, m = 0.12 to 0.10). The value of $\Delta H_{\infty}^{\circ}$ is 1,378 \pm 15 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 1,378 \pm 15$ cal/mole.

(CH₃)₃NHCl(c)

 $\Delta H_{\rm soln}$ has been measured by Berthelot [498] (1880; 18 °C, one determination, m=0.21). The specific heat of the crystal used to correct to 25 °C has been estimated. $\Delta H_{\infty}^{\circ} = 340 \pm 200 \text{ cal/mole.}$

Tentative "Best" value, $\Delta H_{\rm m}^{\circ} = 350 \pm 200$ cal/mole.

$N(CH_3)_4Cl(c)$

 $\Delta H_{\rm soln}$ has been measured by Askew et al., [499] (1934; 20 °C, two determinations, m = 0.01). The ΔC_P used to correct to 25 °C was estimated. $\Delta H_{\infty}^{\circ}$ = 975 ± 75 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 975 \pm 75$ cal/mole.

N(CH₃)₄Br(c)

 ΔH_{soln} has been measured by Samo'ilov [407, 408] (1951; 25 °C, one measurement, m = 0.06) and Askew et al., [499] (1934; 20 °C, three measurements, m=0.01 to 0.006). The ΔC_P used to correct to 25 °C was estimated. Φ_L has been estimated as 10 and 25 cal/mole, respectively. The values, recomputed to infinite dilution and 25 °C, are 5,430 and $5,800 \pm 100$ cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 5,800 \pm 100$ cal/mole.

$N(CH_3)_4I(c)$

 ΔH_{soln} has been measured by Walden [500] (1907; 19 to 20 °C, two determinations, m = 0.22 and 0.13) and Coulter, Pitzer, and Latimer [109] (1940; 25 °C, five determinations, m = 0.06 to 0.04). The latter investigators also measured the specific heat of the crystal; Φ_C was estimated. Φ_L has been estimated as -120 and 10 cal/mole, respectively. The values of $\Delta II_{\infty}^{\circ}$ are $10,080\pm100$ and $10,055\pm20$ cal/ mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 10,055 \pm 20$ cal/mole.

$AgF \cdot nH_2O(c)$

Guntz [187, 501] has measured ΔH_{soln} of AgF

Guntz [187, 301] has measured $\Delta H_{\rm soln}$ of Agr (1884; 10 °C, $\Delta H_{\rm dilute\ solution} = -3,400\ {\rm cal/mole})$ and AgF · 2H₂O, (10 °C, $\Delta H_{\rm dilute\ solution} = 1,500\ {\rm cal/mole})$. Guntz and Guntz [502, 503] measured $\Delta H_{\rm soln}$ of AgF (1913; 16 °C, $\Delta H_{\rm dilute\ solution} = -4,300\ {\rm cal/mole})$, AgF · H₂O (10 °C, $\Delta H_{\rm dilute\ solution} = -850\ {\rm cal/mole})$, and AgF · 4H₂O (13 °C, $\Delta H_{\rm dilute\ solution} = 1,500\ {\rm cal/mole})$, and AgF · 4H₂O (13 °C, $\Delta H_{\rm dilute\ solution} = 4,930\ {\rm cal/mole})$ mole).

"Best" values cannot be given at the present time since there is insufficient information available.

$AgClO_4(c)$

 ΔH_{soin} has been measured by Bruni and Levi [254] (1917; 12 °C, one measurement, m=0.06). The ΔC_p used to correct to 25 °C was estimated. Φ_L has been assumed to be -100 cal/mole. The value of $\Delta H_{\infty}^{\circ}$ is 1,760 ±150 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 1,760 \pm 150$ cal/mole.

$AgNO_2(c)$

 ΔH_{soln} has been measured by Berthelot [216] (1883; ?, one measurement, ?) and Bureau [487] (1937; 20 °C, one measurement, m=0.14). Φ_L has been estimated as -20 cal/mole at m = 0.14. Berthelot reports $\Delta H_{\text{soln}} = 8,740 \text{ cal/mole}$. Bureau's value, recomputed to infinite dilution and 25 °C, is 8,830 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 8,830 \pm 200$ cal/mole.

$AgNO_3(c)$

 ΔII_{soln} has been measured by Thomson [30] (1883; 18 °C, two measurements, m = 0.28), Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.12), Smith, Brown, and Pitzer [504] (1937; 25 °C, one measurement, m = 0.009), Roth [69] (1944; 17.4 to 19 °C, four measurements, m = 0.08 to 0.03), Graham and Hepler [505, 506] (1955; 25 °C, six measurements, m = 0.06 to 0.004), and Terasaki [288] (1957; 25 °C, 20 measurements, m = 1.09 to 0.02). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, 5,377 \pm 50; Lange and Martin, 5,475; Smith, Brown, and Pitzer, 5,365 ± 50; Roth, 5,428 ± 50; Graham and Hepler. $5,440\pm50$; Terasaki, $5,485\pm100$. Although the concentrations in Terasaki's paper were corrected to the most reasonable form, it was not possible to completely eliminate the errors.

Measurements have also been made by Berthelot [377], Favre and Silbermann [262], and Bruni and Levi [254].

"Best" value, $\Delta H_{\infty}^{\circ} = 5,400 \pm 50$ cal/mole.

LiOH(c)

 ΔH_{soln} has been measured by de Forcrand [507, 508] (1907; 15 °C, one measurement, m = 0.50), Ueda [289, 290] (1933; 25 °C, four measurements, m = 0.14), and Reshetnikov [429] (1961; 25 °C, seven measurements, m = 0.14). The values of $\Delta H_{\infty}^{\circ}$ are: de Forcrand, -4,990; Ueda, -5,040; Reshetnikov, $5,632\pm15$ cal/mole.

Truchot [509] has also reported a measurement. "Best" value, $\Delta H_{\infty}^{\circ} = -5.632 \pm 15$ cal/mole.

LiOH:H2O(c)

 ΔH_{soln} has been measured by de Forcrand [510] (1906; 18 °C, one measurement, m = 0.50) and [508] (1908; 15 °C, one measurement, m = 0.50), and Ueda [289, 290] (1933; 25 °C, three measurements, m=0.14). The values of ΔH_{∞}^{o} are: de Forcrand, -1,100 and -950; Ueda, $-1,020 \pm 200$ cal/mole.

The dissociation pressure measurements of Ueda [290] lead to a value of ΔH for LiOH(c) + H₂O(1) = LiOH \cdot H₂O(c) in poor agreement with that calculated from the $\Delta H_{\infty}^{\circ}$ values of the two solids, if $\Delta H_{\infty}^{\circ}$ =-1,020 cal/mole is used. The value is in excellent agreement, though, with that calculated using de Forcrand's measurements on LiOH(c) and LiOH·H₂O(c). We have combined this difference with the value for LiOH(c) to obtain the "best'

"Best" value, $\Delta H_{\infty}^{\circ} = -1,600 \pm 400$ cal/mole.

 ΔH_{soln} has been measured by Kapustinskii and Stakhanova [511] (1956; 25 °C, ?, ?) and Kolesov and Skuratov [512] (1961; 21.5 °C, 15 measurements, m=0.01). de Forcrand [513] thought he measured $\Delta H_{\rm soln}$ of LiF(c); solubility measurements indicate this was not possible, since his compound was much more soluble. Kapustinskii and Stakhanova give a value at infinite dilution of -1,330 cal/mole. Kolesov and Skuratov's measurements correct to 1.130 ± 25 cal/mole. The solubility measurements of Payne [514] indicate that the differential heat of solution at saturation is positive.

"Best" value, $\Delta H_{\infty}^{\circ} = 1{,}130 \pm 25$ cal/mole.

LiCl(c)

 ΔH_{soln} has been measured by Thomsen [30] (1883; 18.5 °C, three measurements, m = 0.28 to 0.19), Pickering [515] (1888; 18 °C, one measurement, m = 0.28), Bonnefoi [280, 516] (1897; 15 °C, one measurement, m = 0.25), Haigh [388] (1912; 21 °C, three measurements, m=0.14), Lange and Dürr [118] (1926; 25 °C, 78 measurements, m = 18.34 to 0.04), Maier [517] (1927; 19.5 °C, one measurement, m = 0.24), Wassermann [277] (1930; 18.3 °C, four measurements, m = 0.12 to 0.08), Askew et al., [499] (1934; 20 °C, one measurement, m = 0.04), Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.20), Samoilov [407, 408] (1951; 25 °C, one measurement, m=0.06), and Samoilov and Buslaeva [444] (1960; 25 and 35 °C, three measurements, m=0.06). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, $-8,704\pm50$; Pickering, $-8,500\pm100$; Bonnefoi, -8,847; Haigh, -8,700±50; Lange and Dürr, -8,813±40; Maier, -8,670; Wassermann, -8,884 ±30; Askew et al., -8,690; Lange and Martin, -8,963; Samoĭlov, -8,780; Samoĭlov and Buslaeva, -8,790 ±100. The precision of Lange and Dürr's values is excellent. The overall uncertainty should be ±10; however, there appears to be a systematic error in the measurements as shown by their specific heat measurements. In considering this, the "best" value has been estimated as $\Delta H_{\infty}^{\circ} = -8,850$ ±50 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -8.850 \pm 50$ cal/mole.

$LiCl \cdot H_2O(c)$

 $\Delta H_{\rm soln}$ has been measured by Bogorodskii [518] (1911, 18 to 23 °C, four measurements, m=0.40 to 0.28) and Maier [517] (1927; 19.5 °C, one measurement, m = 0.17). These values correct to -4,540

±100 and -4,590 cal/mole, respectively.

The measurements of Ueda [290], Maier [517], Appleby and Crawford [636], and Hüttig and Reuscher [637] on the dissociation pressure of LiCl · H₂O(c) lead to a value of ΔH for the reaction, $LiCl(c) + H_2O(1) = LiCl \cdot H_2O(c)$, in good agreement with the difference, $\Delta H_{\infty}^{\circ}$ LiCl(c) $-\Delta H_{\infty}^{\circ}$ LiCl · $H_2O(c)$. "Best" value, $\Delta H_{\infty}^{o^*} = -4,560 \pm 100$ cal/mole.

LiCl · 2H₂O(c)

 ΔH_{soln} has been measured by Bogorodskii [518] (1911; 18 to 23 °C, four measurements, m = 0.39 to 0.34). However, Campbell and Griffiths [638] reported the transition, LiCl \cdot 2H₂O(c) = LiCl \cdot H₂O(c) occurred at 12.5 °C.

Hüttig and Reuscher's [637] measurements of the dissociation pressure lead to a value of ΔH for the reaction, $LiCl \cdot H_2O(c) + H_2O(l) = LiCl \cdot$ 2H₂O(c), in excellent agreement with the difference $\Delta H_{\infty}^{\circ} \operatorname{LiCl} \cdot \operatorname{H}_{2}\operatorname{O}(c) - \Delta H_{\infty}^{\circ} \operatorname{LiCl} \cdot 2\operatorname{H}_{2}\operatorname{O}(c).$

Assuming that the compound exists at 25 °C we have, as "best" value, $\Delta H_{\infty}^{\circ} = -1,120 \pm 40$ cal/ mole.

LiClO₄(c)

 ΔH_{soin} has been measured by Smeets [395] (1933; 18 °C, two measurements, m = 0.24), Markowitz, Harris, and Stewart [519] (1959; 25 °C, one measurement, m = 0.12), Birky and Hepler [481] (1960; 25 °C nine measurements, m = 0.02 to 0.005), and Vorob'ev, Privalova, Monaenkova, and Skuratov [261] (1060, 25° °C) [261] (1960; 25 °C, one measurement, m = 0.13). The ΔC_p used to correct to 25 °C has been estimated. The values correct to: Smeets, $-6,511 \pm 75$; Markowitz et al., $-6,364\pm50$; Birky and Hepler, $-6,325 \pm 50$; Vorob'ev et al., $-6,414 \pm 60$ cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -6,345 \pm 50$ cal/mole.

LiClO₄·3H₂O(c)

 ΔH_{soln} has been measured by Smeets [395] (1933; 18 °C, two measurements, m = 0.17) and Markowitz, Harris, and Stewart [519] (1959; 25 °C, one measurement, m = 0.12). The ΔC_p used to correct to 25 °C has been estimated. Their values correct to 7,896 \pm 75 and 7,795 \pm 50 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 7,795 \pm 50$ cal/mole.

 $\Delta H_{\rm soln}$ has been measured by Bodisko [520] (1889; 17 °C, one measurement, m = 0.34), Bonnefoi [521] (1900; 14 °C, one measurement, m = 0.34), Maier [517] (1927; 17 °C, one measurement, m = 0.12), and Lange and Schwartz [119] (1928; 25 °C, 44 measurements, m = 17.12 to 0.22). Their values of ΔH_{∞}^{c} are: Bodisko, $-11,645 \pm 75$; Bonnefoi, -11,830

 ± 100 ; Maier, -11,850; Lange and Schwartz, -11,668 ± 50 . The precision of Lange and Schwartz' measurements is excellent; the standard deviation is ± 3 . In view of the apparent systematic error in their specific heat measurements, the uncertainty in their solution measurements has been increased to ±50 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -11,670 \pm 50$ cal/mole.

$LiBr \cdot H_2O(c)$

 ΔH_{soln} has been measured by Maier [517] (1927; 17 °C, one measurement, m=0.10). His value corrects to -5,555 cal/mole.

Maier's measurements of the dissociation pressure of LiBr·H₂O(c) lead to a ΔH for the reaction, $LiBr(c) + H_2O(1) = LiBr \cdot H_2O(c)$ in good agreement with the difference, $\Delta H_{\infty}^{\circ} \operatorname{LiBr(c)} - \Delta H_{\infty}^{\circ} \operatorname{LiBr \cdot H_2O(c)}$. "Best" value, $\Delta H_{\infty}^{\circ} = -5,560 \pm 100 \text{ cal/mole}$.

LiBr·2H₂O(c)

 ΔH_{soln} has been measured by Maier [517] (1927; 17 °C, one measurement, m = 0.08). The value of $\Delta H_{\infty}^{\circ}$ is -2,250 cal/mole.

Maier's measurements of the dissociation pressure are discordant and cannot be used. Hüttig and Reuscher's [637] measurements lead to a ΔH for the reaction, $\text{LiBr} \cdot \text{H}_2\text{O}(c) + \text{H}_2\text{O}(1) = \text{LiBr} \cdot 2\text{H}_2\text{O}(c)$, which is in poor agreement with the difference $\Delta H_{\infty}^{\circ} \text{LiBr} \cdot \text{H}_2\text{O(c)} - \Delta H_{\infty}^{\circ} \text{LiBr} \cdot 2\text{H}_2\text{O(c)}$. "Best" value, $\Delta H_{\infty}^{\circ} = -2,250 \pm 200 \text{ cal/mole}$.

$LiBrO_3(c)$

 $\Delta H_{\rm soln}$ has been measured by Boyd and Vaslow [522] (1962; 25 °C, three measurements, m=0.007). Φ_L has been estimated to be 30 cal/mole at this concentration. $\Delta H_{\infty}^{\circ}$ is 338 ± 40 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 340 \pm 40$ cal/mole.

LiI(c)

 ΔH_{soln} has been measured by Bodisko [523, 524, 525] (1888; 17 °C, one measurement, m = 0.18), Mosnier [526] (1897; 15 °C, one measurement, m=0.07), Beketov and Beketov [527] (1902; 18 °C, two measurements, m = 0.24), and Maier [517] (1927; 17 °C, one measurement, m=0.07). Beketov's measurements were on a mechanical mixture of Lil with NaCl and KCl. The values of $\Delta H_{\infty}^{\circ}$ are Bodisko, $-15,100 \pm 100$; Mosnier, $-15,280 \pm 150$; Beketov, $-15,030 \pm 150$; Maier, $-15,130 \pm 100$. "Best" value, $\Delta H_{\infty}^{\circ} = -15,130 \pm 75$ cal/mole.

$LiI \cdot H_2O(c)$

 ΔH_{soin} has been measured by Maier [517] (1927; 17 °C, one measurement, m=0.07). The value corrects to $-7,090\pm100$ cal/mole.

The dissociation pressure measurements of Maier [517] and Hüttig and Pohle [639] lead to a ΔH for the reaction, $LiI(c) + H_2O(1) = LiI \cdot H_2O(c)$, in fair agreement with the difference $\Delta H_{\infty}^{\circ}$ LiI(c) $-\Delta H_{\infty}^{\circ}$ Li $\mathrm{I}\cdot\mathrm{H}_{2}\mathrm{O}(\mathrm{c})$.

"Best" value, $\Delta H_{\infty}^{\circ} = -7,090 \pm 200$ cal/mole.

$LiI \cdot 2H_2O(e)$

 ΔH_{soln} has been measured by Maier [517] (1927; 17 °C, one measurement, m = 0.06). His value corrects to $-3,530 \pm 100$ cal/mole.

Maier's dissociation pressure measurements lead to a ΔH for the reaction LiI · H₂O(c) + H₂O(l) =LiI · 2H₂O(c), in good agreement with the difference, $\Delta H_{\infty}^{\circ}$ LiI · H₂O $-\Delta H_{\infty}^{\circ}$ LiI · 2H₂O. "Best" value, $\Delta H_{\infty}^{\circ}$ = $-3,530\pm100$ cal/mole.

$LiI \cdot 3H_2O(c)$

Maier [517] (1927; 17 °C, one measurement, m = 0.05) measured ΔH_{soln} . The value corrects to 140 cal/mole.

Maier's dissociation pressure measurements lead to a ΔH for the reaction, LiI·2H₂O(c) $+H_2O(1)=LiI\cdot 3H_2O(c)$ that is impossible. We have therefore based $\Delta H_{\infty}^{\circ}$ on the calorimetric value. "Best" value, $\Delta H_{\infty}^{\circ} = 140 \pm 100$ cal/mole.

LiNO₂(c)

 $\Delta H_{\rm soln}$ has been measured by Dodé [306, 528] (1936; 12 °C, three measurements, m=0.13) and Bureau [487] (1937; 20 °C, one measurement, m =0.14). The ΔC_p used to correct to 25 °C has been estimated. Φ_L has been estimated to be 130 cal/mole at the above concentrations. The values of $\Delta H_{\infty}^{\circ}$ are -2,636 and -2,625 cal/mole, respectively. "Best" value, $\Delta H_{\infty}^{\circ} = -2,630 \pm 100$ cal/mole.

$LiNO_2 \cdot H_2O(c)$

Bureau [487] (1937; 20 °C, one measurement, m = 0.14) has measured $\Delta H_{\rm soln}$. The ΔC_p used to correct to 25 °C has been estimated. Φ_L has been estimated as 130 cal/mole at this concentration. The value corrects to 1,680 cal/mole.

Ray and Ogg [640] reported one measurement of the dissociation pressure at 25 °C. This information is insufficient to use as corroborating evidence for the difference, $\Delta H_{\infty}^{\circ} \text{ LiNO}_{2}(c) - \Delta H_{\infty}^{\circ} \text{ LiNO}_{2} \cdot H_{2}O(c)$. The "best" value has been based solely on Bureau's measurement.

"Best" value, $\Delta H_{\infty}^{\circ} = 1,680 \pm 200$ cal/mole.

LiNO₃(c)

 ΔH_{soin} has been measured by Thomsen [30] (1883; 20 °C, two measurements, m=0.56), Pickering [515] (1888; 18 °C, one measurement, m=0.27), and Haigh [388] (1912; 21 °C, three measurements, m=0.12). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, -574 ± 50 ; Pickering, -626; and Haigh, -630 ± 50 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -600 \pm 40$ cal/mole.

NaOH(c)

 ΔH_{soln} has been measured by Thomsen [30] (1883; 17 °C, four measurements, m = 0.34 to 0.29), Berthelot [307] (1875; 10.5 °C, two measurements, m = 0.36), de Forcrand [529] (1902; 21.5 °C, one measurement, m = 0.35), Roth, Wirths, and Berendt [124] (1942; 20 °C, two measurements, m = 10.11 and 0.13), Bobtelsky and Lairsch [530] (1950; 23 °C, one

measurement, m = 0.25), Reshetnikov [429] (1961; 25 °C, five measurements, m = 0.14), and Murch and Giauque [531] (1962; 25 °C, five measurements, m=0.30). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, -10,320; Berthelot, -10,410; de Forcrand, -10,540; Roth and coworkers, -10,400; Bobtelsky and Lairsch, -10,190; Reshetnikov, $-10,555 \pm 20$ cal/mole. Murch and Giauque's value of -10,637 \pm 10 cal/mole was obtained from the extrapolation of their measurement on solid NaOH $\cdot nH_2\hat{O}$ where n varied from 0.97 to 0.12. We have accepted their

"Best" value, $\Delta H_{\infty}^{\circ} = -10.637 \pm 10 \text{ cal/mole.}$

$NaOH \cdot H_2O(c)$

 ΔH_{soln} has been measured by Pickering [532] (1895; 19 °C, two measurements, m = 0.30 and 0.19), de Forcrand [529] (1902; 21.5 °C, one measurement, m = 0.35), and Murch and Giauque [531] (1962; 25 $^{\circ}$ C, five measurements, m = 0.30). The corrected values are $-5,380 \pm 200, -5,400, \text{ and } -5,118 \pm 10$ cal/mole, respectively. Murch and Giauque's value was obtained as for the anhydrous NaOH. We have selected their value.

"Best" value, $\Delta H_{\infty}^{\circ} = -5{,}118 \pm 10 \text{ cal/mole.}$

NaF(c)

 ΔH_{soln} has been measured by Guntz [187] (1884; 12 °C, one measurement, m = 0.14), Hepler, Jolly, and Latimer [533] (1953; 25 °C, one measurement, m = 0.04), and Latimer and Jolly [534] (1953; 25 °C, two measurements, m = 0.04 and 0.02). The values of ∆H_∞ are: Cuntz, 130; Hepler, Jolly, and Latimer, 210; and Latimer and Jolly, 220 cal/mole.

de Forcrand [335] quotes a value which may have been taken from Guntz.

"Best" value, $\Delta H_{\infty}^{\circ} = 218 \pm 10$ cal/mole

NaCl(c)

The values of $\Delta H_{\infty}^{\circ}$ are shown in table XIX. Other measurements of the heat of solution have been reported by: Lehtonen [440], Scholz [316], Staub [542, 543], Berthelot [544], Berthelot and Ilosvay [383], Kessler and Gorbanev [445], Ostwald [443], Richardson and Wells [447], Brönsted [385], Sandonnini and Gerosa [141], Mishchenko and Yakovlev [545], Favre and Silbermann [262], Graham [246], Varali-Thevenet [442], von Stackelberg [275, 614], Zemczuzny and Rambach [441], Varet [479], Jones, Burgess, and Amis [546], Chipman, Johnson, and Maass [133], Allmand and Polack [547], and Koehler [612].

"Best" value, $\Delta H_{\infty}^{\circ} = 928 \pm 5$ cal/mole.

NaClO₂(c)

 $\Delta H_{\rm soln}$ has been measured by Fontana and Latimer [548] (1949; 25 °C, two measurements, m =0.005), Levi and Bisi [549] (1956; 25 °C, three measurements, dilute solution), and Ishi [550] (1961; 18 °C, two measurements, m=0.06). The ΔC_p used to correct to 25 °C has been estimated. Φ_L has been estimated as 25 cal/mole at m = 0.005 and 50 cal/mole at m = 0.06. Fontana and Latimer's value

corrects to 80 ± 10 cal/mole at infinite dilution. Ishi's value corrects to -20 ± 100 cal/mole at infinite dilution. Levi and Bisi give a value of 370 cal/mole for their "dilute solution."

"Best" value, $\Delta H_{\infty}^{\circ} = 80 \pm 10$ cal/mole.

$NaClO_2 \cdot 3H_2O(c)$

 $\Delta H_{\rm soln}$ has been measured by Levi and Bisi [549] (1956; 25 °C, three measurements, dilute solution) and Ishi [550] (1961; 18 °C, three measurements, m=0.06). The ΔC_p used to correct to 25 °C was estimated. Levi and Bisi report a value of 7,020 cal/mole for their dilute solution. Ishi's value corrects to 6,780 ± 100 cal/mole at 25 °C and infinite

The dissociation pressure measurements of Riganti [641] lead to ambiguous results and cannot be used as corroborating evidence for the difference, $\Delta H_{\infty}^{\circ} \text{NaClO}_{2}(c) - \Delta H_{\infty}^{\circ} \text{NaClO}_{2} \cdot 3H_{2}O(c)$.

"Best" value, $\Delta H_{\infty}^{\circ} = 6,830 \pm 100 \text{ cal/mole.}$

NaClO₃(c)

 $\Delta H_{\rm soln}$ has been measured by Colomina and Nicolas [145] (1949; 25 °C, 14 measurements, m=8.54to 0.11), Nelson, Moss, and Hepler [551] (1960; 25 °C, six measurements, m = 0.02 to 0.01), and Ishi [550] (1961; 18 °C, one measurement, m = 0.06). The values of $\Delta H_{\infty}^{\circ}$ are $5,191\pm10,\ 5,191\pm60,\$ and $5,056 \pm 100$ cal/mole, respectively.

Measurements have also been reported by Foote and Saxton [552], Brönsted [553], and Berthelot

"Best" value, $\Delta H_{\infty}^{\circ} = 5{,}191 \pm 10$ cal/mole.

NaClO₄(c)

Measurements of $\Delta H_{\rm soln}$ have been made by Smeets [395] (1933; 18 °C, two measurements, m=0.21), Askew et al., [499] (1934; 20 °C, two measurements, m = 0.005), Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.12), Colomina and Nicolas [145] (1949; 25 °C, 19 measurements, m = 12.22 to 0.13), and Vorob'ev et al., [261] (1960; 25 °C, ?, m = 0.12). The ΔC_p used to correct to 25 °C was estimated. Their values correct to: Smeets, $3,318\pm30$; Askew et al., $3,340\pm20$; Lange and Martin, 3,429; Colomina and Nicolas, 3,317±8; Vorob'ev et al., $3,329 \pm 20$ cal/mole.

Other measurements have been reported by Berthelot [216, 377].

"Best" value, $\Delta H_{\infty}^{\circ} = 3.317 \pm 8$ cal/mole.

$NaClO_4 \cdot H_2O(c)$

 ΔH_{soln} has been measured by Smeets [395] (1933; 18 °C, two measurements, m=0.20). The ΔC_p used to correct to 25 °C has been estimated. This value corrects to 5,384 \pm 50 cal/mole.

The dissociation pressure measurements of Markowitz and Boryta [642] lead to a ΔH for the reaction, $NaClO_4(c) + H_2O(1) = NaClO_4 \cdot H_2O(c)$, in fair agreement with the difference, $\Delta H_{\infty}^{\circ}$ NaClO₄(c) $-\Delta H_{\infty}^{\circ}$ $NaClO_4 \cdot H_2O(c)$.

"Best" value, $\Delta H_{\infty}^{\circ} = 5{,}380 \pm 50$ cal/mole.

NaBr(c)

 ΔH_{soln} has been measured by Thomsen [30] (1883; 17 and 20 °C, ?, m = 0.28), Chipman, Johnson, and Maass [133] (1929; 25.4 °C, 15 measurements, m = 1.42 to 0.11), Askew et al., [499] (1934; 20 °C) one measurement, m=0.01), Wüst and Lange [21] (1925; 25 °C, 15 measurements, m=8.63 to 1.02), Lange and Dürr [389] (1926; 25 °C, four measurements, m = 0.56), Slansky [399] (1940; 25 °C, own extrapolation to infinite dilution), and Wallace [328] (1949; 25 °C, six measurements, m = 0.06 to 0.03). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, -87; Chipman et al., -45 ± 200 ; Askew et al., -180; Wüst and Lange, -149 ± 15 ; Lange and Dürr, -248; Slansky, 40; Wallace, -144 ± 15 .

Other measurements have been reported by Berthelot [377], Favre and Valson [554], and Varet

"Best" value, $\Delta H_{\infty}^{\circ} = -144 \pm 15$ cal/mole.

$NaBr \cdot 2H_2O(c)$

 ΔH_{soln} has been measured by Berthelot [377] (1875; 11 °C, one measurement, m = 0.12), Thomsen [30] (1883; 19 °C, two measurements, m = 0.19), and Samoilov [408] (1952; 25 °C, one measurement, m=0.06). The values of $\Delta H_{\infty}^{\circ}$ are: Berthelot, 4,150; Thomsen, 4,454 \pm 50; Samoilov, 4,572 cal/ mole. Favre and Valson [554] also report a measurement.

The dissociation pressure measurements of Dingemans [643] and Bell [644] lead to a ΔH for the reaction, NaBr(c) + 2H₂O(1) = NaBr · 2H₂O(c), in good agreement with the difference, $\Delta H_{\infty}^{\circ}$ NaBr(c)

 $\Delta H_{\infty}^{\circ} \text{ NaBr} \cdot 2H_2O(c)$.

"Best" value, $\Delta H_{\infty}^{\circ} = 4{,}454 \pm 50$ cal/mole.

NaBrO₃(c)

 $\Delta H_{\rm soin}$ has been measured by Spencer and Hepler [484] (1960; 25 °C, six measurements, m=0.02 to 0.01) and Boyd and Vaslow [522] (1962; 25 °C, three measurements, m = 0.007 to 0.01). The values of $\Delta H_{\infty}^{\circ}$ are: 6,430 ± 60 and 6,350 ± 100 cal/mole, respectively. "Best" value, $\Delta H_{\infty}^{\circ} = 6.430 \pm 60$ cal/mole.

 $\Delta H_{\rm soln}$ has been measured by Berthelot [377] (1875; 11 °C, one measurement, m=0.11), Thomsen [30] (1882; 20.1 °C, two measurements, m=0.28), Pickering [515] (1888; 18 °C, one measurement, m = 0.28), Varet [479] (1896; 16 °C, one measurement, m = 0.50), Mosnier [526] (1897; 15 °C, one measurement, m = 0.07), Wüst and Lange [21] (1925; 25 °C, 14 measurements, m=9.82 to 0.63), Askew et al., [499] (1934; 20 °C, two measurements, m = 0.02), and Mishchenko and Sukhotin [555] (1954; 25 °C, five measurements, m = 0.10 to 0.02). values of $\Delta H_{\infty}^{\circ}$ are: Berthelot, $-1,800 \pm 200$; Thomsen, -1,406±100; Pickering, -1,652; Varet, -1,565; Mosnier, -1,721; Wiist and Lange, -1,808±20; Askew et al., -1,781±30; Mishchenko and Sukhotin, -1.788 ± 25 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -1,800 \pm 20$ cal/mole.

$NaI \cdot 2H_2O(c)$

 ΔH_{soin} has been measured by Berthelot [377] (1875; 11 °C, one measurement, m = 0.11), Thomsen [30] (1882; 19 °C, two measurements, m = 0.18) and Samoĭlov [408] (1952; 25 °C, one measurement, m = 0.06). The values of ΔH_{∞} are: Berthelot, 3,680; Thomsen, $3,855 \pm 50$; Samoilov, 4,184 cal/mole.

Favre and Valson [554] also report a measurement. The dissociation pressure measurements of Dingemans [645] lead to a ΔH for the reaction, NaI(c) $+2H_2O(1)=NaI \cdot 2H_2O(c)$, in excellent agreement with the difference, $\Delta H_{\infty}^{\circ} \text{NaI(c)} - \Delta H_{\infty}^{\circ} \text{NaI} \cdot 2\text{H}_{2}\text{O(c)}$. "Best" value, $\Delta H_{\infty}^{\circ} = 3.855 \pm 50$ cal/mole.

NaIO₃(c)

 ΔH_{soln} has been measured by Spencer and Hepler [484] (1960; 25 °C, four measurements, m = 0.01 to 0.004) and Shidlovskii and Valkina [556] (1961; 25 °C, three measurements, m = 0.06). The values of $\Delta H_{\infty}^{\circ}$ are 4,820 ± 60 and 4,890 ± 60 cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 4,850 \pm 60$ cal/mole.

NaIO₄(c)

Kapustinskii and Yatsimirskii [557] (1949; 25 °C, one measurement, ?) report a value of 8,910 cal/mole for $\Delta H_{\rm soln}$. Correction to infinite dilution is

"Best" value, $\Delta H_{\text{soln}} = 8,910 \text{ cal/mole.}$

NaNO₂(c)

 $\Delta H_{\rm soin}$ has been measured by Swietoslawski [558] (1910; 18 °C, one measurement, m = 0.16), Matignon and Marchal [559] (1920; 20 °C, three measurements, m = 0.16), Dodé [306, 528] (1936; 12 °C, two measurements, m = 0.13), Bureau [487] (1937; 20 °C, one measurement, m = 0.14), Perreu [149] (1941; 14 °C, four measurements, m = 0.14), and Reshetnikov [429] (1961; 25 °C, five measurements, m = 0.17). The specific heat of the crystal used to correct to 25 °C was obtained from Voskresenskaya, Yankovskaya, and Anosov [621]. Φ_L has been assumed to be 50 cal/mole at these concentrations. The values of ΔH_∞ are: Swietoslawski, 3,300; Matignon and Marchal, 3,300; Dodé, 3,110; Bureau, 3,370; Perreu, 3,300; Reshetnikov, 3,320 ±70 cal/mole. Reshetnikov's value is obtained from the ΔH_{soln} of a mechanical mixture of NaNO2 with NaOH. "Best" value, $\Delta H_{\infty}^{\circ} = 3,320 \pm 40$ cal/mole.

$NaNO_3(c)$

 ΔH_{soln} has been measured by Winkelmann [94] (1873; 20 to 28 °C, 16 measurements, m=2.79 to 0.37), Berthelot [377] (1875; 13 °C, one measuremeasu ment, m=0.74), Tilden [560] (1884; 16 °C, three measurements, m = 0.56), Thomsen [30] (1883; 18 C, three measurements, m = 0.28), von Stackelberg [275] (1898; 17 °C, two measurements, m = 0.28 and 0.14), Haigh [388] (1912; 21 °C, three measurements, m = 0.14), Mondain-Monval [268] (1923; 16 °C, one measurement, m=0.37), Voskresenskaya and Ponomareva [400] (1946; 25 °C, one measurement, m = 0.18), Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.12), and Van Tassel and Wendlandt [561] (1959; 25 °C, five measurements, m = 0.28). The values of $\Delta H_{\infty}^{\circ}$ are: Winkelmann, 4,840 ±150; Berthelot, 4,700; Tilden, 4,772; Thomsen, 4,904±50; von Stackelberg, 4,965±100; Haigh, 4,790; Mondain-Monval, 4,640; Voskresenskaya and Ponomareva, 4,825; Lange and Martin, 4,892; Van Tassel and Wendlandt, 4,980 ± 100 cal/mole. Measurements have also been reported by Scholz [316], Varali-Thevenet [442], Mondain-Monval [276], and Zawidzki and Schagger [562].

"Best" value, $\Delta H_{\infty}^{\circ} = 4,900 \pm 50$ cal/mole.

$NaC_2H_3O_2(c)$

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, two measurements, m = 0.28), Pickering [384] (1887; 15 to 25 °C, 39 measurements, m = 0.28) and Jolly [563, 564] (1952; 25 °C, three measurements, m = 0.008 to 0.005). The specific heat of the crystal used to correct to 25 °C was obtained from Strelkov [622]. The values of ΔH_2° are: Thomsen, $-4{,}136{\pm}50$; Pickering, $-4{,}200{\pm}50$; and Jolly, $-4{,}144{\pm}100$ cal/mole.

 $\Delta H_{\rm soln}$ has also been measured by Berthelot [377,

"Best" value, $\Delta H_{\infty}^{\circ} = -4,140 \pm 50$ cal/mole.

$NaC_2H_3O_2 \cdot 3H_2O(c)$

 ΔH_{soln} has been measured by Thomsen [30] (1883; 19 °C, two measurements, m=0.14), Pickering [384] (1887; 15 to 25 °C, 33 measurements m, = 0.28), Bindel [154] (1890; 21 °C, four measurements, m = 5.55 to 1.11), Calvet [250] (1933; 17 °C, three measurements, m = 0.28), and Perreu [153] (1940; 18 °C, one measurement, m = 0.12). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, $4,800\pm50$; Pickering, $4,715\pm50$; Bindel, $4,534 \pm 75$; Calvet, $4,682 \pm 75$; Perreu, $4,663 \pm 100$ cal/mole.

Other measurements have been reported by Berthelot [476] and Gnesotto and Fabris [156].

The dissociation vapor pressure measurements of Baxter and Cooper [646] lead to a ΔH for the reaction, NaC₂H₃O₂(c) + 3H₂O(1) = NaC₂H₃O₂ $^{\circ}3H_2O(c)$, in good agreement with the difference $\Delta H_{\infty}^{\circ}$ NaC₂H₃O₂(c) $-\Delta H_{\infty}^{\circ}$ NaC₂H₃O₂ $^{\circ}3H_2O(c)$. "Best" value, $\Delta H_{\infty}^{\circ}=4,700\pm75$ cal/mole.

NaCN(c)

 ΔII_{soln} has been measured by Joannis [496] (1882; 9 °C, three measurements, m = 0.56). The specific heat of the solid used to correct to 25 °C was obtained from Messer and Ziegler [623]. Φ_c was estimated. Φ_L has been estimated as -30cal/mole at m=0.56. The value of $\Delta H_{\infty}^{\circ}$ is 290 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 290 \pm 200 \text{ cal/mole.}$

$NaCN \cdot \frac{1}{2}H_2O(c)$

 ΔH_{soln} has been measured by Joannis [496] (1882; °C, one measurement, m=0.56). Φ_L has been stimated as -30 cal/mole at m=0.56. The value, estimated as -30 cal/mole at m = 0.56. recomputed to $\Delta H_{\infty}^{\circ}$, is 790 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 790 \pm 200$ cal/mole.

$NaCN \cdot 2H_2O(c)$

 ΔH_{soln} has been measured by Joannis [496] (1882; 9 °C, three measurements m = 0.56). Φ_L has been estimated as -30 cal/mole at m = 0.56. The value of ΔH_{∞}^{o} is 4,440 cal/mole. "Best" value, $\Delta H_{\infty}^{o} = 4,440 \pm 200$ cal/mole.

NaCNO(c)

 ΔH_{soln} has been measured by Lord and Woolf [565] (1954; 25 °C, one measurement, m=0.03) and Lemoult [566] (1899; 13 °C, four measurements, m=0.05 to 0.02). Φ_L has been estimated as 50 cal/mole at these concentrations. The values of $\Delta H_{\infty}^{\circ}$ are: 4,590 \pm 40 and 4,630 \pm 100 cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 4,590 \pm 40$ cal/mole.

NaCNS(c)

 $\Delta H_{\rm soln}$ has been measured by Vrzhesnevskii [448] (1912; 18 °C, one measurement, m=0.56) and Partington and Soper [334] (1929; 25 °C, 19 measurements, m=1.23 to 0.17). The ΔC_p used to correct to 25 °C was estimated. The values of ΔH_{∞}^{*} are 1,670 and $1,632 \pm 20$ cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 1,632 \pm 20$ cal/mole.

KOH(c)

 $\Delta H_{\rm soln}$ has been measured by Berthelot [307, 377] (1875; 11.4 °C, one measurement, m = 0.21), Thomsen [30] (1883; 18 °C, one measurement, m = 0.22), de Forcrand [567] (1902; 21.5 °C, one measurement, m=0.33), Voskresenskaya and Ponomareva [400] (1946; 25 °C, two measurements, m=0.18), and Reshetnikov [429] (1961; 25 °C, six measurements, m=0.09). The values of $\Delta H_{\infty}^{\circ}$ are: Berthelot, -13,270; Thomsen, $-13,750\pm100$; de Forcrand, -13,257; Voskresenskaya and Ponomareva, -7,660; Reshetnikov, $-13,769 \pm 20$ cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -13,769 \pm 20$ cal/mole.

$KOH \cdot nH_2O(c)$

de Forcrand [567] (1902; 21.5 °C, one measurement for each composition, m = 0.28) measured ΔH_{soln} of various compositions of KOH $\cdot nH_2O(c)$, where n = 0.25, 0.535, 0.91, 1.16,and 1.55. Berthelot's [307] measurements (1875; 11.4 °C, one measurement for each composition, m = 0.33) were on compositions where n=0.875 and 2. The $\Delta H_{\infty}^{\circ}$ values obtained were plotted, together with the values of $\Delta H_{\infty}^{\circ}$ KOH(c) to obtain the $\Delta H_{\infty}^{\circ}$ values for the hydrates, $KOH \cdot H_2O(c)$ and $KOH \cdot 1.5H_2O(c)$. Berthelot's measurements on KOH · 2H₂O(c) appear to be seriously in error. No value for this is given.

The dissociation pressure measurements of Shibata, Oda, and Furukawa [650] and Bolte [651] are inconclusive.

 $KOH \cdot H_2O(c)$. "Best" value, $\Delta H_{\infty}^{\circ} = -3,500 \pm 300$

 $KOH \cdot 1.5H_2O(c)$. "Best" value, $\Delta H_{\infty}^{\circ} = -2,500$ ± 300 cal/mole.

KF(c)

 ΔH_{soln} has been measured by Guntz [187] (1884; 18 °C, one measurement, m = 0.28), de Forcrand [335, 513] (1911; 15 °C, one measurement, m = 0.50), Lange and Eichler [338] (1927, 25 °C, eight measurements, m=3.16 to 0.21), and Lange and Martin [397] (1937; 25 °C, one measurement, m=0.31). The values, recomputed to $\Delta H_{\infty}^{\circ}$ are: Guntz, -3,950; de Forerand, -4,540; Lange and Eichler, -4,238 ± 40 (standard deviation is ± 4); Lange and Martin, 4,007 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -4,238 \pm 40$ cal/mole.

$KF \cdot 2H_2O(c)$

 ΔH_{soln} has been measured by Guntz [187] (1884; 20 °C, one measurement, m=0.28), de Forcrand [335, 513] (1911; 15 °C, one measurement, m = 0.50), and Lange and Eichler [338] (1937; 25 °C, two measurements, m = 0.47 and 0.40). The values of $\Delta H_{\infty}^{\circ}$ are: Guntz, 830; de Forcrand, 1,890; Lange and Eichler, $1,666 \pm 20$ cal/mole.

The dissociation pressure measurements of Bell [647] lead to a ΔH for the reaction, $KF(c) + 2H_2O(c)$ =KF·2H₂O(c), in excellent agreement with the difference $\Delta H_{\infty}^{\circ} \text{KF}(c) - \Delta H_{\infty}^{\circ} \text{KF} \cdot 2\text{H}_2\text{O}(c)$. "Best" value, $\Delta H_{\infty}^{\circ} = 1,666 \pm 20 \text{ cal/mole}$.

$KF \cdot 4H_2O(c)$

de Forcrand's reported value [335, 513] (1911; 15 °C, one measurement, m=0.51) for ΔH_{soln} is 6,160 cal/mole. The melting point, reported by Yatlov and Polyakov [635] is 17.5 °C. A hypothetical $\Delta H_{\infty}^{\circ}$ at 25 °C would be 6,080 ± 200 cal/mole.

KClO₃(c)

 $\Delta H_{\rm soln}$ has been measured by Berthelot [377] (1875; 10 °C, one measurement, m=0.18), Thomsen [30] (1882; 19 °C, two measurements, m=0.14), von Stackelberg [275] (1898; 16 °C, five measurements, m=0.50 to 0.11) and Nelson, Moss, and Hepler [551] (1960; 25 °C, six measurements, m=0.02 to 0.005). The values of $\Delta H_{\infty}^{\circ}$ are: Berthelot, 9,610; Thomsen, 9,916 ±50; von Stackelberg, $10,000 \pm 100$; Nelson, Moss, and Hepler, 9.870 ± 50 cal/mole.

Bindel [154] has also reported a measurement. "Best" value, $\Delta H_{\infty}^{\circ} = 9.890 \pm 50$ cal/mole.

KClO₄(c)

 ΔH_{soin} has been measured by von Stackelberg [275] (1898; 16 °C, four measurements, m=0.22to 0.06), Noyes and Sammet [568] (1903; 21 °C, seven measurements, m=0.12), Latimer and Ahlberg [569] (1930; 25 °C, ?, m=0.03), Birky and Hepler [481] (1960; 25 °C, seven measurements, m = 0.009 to 0.004) and Vorob'ev et al., [261] (1960; 25 °C, ?, m=0.01). The values of $\Delta H_{\infty}^{\circ}$ are: von Stackelberg, 12,310 \pm 100; Noyes and Sammet, 12,154 \pm 30; Latimer and Ahlberg, 12,110 \pm 50; Birky and Hepler, 12,300 ±60; Vorob'ev et al., $12.127 \pm 60 \text{ cal/mole.}$

Berthelot [215, 216] has also reported measure-

"Best" value, ΔH_{∞} = 12,200 ± 60 cal/mole.

KBr(c)

 $\Delta H_{\rm soln}$ has been measured by Thomsen [30] (1883; 19.6 °C, two measurements, m=0.28), Brönsted [385] (1906; 20 °C, eight measurements, m=0.08), Walden [500] (1907; 16.6 °C, two measurements, m = 0.28), Wüst and Lange [21] (1925; 25 °C, 31 measurements, m = 5.25 to 0.37), Popov, Bundel, and Choller [165] (1930; 20 °C, seven measurements, m=0.19), Chipman, Johnson, and Maass [133] (1929; 15.8 to 27 °C, 30 measurements, m=0.43 to 0.09), Fedorov and Sil'chenko [271] (1933; 23.5 °C, three measurements, m=0.56 to 0.14), Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.13), Popov, Skuratov, and Strel'tsova [172] (1940; 20.5 °C, m = 0.12), and Hietala [426] (1960; 25 °C, three measurements, m = 0.28). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, 4,769 ± 50; Brönsted, 4,756 \pm 50; Walden, 4,727 \pm 50; Popov, Bundel, and Choller, 4,685 \pm 50; Wüst and Lange, 4,777 \pm 40; Chipman, Johnson, and Maass, 4,753 ± 100; Federov and Sil'chenko, $4,889 \pm 100$; Lange and Martin, 4,783; Popov, Skuratov, and Strel'tsova, 4,542; Hietala, $4,739 \pm 20$ cal/mole. Measurements have also been reported by Scholz [316], Berthelot [348, 377], Berthelot and Ilosvay [383], and Balarew and Kolarow [570]. Wüst and Lange's measurements appear to run high, as shown by their work on KCl. In view of this and Hietala's more recent measurements on KBr and KCl, a value of 4,750 ± 20 cal/mole has been chosen.

"Best" value, $\Delta H_{\infty}^{\circ} = 4.750 \pm 20$ cal/mole.

KBrO₃(c)

 $\Delta H_{\rm soln}$ has been measured by Berthelot [216, 571] (1878; 11 °C, one measurement, m = 0.12), Thomsen [30] (1883; 19 °C, two measurements, m = 0.28), von Stackelberg [275] (1898; 16 °C, three measurements, $m = 0.2\tilde{2}$ and 0.11), Lange and Martin [397] (1937; 25 °C, one measurement, m=0.03), Mel, Jolly, and Latimer [572, 573] (1953; 25 °C, three measurements, m=0.02 to 0.009), and Boyd and Vaslow [522] (1962; 25 °C, one measurement, m=0.003). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, 9,832 ± 50; Berthelot, 9,300; von Stackelberg, $9,710 \pm 100$; Lange and Martin, 9,888; Mel, Jolly, and Latimer, 9,780 ±30; Boyd and Vaslow, 9,700 ± 100 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 9,830 \pm 50$ cal/mole.

KI(c)

 ΔH_{soln} has been measured by Thomsen [30] (1882; 19 °C, two measurements, m = 0.28), Wüst and Lange [21] (1925; 25 °C, 24 measurements, m = 7.95to 0.29), Lange and Martin [397] (1937; 25 °C, one measurement, m=0.11), Popov, Skuratov, and Strel'tsova [172] (1940; 20.5 °C, one measurement, m = 0.09), Bobtelsky and Lairsch [530] (1950; 24.5 °C, one measurement, m = 0.30), and Kapustinskii

and Drakin [411] (1952; 25 °C, one measurement, m = 0.0008). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, $4,830 \pm 50$; Wüst and Lange, $4,897 \pm 40$ (standard deviation is ±2); Lange and Martin, 4,873; Popov, Skuratov, and Strel'tsova, $4,790 \pm 100$; Bobtelsky and Lairsch, 5,020; Kapustinskii and Drakin, 4,830 ±70 cal/mole. Measurements have also been reported by Berthelot [377], Berthelot and Ilosvay [383], Scholz [316], Walden [500], Kantola [574], and Hieber and Mühlbauer [575].

Wüst and Lange's measurements tend to run slightly high. In view of this, a value of $4,860 \pm 30$

cal/mole has been selected.

"Best" value, $\Delta H_{\infty}^{\circ} = 4,860 \pm 30$ cal/mole.

$KIO_3(c)$

 ΔH_{soin} has been measured by Berthelot [349] (1878; 12 °C, one measurement, m = 0.12), Thomsen [30] (1882; 19 °C, three measurements, m = 0.13 to 0.10), von Stackelberg [275] (1898; 16 °C, four measurements, m=0.22 to 0.11), and Spencer and Hepler [484] (1960; 25 °C, five measurements, m = 0.01 to 0.006). The values of $\Delta H_{\infty}^{\circ}$ are; Berthelot, 5,600; Thomsen, 6,640 ±50; von Stackelberg, $6,610 \pm 100$; Spencer and Hepler, $6,630 \pm 50$ cal/

"Best" value, $\Delta H_{\infty}^{\circ} = 6.630 \pm 50$ cal/mole.

KNO₂(c)

 ΔH_{soln} has been measured by Dodé [306, 528] (1936; 12 °C, four measurements, m = 0.14), Bureau [487] (1937; 20 °C, one measurement, m = 0.14), and Reshetnikov [429] (1961; 25 °C, three measurements, m=0.17). The ΔC_p used to correct to 25 °C was estimated. Φ_L has been estimated as -50 cal/mole at the above concentrations. The values of ΔH_{\perp}° are: Dodé, $3,240 \pm 150$; Bureau, $3,270 \pm 150$; Reshetnikov, $3,190 \pm 70$ cal/mole. Reshetnikov's value is obtained from the $\Delta H_{\rm soln}$ of a mechanical mixture of KOH with KNO₂.

"Best" value, $\Delta H_{\infty}^{\circ} = 3{,}190 \pm 70$ cal/mole.

$KNO_3(c)$

Values for $\Delta H_{\infty}^{\circ}$ are shown in table XX. Other measurements have been reported by Lloyd and Wyatt [71], Scholz [316], Person [584], Hieber and Mühlbauer [575], Varali-Thevenet [442], Staub [543], Berthelot [225, 377], Anderson and Noyes [478], Leval't-Ezerskii [585], Graham [352], Colson [350], and Zawidzki and Schagger [562].

Part of the discrepancy in the tabulated values may be due to the presence of a few percent of the metastable form.

"Best" value, ΔH_{∞} = 8,340 ±30 cal/mole.

$KC_2H_3O_2(c)$

 $\Delta H_{\rm soln}$ has been measured by Thomsen [30] (1883; 19 °C, one measurement, m=0.28). The specific heat of the crystal used to correct to 25°C was estimated. The value of $\Delta H_{\infty}^{\circ}$ is $-3,665 \pm 50$ cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -3.665 \pm 50$ cal/mole.

KCN(c)

Thomsen [30] (1883; 18 °C, two measurements, m = 0.32) and Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.32) have measured ΔH_{soln} . Messer and Ziegler [623] measured the specific heat of the crystal. Φ_C was estimated. Φ_L has been estimated to be 10 and 60 cal/mole, respectively, at the measured concentrations. The values of ΔH_{∞}^{*} are 2,860±75 and 2,745 cal/mole, respectively.

Berthelot has also measured ΔH_{soln} [216, 586]. "Best" value, $\Delta H_{\infty}^{\circ} = 2,800 \pm 100$ cal/mole.

KCNO(c)

 $\Delta H_{\rm soln}$ has been measured by Myers [203] (1958; 25 °C, four measurements, m=0.01); Vanderzee and Myers [204] report the same measurements. Φ_L has been estimated as 40 cal/mole at m=0.01. Their value of $\Delta H_{\infty}^{\circ}$ is 4,840 \pm 25 cal/mole.

Berthelot [495] also has measured $\Delta H_{
m soln}$. "Best" value, ΔH_{∞} = 4,840 ±25 cal/mole.

KCNS(c)

 ΔH_{soln} has been measured by Joannis [496] (1882; 13 °C, three measurements, m=0.28), Vrzhesnevskii [448] (1912; 18 °C, one measurement, m = 0.56), and Partington and Soper [334] (1929; 25 °C, 14 measurements, m = 2.98 to 0.25). The specific heat of the crystal used to correct to 25 °C was estimated. The values of $\Delta H_{\infty}^{\circ}$ are: Joannis, 5,930; Vzzhesnevskii, 6,090; Partington and Soper, 5,789 \pm 50 cal/mole. "Best" value, $\Delta H_{\omega}^{\circ} = 5,790 \pm 50$ cal/mole.

KMnO₄(c)

 ΔH_{soln} has been measured by Thomsen [30] (1883; 18 °C, two measurements, m-0.11), Roth and Becker [587] (1932; 16 to 21 °C, six measurements, m = 0.08 to 0.01), Perreu [174] (1940; 18 °C, one measurement, m=0.05), and Nelson, Moss, and Hepler [551] (1960; 25 °C, six measurements, m=0.01 to 0.005). The values of $\Delta H_{\infty}^{\circ}$ are: Thomsen, $10,260 \pm 150$; Roth and Becker, $10,340 \pm 100$; Perreu, 10,180; Nelson, Moss, and Hepler, 10,410 ± 60 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 10,410 \pm 60$ cal/mole.

RbOH(c)

 ΔH_{soln} has been measured by de Forcrand [510] (1906; 20 °C, one measurement, m=0.50). The specific heat of the crystal used to correct to 25 °C was estimated. Φ_L has been estimated as 200 cal/mole at this concentration. The value of $\Delta H_{\infty}^{\circ}$ is -14,900 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -14,900 \pm 200$ cal/mole.

RbOH·H₂O(c)

 ΔH_{soln} has been measured by de Forcrand [510] (1906; 20 °C, one measurement, m = 0.50). Φ_L has been estimated as 200 cal/mole at this concentration. The value of $\Delta H_{\infty}^{\circ}$ is -4,310 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -4,310 \pm 200$ cal/mole.

RbOH·2H₂O(c)

 $\Delta H_{\rm soln}$ has been measured by de Forcrand [510] (1909; 15 °C, one measurement, m = 0.50). Φ_L has been estimated as 200 cal/mole at this concentration. The value of $\Delta H_{\infty}^{\circ}$ is 210 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 210 \pm 200$ cal/mole.

RbF(c)

de Forcrand [513, 588] (1911; 15 °C, one measurement, m = 0.50), measured ΔH_{soln} . The value of $\Delta H_{\infty}^{\circ}$ is -6,240 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -6,240 \pm 200$ cal/mole.

$RbF \cdot H_2O(c)$

de Forcrand [589] (1911; 15 °C, one measurement, m=0.50), measured $\Delta H_{\rm soln}$ of the substance with the composition RbF·1/3H₂O. Lannung [648] showed the existence of RbF·H₂O. The $\Delta H_{\infty}^{\circ}$ RbF(c) and the $\Delta H_{\infty}^{\circ}$ for RbF·1/3H₂O were used to obtain $\Delta H_{\infty}^{\circ}$ RbF·H₂O(c) = -100 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ}$ = -100 ±300 cal/mole.

$\mathbf{RbF} \cdot \mathbf{l} \cdot \mathbf{l} \cdot \mathbf{H}_2 \mathbf{O}(\mathbf{c})$

de Forcrand [589] (1911; 15 °C, one measurement, m = 0.50), measured ΔH_{soln} . The value of ΔH_{∞}^{0} is 320 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 320 \pm 200$ cal/mole.

RbCl(c)

 $\Delta H_{\rm soln}$ has been measured by de Forcrand [354] (1906; 19 °C, one measurement, m = 0.25) and [513, 588] (1911; 15 °C, one measurement, m = 0.50), Zemczuzny and Rambach [441] (1910; 18 °C, one measurement, m = 0.56), Haigh [388] (1912; 21 °C. three measurements, m = 0.14), Samollov [590] (1956; 25 °C, one measurement, m = 0.06), and Samoilov and Buslaeva [444] (1960; 15 and 35 °C, two measurements, m=0.06). The values of $\Delta H_{\infty}^{\circ}$ are: de Forcrand, 4,028 and 4,100; Zemczuzny and Rambach, 4,127; Haigh, 4,000 ±50; Samoilov, 4,112 ±50; Samoilov and Buslaeva, 4,151 ±75 cal/mole.

"Best" value, $\Delta H_{\infty} = 4,130 \pm 75$ cal/mole.

RbClO₃(c)

 ΔH_{soln} has been measured by Pitzer [591] (1938; 25 °C, two measurements, m = 0.01). Φ_L has been estimated as 25 cal/mole at this concentration. The value of $\Delta H_{\infty}^{\circ}$ is 11,410 \pm 60 cal/mole.

'Best' value, $\Delta H_{\infty}^{\circ} = 11,410 \pm 60 \text{ cal/mole.}$

RbClO₄(c)

 ΔH_{soln} has been measured by Pitzer [591] (1938; 25 °C, two measurements, m = 0.01). Φ_L has been estimated as 10 cal/mole at this concentration. The value of ΔH_{∞}^{*} is 13,565 ± 60 cal/mole.

"Best" value, $\Delta H_{\infty} = 13,560 \pm 60$ cal/mole.

RbBr(c)

 ΔH_{soln} has been measured by de Forcrand [513, 588] (1911; 15 °C, one measurement, m = 0.50) and Lange and Martin [397] (1937; 25 °C, one measurement, m = 0.11). The values of $\Delta H_{\infty}^{\circ}$ are 5,618 and 5,230 cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 5,230 \pm 100$ cal/mole.

RbBrO₃(c)

 ΔH_{soln} has been measured by Boyd and Vaslow [522] (1962; 25 °C, three measurements, m = 0.004). Φ_L has been estimated to be 20 cal/mole at this concentration. The value of ΔH_{∞}^{o} is $11,700 \pm 100$ cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 11,700 \pm 100 \text{ cal/mole.}$

RbI(c)

 ΔH_{soln} has been measured by Mosnier [526] (1897; 15 °C, one measurement, m = 0.06) and de Forcrand [513, 588] (1911; 15 °C, one measurement, m = 0.50). The values of $\Delta H_{\infty}^{\circ}$ are 5,675 and 6,225 cal/mole, respectively. "Best" value, $\Delta H_{\infty}^{\circ} = 6,000 \pm 300$ cal/mole.

RbNO₃(c)

 $\Delta H_{\rm soln}$ has been measured by Haigh [388] (1912; 21 °C, three measurements, m = 0.14) and Vorobiev, Skuratov, and Ebragen [592] (1961; ?, ?, m = 0.03). The specific heat of the crystal used to correct to 25 °C was obtained from Mustajoki [624]. Haigh's value corrects to 8,716 ±60 cal/mole at 25 °C and infinite dilution. Vorobiev's reported value of 8,910 for ΔH_{soln} is a preliminary one; hence it is given little weight in the selection of $\Delta H_{\infty}^{\circ}$.
"Best" value, $\Delta H_{\infty}^{\circ} = 8,720 \pm 60$ cal mole.

CsOH(c)

 ΔH_{soln} has been measured by Beketov [593] (1892; 16 °C, one measurement, m=0.17) and de Forcrand [510] (1906; 15 °C, one measurement, m=0.50). The specific heat of the solid used to correct to 25 °C was estimated. Φ_L has been estimated as 160 and 230 cal/mole, respectively, at the measured concentrations. The value of $\Delta H_{\infty}^{\circ}$ arc -16,470 and -17,090 ±200 cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = -17,100 \pm 200$ cal/mole.

$CsOH \cdot H_2O(c)$

 ΔH_{soln} has been measured by de Forcrand [510] (1906; 15 °C, one measurement, m = 0.50). Φ_L has been estimated as 230 cal/mole at this concentration. The value of $\Delta H_{\infty}^{\circ}$ is $-4,900\pm200$ cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = -4,900\pm200$ cal/mole.

CsF(c)

 $\Delta H_{\rm soln}$ has been measured by de Forcrand [513, 588] (1911; 15 °C, one measurement, m=0.50). The value of $\Delta H_{\infty}^{\circ}$ is -8.810 ± 200 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -8.810 \pm 200$ cal/mole.

$CsF \cdot H_2O(c)$

de Forcrand [589] (1911; 15 °C, one measurement, m=0.50) measured $\Delta H_{\rm soln}$ of the substance with the composition CsF \cdot 2/3H₂O. Lannung [648] showed the existence of CsF $H_2O(c)$. $\Delta H_{\infty}^{\circ}$ CsF

and ΔH_m° for CsF · 2/3H₂O were used to obtain ΔH_{m}° of CsF · H₂O = -2,500 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = -2,500 \pm 300 \text{ cal/mole.}$

$CsF \cdot l \frac{1}{2} H_2O(c)$

de Forcrand [589] (1911; 15 °C, one measurement, m = 0.50) has measured $\Delta H_{\rm soln}$. The value of $\Delta H_{\infty}^{\circ}$ is -1,300 cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} - -1,300 \pm 200$ cal/mole.

CsCl(c)

 ΔH_{soln} has been measured by de Forcrand [354] (1906; 19 °C, one measurement, m = 0.25) and [513, 588] (1911; 15 °C, one measurement, m = 0.50), Haigh [388] (1912; 21 °C, three measurements, m = 0.14), Samollov [590] (1960; 25 °C, one measurements, m = 0.14), Samollov [590] (1960; 25 °C, one measurements). ment, m = 0.05), Rodnikova [594] (1958; 15 and 35 °C, six measurements, m = 0.05), and Samoilov and Buslaeva [444] (1960; 15 and 35 °C, three measurements, m = 0.06). The values of $\Delta H_{\infty}^{\circ}$ are: de Forcrand, 4,300 and 4,320; Haigh, 4,077 \pm 50; Samoĭlov, 4,254; Rodnikova, 4,180 ± 100; Samoĭlov and Buslaeva, $4{,}180 \pm 100 \text{ cal/mole.}$

Beketov and Beketov [527] also measured ΔH_{soin} , using mechanical mixtures of CsCl with LiI, NaI,

"Best" value, $\Delta H_{\infty}^{\circ} = 4,250 \pm 100$ cal/mole.

$CsClO_4(c)$

Pitzer [591] (1938; 25 °C, four measurements, m = 0.002 to 0.009) has measured $\Delta H_{\rm soln}$. Φ_L has been estimated as 15 cal/mole in this concentration range. The value of $\Delta H_{\infty}^{\circ}$ is $13,250 \pm 100$ cal/mole. "Best" value, $\Delta H_{\infty}^{\circ} = 13,250 \pm 100$ cal/mole.

CsBr(c)

 ΔH_{soln} has been measured by de Forcrand [513, 588] (1911; 15 °C, one measurement, m = 0.50) and Lange and Martin [397] (1937; 25 °C, one measurement, m=0.11). The values of $\Delta H_{\infty}^{\circ}$ are 6,420 and 6,210 cal/mole, respectively.

"Best" value, $\Delta \hat{H}_{\infty}^{\circ} = 6,210 \pm 100$ cal/mole.

$CsBrO_3(e)$

 $\Delta H_{\rm soln}$ has been measured by Boyd and Vaslow [522] (1962; 25 °C, five measurements, m = 0.005 to 0.001). Φ_L has been estimated as 20 cal/mole in this concentration range. The value of $\Delta H_{\infty}^{\circ}$ is $12,060 \pm 100 \text{ cal/mole.}$

"Best" value, $\Delta H_{\infty}^{\circ} = 12,060 \pm 100$ cal/mole.

CsI(c)

 $\Delta H_{\rm soln}$ has been measured by Beketov and Beketov [527] (1904; 18 °C, three measurements, m = 0.16 to 0.12) and de Forcrand [513, 588] (1911; 15 °C, one measurement, m = 0.50). Beketov and Beketov's measurements were with mechanical mixtures of CsI with LiCl, NaCl, and KCl. The values of $\Delta H_{\infty}^{\circ}$ are 7,960 ± 150 cal/mole and 7,970 cal/mole, respectively.

"Best" value, $\Delta H_{\infty}^{\circ} = 7,970 \pm 100 \text{ cal/mole.}$

$CsNO_3(c)$

 $\Delta H_{\rm soln}$ has been measured by Haigh [388] (1912; 21 °C, four measurements, m=0.14). The value of $\Delta H_{\infty}^{\circ}$ is 9,560 ±50 cal/mole.

"Best" value, $\Delta H_{\infty}^{\circ} = 9,560 \pm 50$ cal/mole.

VIII. Review Papers

Besides those mentioned in the introduction, there are numerous review and compilation papers on the thermal properties of the aqueous uni-univalent electrolytes. Among these are Eigen and Wicke [625], Latimer [626], Mishchenko [627], Gucker [628], Wüst and Lange [629], Vasil'ev et al. [630], Robinson and Frank [631], and Kapustinskii, Drakin and Yakushevskii [632]. Particular note should be made of Robinson and Stoke's [633] correlation of activity coefficients, conductivities, osmotic coefficients, etc.

No attempt is made here to list all review papers on the thermal properties of specific aqueous uniunivalent electrolytes. The following lists some for the more important compounds.

HF

Elmore, Hatfield, Mason, and Jones [197] have calculated the thermal properties at 25 °C. They include the activity, free energy, enthalpy, entropy, and molal heat capacity. Franck and Spalthoff [611] correlate the specific heat at constant volume with the density and temperature.

HCl

Van Nuys [596] presents charts summarizing the enthalpy and heat of dilution data from 0 to 100 °C.

HNO:

Chédin [597] reviews the properties of aqueous HNO₃, its hydrates, equilibrium equations, and ionization. McCurdy and McKinley [598] correlate the enthalpy-temperature data for the HNO₃-H₂SO₄-H₂O system. Kharbanda [619] provides a monogram which permits the calculation of the specific heat of aqueous solutions from -40 to 192 °C. Bump and Sibbitt [620] correlate the specific heat data of aqueous mixtures of HNO₃ and H₂SO₄.

$C_2H_4O_2$

Lemlich, Gottschlich, and Hoke [595] give an enthalpy-concentration chart for acetic acid-water mixtures.

NH₃

Rozenfel'd [599] gives a temperature-entropy diagram for the aqueous system.

NH₄NO₃

Othmer and Frohlich [600] construct an enthalpyconcentration diagram.

AgNO₃

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=15 to 1.

LiOH

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=5 to 1.

LiCl

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \overline{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=18 to 1.

LiR

Löwer [602] and Rozenfel'd and Karnaukh [603] give isotherms for the enthalpy-concentration of aqueous solutions.

LiNO₂

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \overline{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=20 to 1.

NaOH

Rant [604, 605] gives diagrams for the enthalpy-entropy-concentration-temperature properties. Ginzburg [606, 610] gives the thermal properties of concentrated solutions. Haltenberger [607] gives an enthalpy-concentration chart. Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=27 to 2.

NaCl

Fábry [608] constructs a temperature-enthalpy-concentration chart. Shields [609] reviews the physical and thermal properties of brine solutions.

NaNO₃

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=9 to 1.

KOH

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=21 to 1.

KNO₃

Kangro and Groeneveld [601], using vapor pressure data at 20 and 25 °C, calculate values of \bar{L}_1 at 22.5 °C and the activity coefficients at 25 °C from m=3 to 1.

IX. References

- Rossini, F. D., BS. J. Research 4, 313-327 (1930) RP151.
- Rossini, F. D., BS. J. Research 6, 791-806 (1931) RP305. Rossini, F. D., BS. J. Research 6, 847-856 (1931) RP309.
- [4] Rossini, F. D., BS. J. Research 7, 47-55 (1931) RP331.
 [5] Rossini, F. D., BS. J. Research 9, 679-702 (1932) RP499.
 [6] Bichowsky, F. R., and Rossini, F. D., The thermochemistry of the chemical substances (Reinhold Publ. Corp., New
- York, N.Y., 1936).
 [7] Rossini, F. D., Wagman, D. D., Evans, W. H., Levine, S., and Jaffe, I., Selected values of chemical thermodynamic properties, Natl. Bur. Standards Circ. 500, (U.S. Government Printing Office, Washington, D.C., 20402 1952).

[8] Harned, H. S., and Owen, B. B., The physical chemistry of electrolytic solutions, 2d edition (Reinhold Publ. Corp.,

New York, N.Y., 1950).
[9] Harned, H. S., and Owen, B. B., The physical chemistry of electrolytic solutions, 3d edition (Reinhold Publ. Corp., New York, N.Y., 1958). [10] Lewis, G. N., Randall, M., Pitzer, K. S., and Brewer, L.,

- [10] Lewis, G. N., Randan, M., Titzel, R. S., and Brewer, E., Thermodynamics, 2d edition (McGraw-Hill Book Co., Inc., New York, N.Y., 1961).
 [11] Britske, E. V., Kapustinskii, A. F., Veselovskii, B. K., Shamovskii, L. M., Chentsova, L. G., and Anvaer, B. I., Termicheskie konstanty neorganicheskik veshchesty (Izdatel'stvo Akademii Nauk S.S.S.R., Moskva, Leningrad, 1949).
- [12] Lange, E., and Robinson, A. L., Chem. Revs. 9, 89-116
- (1931). [13] Lange, E., and Monheim, J., Z. Elektrochem. **36**, 772–777 (1930).
- [14] Lewis, G. N., and Randall, M., Thermodynamics and the [14] Lewis, G. N., and Kandali, M., Thermodynamics and the free energy of chemical substances (McGraw-Hill Book Co., Inc., New York, N.Y., 1923).
 [15] Glasstone, S., Thermodynamics for chemists (D. Van Nostrand Co., Inc., New York, N.Y., 1947).
 [16] Cameron, A. E., and Wichers, E., J. Am. Chem. Soc. 84, 4175-4197 (1962).
 [17] Signet H. E. A., J. Phys. 22, 614, 622 (1955).

- [17] Stimson, H. F., Am. J. Phys. 23, 614-622 (1955).
 [18] Natl. Bur. Standards Circ. 459, (U.S. Government Printing
- Office, Washington, D.C., 20402 1947).
 [19] Kelley, K. K., Contributions to the data on theoretical metallurgy. XIII. High temperature heat-content, heat capacity, and entropy data for the elements and inorganic compounds, U.S. Bur. Mines Bull. 584, 232
- pp. (1960).

 [20] Kelley, K. K., and King, E. G., Contributions to the data on theoretical metallurgy. XIV. Entropies of the elements and inorganic compounds, U.S. Bur. Mines
- [21] Wüst, J., and Lange, E., Z. physik. Chem. 116, 161-214 (1925).
- [22] Guggenheim, E. A., and Prue, J. E., Trans. Faraday Soc. **50,** 710-718 (1954).
- 50, 710-718 (1954).
 [23] Baud, E., Ann. chim. et phys. [8] 1, 8-71 (1904).
 [24] Pranschke, A., and Schwiete, H. E., Z. anorg. u. allgem. Chem. 223, 225-233 (1935).
 [25] Kolesov, V. P., Skuratov, S. M., and Uvarov, V. Ya., Zhur. Neorg. Khim. 5, 1934-1937 (1960).
 [26] Thorvaldson, T., and Bailey, E. C., Can. J. Res. 24B, 51-56 (1946).

- (1946). [27] Roth, W. A., and Chall, P., Z. Elektrochem. 34, 185-199
- (1928). [28] Roth, W. A., Pahlke, H., Bertram, A. Elektrochem. 43, 350-355 (1937). A., and Börger, E., Z.
- 29] Mulert, O., Z. anorg. Chem. 75, 198-240 (1912). [30] Thomsen, J., Thermochemische Untersuchungen. Vol. I Neutralisation, Vol. II Metalloide, Vol. III Metalle, Vol. IV Organische Verbindungen (J. Barth Verlag, Leipzig, 1882–1886).
- [31] Marignac, C., Oeuvres Completes II (1860-1887) Bibl. Univ. Archives, t. XXXIX 217-249 (1870).
- [32] Marignac, C., Ann. chim. et phys. [5] **8**, 410–430 (1876). [33] Richards, T. W., and Rowe, A. W., Proc. Am. Acad. Arts Sci. 43, 475-488 (1908).
- [34] Richards, T. W., and Rowe, A. W., Z. physik. Chem. 64, 187-200 (1908).

- [35] Richards, T. W., and Rowe, A. W., Z. physik. Chem. 84,
- [36] Richards, T. W., and Rowe, A. W., Proc. Am. Acad. Arts Sci. 49, 173-199 (1913).
 [37] Richards, T. W., and Rowe, A. W., J. Am. Chem. Soc. 42, 173-199 (1913).
- [31] Incharus, I. W., and Rowe, A. W., J. Am. Gleffi. Soc. 42, 1621-1635 (1920).
 [38] Richards, T. W., Rowe, A. W., and Burgess, L. L., J. Am. Chem. Soc. 32, 1176-1186 (1910).
 [39] Ackermann, T., Z. Elektrochem. 62, 411-419 (1958).
 [40] Vrevskii, M. S., and Kaigorodov, A. N., Zhur. Russ. Fiz.-Village Check Philips Phil

- Khim. Ohshchestva, Chast' Khim. 54, 336-347 (1923).
 [41] Vrevskii, M. S., and Kaigorodov, A. N., Z. physik. Chem. 112, 83-89 (1924).
- [42] Tucker, W. S., Phil. Trans. Roy. Soc. London A215, 319-351 (1915).
- [43] Richards, T. W., Mair, B. J., and Hall, L. P., J. Am. Chem. Soc. 51, 727-730 (1929).
 [44] Thorvaldson, T., Brown, W. G., and Peaker, C. R., J. Am. Chem. Soc. 52, 3927-3936 (1930).
- [45] Drucker, C., Arkiv Kemi, Mineral., Geol. 11A, No. 18, 27 pp. (1935). [46] Radulescu, D., and Jula, O., Z. physik. Chem. **B26**, 390–
- 394 (1934). [47] Wicke, E., Eigen, M., and Ackermann, T., Z. physik.
- Chem. [NF] 1, 340-364 (1954).

 [48] Randall, M., and Ramage, W. D., J. Am. Chem. Soc. 49, 93-100 (1927).
- [49] Gucker, F. T., Jr., and Schminke, K. H., J. Am. Chem. Soc. 54, 1358-1373 (1932).
- [50] Sturtevant, J. M., J. Am. Chem. Soc. 64, 762-768 (1942).
 [51] Bates, R. C., and Bower, V. E., J. Res. NBS 53, 283-290
- (1954) RP2546.
- [52] Hayes, J. C., and Lietzke, M. H., J. Phys. Chem. 64, 374-376 (1960).
- [53] Greeley, R. S., Ph. D. thesis, Univ. of Tennessee, (1959), taken from Dissertation Abstr. 20, 1190 (1959).
- [54] Covington, A. K., and Prue, J. E., J. Chem. Soc. 1955, 3701-3704.
- [55] Covington, A. K., and Prue, J. E., J. Chem. Soc. 1955,
- 3696-3700.
 [56] Oiwa, I. T., J. Phys. Chem. **60**, 754-759 (1956).
 [57] Taylor, J. K., and Smith, E. R., in Katz, J. J., and Rabinowitz, E., eds., U.S. AEC TID **5299**, 429-436 (1958).
- [58] Aston, J. G., and Gittler, F. L., J. Am. Chem. Soc. 77, 3173-3174 (1955).
- [59] Berthelot, M., Ann. chim. et phys. [5] 27, 222-225 (1882).
 [60] Berthelot, M., Compt. rend. 93, 291-292 (1881).
 [61] Tolloczko, S., and Meyer, M., Kosmos 35, 641-648
- (1910).
- [62] Roozeboom, H. W. B., Rec. trav. chim. 5, 323-334 (1886).
 [63] Johnson, W. II., Gilliland, A. A., and Prosen, E. J., J. Res. NBS 63A (Phys. and Chem.) No. 2, 161-166
- [64] Biermann, W. J., and Yamasaki, R. S., J. Am. Chem. Soc. 77, 241 (1955).
- [65] Randall, M., and Taylor, M. D., J. Phys. Chem. 45, 959-967 (1941). [66] Marignac, C., Arch. sci. phys. et nat. [2] 55, 113-135
- (1876).
- [67] Mishchenko, K. P., and Ponomareva, A. M., Zhur. Fiz. Khim. 26, 998-1006 (1952).
 [68] Forsythe, W. R., and Giauque, W. F., J. Am. Chem. Soc.
- 64, 48-61 (1942).
 [69] Roth, W. A., Z. Elektrochem. 50, 107-110 (1944).
 [70] Richards, T. W., and Rowe, A. W., J. Am. Chem. Soc. 43, 770-796 (1921).
 [71] Lloyd, L., and Wyatt, P. A. H., J. Chem. Soc. 1957, 4262-
- [72] Bump, T. R., Pagerey, P. F., Kern, J. P., Fyfe, D. W., St. Clair, C. R., and Sibbitt, W. L., Jet Propulsion 25, 170–172, 180 (1955).
- [73] Booman, K., Elverum, G. W., Jr., and Mason, D. M., J. Phys. Chem. 59, 516-519 (1955).
 [74] Bury, C. R., and Davies, D. G., J. Chem. Soc. 1932, 2413-
- [75] Ackermann, T., and Schreiner, F., Z. Elektrochem. 62, 1143-1151 (1958).
- [76] Glagoleva, A. A., and Cherbov, S. I., Zhur. Obshcheï Khim. 6, 685-690 (1936).

[77] von Reis, M. A., Ann. Physik [3] 13, 447-465 (1881).

[78] Berthelot, M., and Ogier, J., Ann. chim. et phys. [5] 23, 201-209 (1881).

[79] Pettersson, O., J. prakt. Chem. [2] 24, 293-310 (1881).
[80] Gibson, G. E., Latimer, W. M., and Parks, G. S., J. Am. Chem. Soc. 42, 1533-1542 (1920).

[81] Stout, J. W., and Fisher, L. H., J. Chem. Phys. 9, 163-168 (1941).

- [82] Neiman, M. B., Zhur. Obshchei Khim. 2, 1-8 (1932).
 [83] Neiman, M. B., Z. physik. Chem. A158, 258-264 (1932).
 [84] Richards, T. W., and Gucker, F. T., Jr., J. Am. Chem. Soc. 47, 1876-1893 (1925).
- [85] von Reis, M. A., quoted in Roth, W. A., and Scheel, K., eds., Landolt-Bornstein Physikalisch-chemische Tab-ellen, V Aufl., II Ergzb., 1269 (J. Springer Verlag, Berlin, 1923).

- [86] Parks, G. S., and Kelley, K. K., J. Am. Chem. Soc. 47, 2089-2097 (1925).
 [87] Swietoslawski, W., and Zielenkiewicz, A., Bull. acad. polon. sci., Ser. sci., chim., geol. et geograph. 6, 365-466 (1972). 366 (1958).
- [88] Berthelot, M., Ann. chim. et phys. [5] 12, 529-535 (1877).
 [89] Rühlemann, F., Dissertation, Jena, (1908), quoted in Meyer, R. J., and Pietsch, E., Gmelins Handbuch der anorganischen Chemie, 8th Aufl. System No. 23 (Verlag Chem.,

Rerlin, 1936). [90] Hildenbrand, D. L., and Giauque, W. F., J. Am. Chem.

Soc. 75, 2811–2818 (1953).

[91] Labowitz, L. C., and Westrum, E. F., Jr., J. Phys. Chem. **65**, 408-414 (1961).

65, 408-414 (1961).
 [92] Mishchenko, K. P., and Ponomareva, A. M., Zhur. Obshchei Khim. 26, 1296-1310 (1956).
 [93] D'Ans, J., and Tollert, H., Z. Elektrochem. 43, 81-91 (1937).
 [94] Winkelmann, A., Ann. Physik [2] 149 1-32 (1873)
 [95] Faasch, H., Dissertation, Rostock (1911), quoted in Roth, W. A., and Scheel, K., eds., Landolt-Bornstein Physikalisch-chemische Tabellen, V Aufl., II Ergzb., 1262-1263 (J. Springer Verlag, Berlin, 1923).
 [96] Urban, F., J. Phys. Chem. 36, 1108-1122 (1932).
 [97] Tollinger, J., Sitzber. Akad. Wiss. Wien., Math.-Nat. Kl. II, 72, 535-577 (1875).
 [98] Zwicky, F., Physik, Z. 27, 271-286 (1926).

[98] Zwicky, F., Physik. Z. 27, 271–286 (1926).
[99] Lerner-Steinberg, B., Z. physik. Chem. 122, 121–125

- [100] Chauvenet, R., Compt. rend. 207, 1216-1218 (1938).
 [101] Rutskov, A. P., Sbornik Nauch.-Issledovatel'. Rabot. Arkhangel'sk. Lesotekh. Inst. im. V. V. Kuibysheva 8, 85-94 (1946).
- [102] Rutskov, A. P., Zhur. Priklad. Khim. 21, 820-823 (1948).
 [103] Cohen, E., Helderman, W. D., and Moesveld, A. L. T., Z. physik. Chem. 112, 135-140 (1924).
 [104] Cohen, E., Helderman, W. D., and Moesveld, A.L.T., Koninkl. Ned. Akad. Wetenschap. Proc. 27, 565-571
- [105] Gucker, F. T., Jr., Ayres, F. D., and Rubin, T. R., J. Am.
- [105] Gucker, F. 1., Jr., Ayres, F. D., and Rubin, T. R., J. Am. Chem. Soc. **58**, 2118–2126 (1936).
 [106] Gucker, F. T., Jr., Ayres, F. D., and Rubin, T. R., J. Am. Chem. Soc. **58**, 2661 (1936).
 [107] Whittow, E. P., and Felsing, W. A., J. Am. Chem. Soc. **66**,
- 2028-2033 (1944).
- [108] Aston, J. G., and Ziemer, C. W., J. Am. Chem. Soc. 68, 1405-1413 (1946).
- [109] Coulter, L. V., Pitzer, K. S., and Latimer, W. M., J. Am.
- Chem. Soc. **62**, 2845–2851 (1940). [110] Jahn-Held, W., and Jellinek, K., Z. Elektrochem. **42**, 608– 623 (1936).
- [111] Kapustinskii, A. F., Yakushevskii, B. M., and Drakin, S. I., Zhur. Fiz. Khim. 27, 588-595 (1953).
 [112] Gucker, F. T., Jr., and Schminke, K. H., J. Am. Chem. Soc. 55, 1013-1019 (1933).
- 113] Chernyak, E. L., Zhur. Obshchei Khim. 8, 1341-1352 (1938) [114] Bennewitz, K., and Kratz, L., Physik. Z. 37, 496-511 (1936).
- [115] White, C. M., J. Am. Chem. Soc. **58**, 1615–1620 (1936). [116] Jauch, K., Z. Physik **4**, 441–447 (1921). [117] Hess, C. B., J. Phys. Chem. **45**, 755–761 (1941).

[118] Lange, E., and Dürr, F., Z. physik. Chem. 121, 361-384 (1926).

- [119] Lange, E., and Schwartz, E., Z. physik. Chem. 133, 129-150 (1928).
- [120] Eigen, M., and Wicke, E., Z. Elektrochem. 55, 354-363 (1951).
- [121] Bertetti, J. W., and McCabe, W. L., Ind. Eng. Chem. 28, 375-378 (1936).
- [122] Ackermann, T., Discussions Faraday Soc. No. 24, 180-193
- [123] Richards, T. W., and Hall, L. P., J. Am. Chem. Soc. 51, 707-712 (1929). [124] Roth, W. A., Wirths, G., and Berendt, H., Z. Elektrochem.
- 48, 264-267 (1942)
- [125] Wilson, H. R., and McCabe, W. L., Ind. Eng. Chem. 34, 558-566 (1942)
- [126] Akerlöf, G., and Kegeles, G., J. Am. Chem. Soc. 62, 620-640 (1940).
- [127] Stakhanova, M. S., and Vasilev, V. A., Izvest. Vysshikh. Ucheb. Zavedenii, Khim. i Khim. Tekhnol. 3, 829-833 (1960).
- [128] Biron, E., Zhur. Russ. Fiz.-Khim. Obshchestva 30, 355-366 (1898).
- [129] Blaszkowska, H., Roczniki Chem. 4, 120–148 (1924).
- 130] Zdanovski i, A. B., Zhur. Fiz. Khim. 12, 106-108 (1938). [131] Zdanovskii, A. B., and Suslina, K. D., Zhur. Fiz. Khim. 12, 109-112 (1938).
- [132] Zdanovskii, A. B., an 11, 861-863 (1938). and Matsenok, E. A., Zhur. Fiz. Khim.
- [133] Chipman, H. R., Johnson, F. M. G., and Maass, O., Proc. Trans. Nova Scotian Inst. Sci. 17, pt. 3, 149-167 (1929).
- Trans. Nova Scottan Inst. Sci. 14, pt. 3, 147-101 (1227).

 [134] Nikolaev, V. I., Kogan, A. G., and Ogorodnikov, G. F., Levest. Akad. Nauk S.S.S.R., Otdel. Mat. i Estestven. Nauk, Ser. Khim. 1936, 193-214.

 [135] Leitner, I. A., Arch. Wärmewirtsch. u. Dampfkesselw. 9, 233-242 (1928).

[136] White, C. M., J. Phys. Chem. 44, 494-512 (1940).

- [137] Hess, C. B., and Gramkee, B. E., J. Phys. Chem. 44, 483-494 (1940).
- [138] Voskresenskaya, N. K., and Yankovskaya, G. N., Izvest. Akad. Nauk S.S.S.R., Otdel. Khim. Nauk No. 1, 3-13 (1945).
- [139] Lipsett, S. G., Johnson, F. M. G., and Maass, O., J. Am. Chem. Soc. 50, 2701-2703 (1928).
 [140] Bousfield, W. R., and Bousefield, C. E., Phil. Trans. Roy. Soc. London A218, 119-156 (1919).
- [141] Sandonnini, C., and Gerosa, G., Gazz. chim. ital. 55, 916-938 (1925).
- [142] Person, C. C., Ann. chim. et phys. [3] 33, 437-448 (1851). [143] Randall, M., and Bisson, C. S., J. Am. Chem. Soc. 42, 347-367 (1920).
- [144] Randall, M., and Rossini, F. D., J. Am. Chem. Soc. 51, 323-345 (1929).
- [145] Colomina, M., and Nicolas, J., Anales real. soc. espan. fis. y quim. (Madrid) 45B, 137-170 (1949).

- [146] Ferrer, P., Thesis, Madrid (1950).
 [147] Bender, P., and Kaiser, A. D., Jr., J. Am. Chem. Soc. 76, 3084-3085 (1954).
- Rossini, F. D., Ph. D. Thesis, Univ. of California, (1928). Perreu, J., Compt. rend. **212**, 442-444 (1941). Magie, W. F., Phys. Rev. [1] **25**, 171-183 (1907). Winkelmann, A., Ann. Physik [2] **149**, 492-521 (1873). Gerlach, H., Caliche **8**, 298 (1926).

- [153] Perreu, J., Compt. rend. 211, 256-258 (1940).
 [154] Bindel, K., Ann. Physik [3] 40, 370-398 (1890).
 [155] Kapustinskii, A. F., and Stakhanova, M. S., Izvest. Vysshikh Ucheb. Zavedenii Khim. i Khim. Tekhnol. 1958, No. 2, 7, 14. No. 3, 7-14.
- [156] Gnesotto, T., and Fabris, C., Atti reale ist. veneto, lettere ed arti 70II, 471-481 (1911).
 [157] Richards, T. W., and Gucker, F. T., Jr., J. Am. Chem. Soc. 51, 712-727 (1929).

- [158] Jaquerod, A., J. chim. phys. 7, 129-151 (1909).
 [159] Gawlik, H., Z. Ver. deut. Ing. 79, 1089 (1935).
 [160] Voskresenskaya, N. K., Izvest. Sektora Fiz. Khim. Anal., Inst. Obshchei i Neorg. Khim., Akad. Nauk S.S.S.R. 10, 259-273 (1938).
- [161] Sabatier, P., Ann. chim. et phys. [5] 22, 5-98 (1881).
 [162] Andrews, T., Ann. chim. et phys. [3] 14, 68-92 (1845).

- [163] Voskresenskaya, N. K., and Rutskov, A. P., Izvest. Akad. Nauk S.S.S.R., Otdel. Khim. Nauk 1940, 795-810.
 [164] Cobb, A. W., and Gilbert, E. C., J. Am. Chem. Soc. 57, 35-39 (1935).
- [165] Popov, M. M., Bundel, A., and Choller, V., Z. physik. Chem. A147, 302-318 (1930).
- [166] Popov, M. M., Khomyakov, K. G., Feodos'ev, N. N., and Schirokich, P. K., Z. physik. Chem. A167, 29-34 (1933).
- [167] Clews, C. J. B., Proc. Phys. Soc. 48, 95-99 (1936).
 [168] Gucker, F. T., Jr., J. Am. Chem. Soc. 50, 1005-1016
- [169] Varasova, E. N., Mishchenko, K. P., and Frost, O. I., Zhur. Obshcheī Khim. 7, 1284-1291 (1937).
- [170] Cohen, E., and Moesveld, A. L. T., Z. physik. Chem. 95, 305-327 (1920).
- [171] Parshikov, I. A., Zhur. Priklad. Khim. 34, 2649-2658 (1961).
- [172] Popov, M. M., Skuratov, S. M., and Strel'tsova, M. M., Zhur. Obshcheĭ Khim. 10, 2023-2027 (1940).
- [173] Lange, E., and Monheim, J., Z. physik. Chem. A150, 349-369 (1980).

- 349-369 (1980).
 [174] Perreu, J., Compt. rend. 211, 182-185 (1940).
 [175] Kapustinskii, A. F., and Klokman, V. R., Izvest. Akad. Nauk. S.S.S.R., Otdel. Khim. Nauk 1943, 259-263.
 [176] Kapustinskii, A. F., Yakushevskii, B. M., and Drakin, S. I., Zhur. Fiz. Khim. 27, 793-798 (1953).
- S. I., Zhur. Fiz. Khuin. 21, 190-190 (1905).
 Kapustinskii, A. F., Lipilina, I. I., and Samoilov, O. Ya., J. chim. phys. 54, 343-347 (1957).
 Kapustinskii, A. F., Lipilina, I. I., and Samoilov, O. Ya., Zhur. Fiz. Khim. 30, 896-900 (1956).
 Kapustinskii, A. F., Acta Chim. Acad. Sci. Hung. 15, 2023 (1978).
- 351-364 (1958).
- [180] Young, T. F., and Groenier, W. L., J. Am. Chem. Soc. 58, 187-191 (1936).
- [181] Young, T. F., and Machin, J. S., J. Am. Chem. Soc. 58, 2254-2260 (1936).
- [182] Young, T. F., and Seligmann, P., J. Am. Chem. Soc. **60**, 2379–2383 (1938).
- [183] Young, T. F., and Vogel, O. G., J. Am. Chem. Soc. 54, 3030-3040 (1932).
- [184] Guggenheim, E. A., and Prue, J. E., Trans. Faraday Soc. 50, 231-236 (1954).

- 30, 251-250 (1954).

 [185] Petersen, E., Z. physik. Chem. 11, 174-184 (1893).

 [186] Roth, W. A., Ann. 542, 35-43 (1939).

 [187] Guntz, A., Ann. chim. et phys. [6] 3, 5-66 (1884).

 [188] Schäfer, H., and Kahlenberg, F., Z. anorg. u. allgem.

 Chem. 305, 291-326 (1960).
- Chem. 305, 291–326 (1960).

 [189] Schäfer, H., and Kahlenberg, F., Z. anorg. u. allgem. Chem. 294, 242–253 (1958).

 [190] Broene, H. H., and De Vries, T., J. Am. Chem. Soc. 69, 1644–1646 (1947).

 [191] Wooster, C. B., J. Am. Chem. Soc. 60, 1609–1613 (1938).

 [192] von Steinwehr, H., Z. physik. Chem. 38, 185–199 (1901).

 [193] Petersen, E., Z. physik. Chem. 4, 384–412 (1889).

 [194] Arrhenius, S., Z. physik. Chem. 4, 96–116 (1889).

 [195] Roth, W. A., and Becker, G., quoted in Roth, W. A., and Scheel, K., eds., Landolt-Bornstein Physikalisch-chem.

- Scheel, K., eds., Landolt-Bornstein Physikalisch-chemische Tabellen, V. Aufl., II Ergzb., 1524 (J. Springer Verlag, Berlin, 1923).
- [196] Berthelot, M., and Guntz, A., Ann. chim. et phys. [6] 3, 362-368 (1884).
- [197] Elmore, K. L., Hatfield, J. D., Mason, C. M., and Jones, A. D., J. Am. Chem. Soc. 71, 2710-2714 (1949).
- [198] Berthelot, M., Ann. chim. et phys. [5] 4, 467-488 (1875).
 [199] Naude, S. M., Z. physik. Chem. 135, 209-236 (1928).
 [200] Becker, G., and Roth, W. A., Z. physik. Chem. A167, 1-15 (1933).
- [201] Muller, J. A., Bull. soc. chim. France [4] 13, 1053-1056 (1913).
- [202] Vrevskii, M. S., and Zavaritskii, N., Z. physik. Chem 112, 90-96 (1924).
- [203] Myers, R. A., Ph. D. thesis, Univ. of Nebraska, (1958). [204] Vanderzee, C. E., and Myers, R. A., J. Phys. Chem. 65, 153-158 (1961).
- [205] Sacconi, L., Paoletti, P., and Ciampolini, M., Ricerca sci. **29**, 2412–2419 (1959).
- [206] Pitzer, K. S., J. Am. Chem. Soc. 59, 2365-2371 (1937).
- [207] Sturtevant, J. M., J. Am. Chem. Soc. 62, 584-587 (1940).

- [208] Sturtevant, J. M., J. Am. Chem. Soc. 62, 3265-3266 (1940)
- [209] Ellis, J. H., J. Am. Chem. Soc. 38, 737-762 (1916).
- [210] Roth, W. A., and Wienert, F., Arch. Eisenhüttenw. 7, 455-460 (1934).
- [211] Schäfer, H., Wittig, F. E., and Jori, M., Z. anorg. u. allgem. Chem. 287, 61-70 (1956).
- [212] Payn, R. C., and Perman, E. P., Trans. Faraday Soc. 25. 599-610 (1929)

- [213] Bender, P., and Biermann, W. J., J. Am. Chem. Soc. 74, 322-325 (1952).
 [214] Vrevskii, M. S., and Zavaritskii, N. N., Zhur. Russ. Fiz.-Khim. Obshchestva, Chast' Khim. 54, 348-360 (1923).
 [215] Berthelot, M., Ann. chim. et phys. [5] 27, 214-222 (1882).
 [216] Berthelot, M., Sur la force des matières explosives d'apres la thermoshimia. Traisième adition. Tomps I et al. la thermochimie. Troisième (Gauthier-Villars, Paris, 1883). Troisième edition, Tome I et II,
- [217] Krivtsov, N. V., Rosolovskii, V. Ya., and Zinov'ev, A. A., Zhur. Neorg. Khim. 5, 772-774 (1960).
 [218] Wood, R. H., U.S. AEC UCRL 3751, 56 pp. (Apr. 9,
- [219] Austin, J. M., and Mair, A. D., J. Phys. Chem. 66, 519-521 (1962).
- [220] Berthelot, M., Compt. rend. 93, 240-246 (1881).
- [221] Vanderzee, C. E., and Swanson, J. A., J. Phys. Chem. 67, 285-291 (1963).
- [222] Louguinine, W., and Schukareff, A., Arch. sci. phys. et nat. [4] 15, 49-77 (1903).
 [223] Biermann, W. J., and Weber, N., J. Am. Chem. Soc. 76, 4289-4291 (1954).
 [224] Richards, T. W., and Rowe, A. W., J. Am. Chem. Soc. 44, 684-707 (1922).

- [225] Berthelot, M., Ann. chim. et phys. [5] 4, 21-46 (1875). [226] Berthelot, M., Ann. chim. et phys. [5] 4, 445-60 (1875). [227] Berthelot, M., Compt. rend. 78, 769-777 (1874). [228] Becker, G., and Roth, W. A., Z. physik. Chem. A174, 104-114 (1935).
- [229] Good, W. D., Scott, D. W., Lacina, J. L., and McCullough, J. P., J. Phys. Chem. 63, 1139-1142 (1959).
 [230] Wilson, G. L., and Miles, F. D., Trans. Faraday Soc. 36,
- 356-363 (1940).
- [231] Roth, W. A., J. prakt. Chem. [2] 158, 117-124 (1941).
 [232] Chédin, J., and Dhers, J., Mem. serv. chim. état (Paris) 38, 381-387 (1953).
- [233] Berthelot, M., Ann. chim. et phys. [5] 6, 334–352 (1875). [234] Faucon, A., Ann. chim. et phys. [8] 19, 70–152 (1910). [235] Glagoleva, A. A., Zhur. Obshchei Khim. 6, 1769-1777
- $(1\overline{9}36).$ [236] Everett, D. H., and Wynne-Jones, W. F. K., Trans. Faraday
- Soc. 35, 1380-1401 (1939).
 [237] Canady, W. J., Papée, H. M., and Laidler, K. J., Trans. Faraday Soc. 54, 502-506 (1958).
 [238] Berthelot, M., Ann. chim. et phys. [5] 4, 8-20 (1875).
 [239] Berthelot, M., Compt. rend. 80, 700-707 (1875).

- [240] Klibanova, Ts. M., and Neiman, M. B., Zhur. Fiz. Khim. 4,
- [241] Richards, T. W., and Mair, B. J., J. Am. Chem. Soc. 51, 737-740 (1929).

- [242] Berthelot, M., Ann. chim. et phys. [5] 6, 325–334 (1875).
 [243] Lundén, H., J. chim. phys. 6, 681–698 (1908).
 [244] Devoto, G., Atti accad. nazl. Lincei Rend., Classe sci. fis.
 mat. e nat. 19, 50–52 (1934).
- [245] Jordan, J., and Dumbaugh, W. H., Jr., Bull. Chem. Thermodynam. No. 2, 9-11 (1959).
 [246] Craham, T., Phil. Mag. [3] 24, 401-420 (1844).
 [247] Favre, P. A., Compt. rend. 73, 767-778 (1871).
 [248] Andrews, T., J. Chem. Soc. 23, 432-441 (1870).
 [240] Retablet, M. Annehime et phys. [5] 4, 526-531 (1875).

- [249] Berthelot, M., Ann. chim. et phys. [5] 4, 526-531 (1875).
 [250] Calvet, E., J. chim. phys. 30, 140-166 (1933).
 [251] Ramstetter, H., and Hantke, G., Z. physik. Chem. Bodenstein Festband 662-668 (1931).
- [252] Baud, E., and Gay, L., Ann. chim. et phys. [8] 17, 398-418 (1909).
- [253] Berthelot, M., Compt. rend. 73, 745-749 (1871).
- [254] Bruni, G., and Levi, G., Gazz. chim. ital. 47, 259-272 (1917). [255] Berthelot, M., Compt. rend. 80, 592-598 (1875).
- [256] Berthelot, M., Ann. chim. et phys. [4] 29, 328-351 (1873).
- [257] Berthelot, M., Ann. chim. et phys. [4] 29, 433-514 (1873).

[258] Biltz, W., and Messerknecht, C., Z. anorg. u. allgem. Chem. 148, 157–169 (1925). [259] Hess, H., Ann. Physik [2] 52, 97–114 (1841).

- [260] Bouzat, A., Ann. chim. et phys. [7] 29, 305-383 (1903).
- [261] Vorob'ev, A. F., Privalova, N. M., Monaenkova, A. S., and Skuratov, S. M., Doklady Akad. Nauk S.S.S.R. 135, 1388-1390 (1960).
- [262] Favre, P. A., and Silbermann, J. T., Ann. chim. et phys. [3] 37, 406–508 (1853).
- 57, 400-508 (1855).

 [263] Andrews, T., Ann. chim. et phys. [3] 4, 316-329 (1842).

 [264] Delépine, M., Ann. chim. et phys. [7] 15, 469-574 (1898).

 [265] Moore, F. S., and Winmill, T. F., J. Chem. Soc. 101, 1635-1676 (1912).

[266] Pollok, J. H., Chem. News 79, 56 (1899).

- [267] Mondain-Monval, P., Ann. chim. (Paris) [10] 3, 72-120 (1925)

- [268] Mondain-Monval, P., Compt. rend. 176, 889-892 (1923). [269] Pratt, F. R., J. Franklin Inst. 185, 663-695 (1918). [270] Magie, W. F., Phys. Rev. [1] 35, 265-271 (1912). [271] Fedorov, A. S., and Sil'chenko, G. F., Ukrain. Khim. Zhur.
- 12, 53-60 (1937).
 [272] Streeck, H., Z. physik. Chem. A169, 103-112 (1934).
 [273] Dunnington, F. P., and Hoggard, T., Am. Chem. J. 22, 207-211 (1899).

- [274] Rümelin, G., Z. physik. Chem. 58, 449–466 (1907).
 [275] von Stackelberg, E., Z. physik. Chem. 26, 533–563 (1898).
 [276] Mondain-Monval, P., Compt. rend. 176, 301–305 (1923).
- [277] Wassermann, A., Z. physik. Chem. A146, 409-417 (1930).
 [278] von Wartenberg, H., and Lerner-Steinberg, B., Z. physik. Chem. 122, 113-120 (1926).
 [279] Fricke, R., and Havestadt, L., Z. Elektrochem. 33, 441-455
- (1927).
- [280] Bonnefoi, J., Ann. chim. et phys. [7] 23, 317-378 (1901).
 [281] Felsing, W. A., and Wohlford, P. H., J. Am. Chem. Soc. 54, 1442-1445 (1932).
- [282] Patterson, A., Jr., and Felsing, W. A., J. Am. Chem. Soc. **60**, 2693–2695 (1938).
- [283] Connick, R. E., and Paul, A. D., J. Phys. Chem. 65, 1216-1220 (1961).
- [284] Paul, A. D., U.S. AEC UCRL **2926**, 163 pp. (Apr. 1955).
 [285] Leden, I., and Marthen, L. E., Acta Chem. Scand. **6**, 1125 (1952).
- [286] von Wartenberg, H., Z. Elektrochem. 53, 343-346 (1949).
 [287] Jackson, N. S., Smith, A. E. C., Gatty, O., and Wolfenden, J. H., J. Chem. Soc. 1934, 1376-1379.
 [288] Terasaki, Y., Nippon Kagaku Zasshi 78, 1777-1780 (1957).
 [289] Ueda, Y., Sci. Repts. Tohoku Imp. Univ. [1] 22, 448-474 (1932).
- (1933). [290] Ueda, Y., Nippon Kagaku Zasshi **52**, 740–755 (1931).
- [291] Juza, R., and Laurer, P., Z. anorg. u. allgem. Chem. 275, 79-93 (1954).
- [292] Leighton, P. A., Unpublished measurements, quoted in Lange, E., and Monheim, J., Z. Elektrochem. 36, 772-777 (1930).
- [293] Lange, E., and Leighton, P. A., quoted in Lange, E., and Robinson, A. L., Chem. Revs. 9, 89-116 (1931).
 [294] Nernst, W., and Orthmann, W., Z. physik. Chem. 135, 199-208 (1928).
- [295] Nernst, W., and Orthmann, W., Sitzber. preuss. Akad.
- Wiss., Physik. math. Kl. 1927, 136–141. [296] Lemoine, G., Compt. rend. 125, 603–605 (1897). [297] Lange, E., and Messner, G., Naturwiss. 15, 521–522 (1927).
- [298] Lange, E., and Messner, G., Z. Elektrochem. 33, 431-440 (1927).
- [299] Birnthaler, W., and Lange, E., Z. Elektrochem. 44, 679-693 (1938).
- [300] Monheim, J., and Streeck, H., unpublished measurements. quoted in Lange, E., and Meixner, J., Physik. Z. 30, 670-678 (1929).
- [301] Johnson, E. F., Jr., and Molstad, M. C., J. Phys. Chem. 55,
- 257-281 (1951).
 [302] Jones, F. R., Ph. D. thesis, Univ. of Delaware, (1961).
 [303] Mishchenko, K. P., Z. Elektrochem. 36, 777-783 (1930).
 [304] Lange, E., and Mishchenko, K. P., Z. physik. Chem. A148, 161-176 (1930).
- [305] Pennington, W., Refrig. Eng. 63, No. 5, 57-61 (1955).[306] Dodé, M., Compt. rend. 203, 365-367 (1936).

- [307] Berthelot, M., Ann. chim. et phys. [5] 4, 513-525 (1875).
 [308] Fricke, R., Z. Elektrochem. 35, 631-640 (1929).
 [309] Sturtevant, J. M., J. Am. Chem. Soc. 62, 2276-2280 (1940).
 [310] Sabatier, P., Compt. rend. 89, 234-236 (1879).
 [311] Bertetti, J. W., and McCabe, W. L., Ind. Eng. Chem. 28, 237-240 (1936). 247-248 (1936).
- [312] de Forcrand, R., Compt. rend. **149**, 1341-1344 (1909). [313] de Forcrand, R., Ann. chim. et phys. [6] **20**, 433-447 (1890).
- [314] Richards, T. W., and Hall, L. P., J. Am. Chem. Soc. 51,

- [314] Richards, T. W., and Hall, L. P., J. Am. Chem. Soc. 51, 731-736 (1929).
 [315] Hayward, A. M., and Perman, E. P., Trans. Faraday Soc. 27, 59-69 (1931).
 [316] Scholz, R., Ann. Physik [3] 45, 193-206 (1892).
 [317] Backlund, J., Öfvers, Finska Vetens.-Soc. Förh. 54, No. 22, 1-18 (1911).
 [318] van Deventer, C. M., and van de Stadt, H. J., Z. physik. Chem. 9, 43-56 (1892).
 [319] Naude, S. M., Z. Elektrochem. 33, 532-534 (1927).
 [320] Malcolm, G. N. Parton, H. N. and Watson, I. D. I.

- [320] Malcolm, G. N., Parton, H. N., and Watson, I. D., J. Phys. Chem. 65, 1900–1902 (1961).
 [321] Robinson, A. L., J. Am. Chem. Soc. 54, 1311–1318 (1932).
 [322] Gulbransen, E. A., and Robinson, A. L., J. Am. Chem. Soc.
- **56**, 2637-2641 (1934).
- [323] Lipsett, S. G., Johnson, F. M. G. and Maass, O., J. Am. Chem. Soc. 49, 925-943 (1927).
 [324] Benson, G. C., and Benson, G. W., Rev. Sci. Instr. 26, 477-481 (1955).
- [325] Lange, E., and Robinson, A. L., Z. physik. Chem. A148, 97-111 (1930).
- [326] Lange, E., and Robinson, A. L., unpublished measure-
- ments, quoted in Lange, E., and Robinson, A. L., Chem. Revs. 9, 89-116 (1931).

 [327] Lange, E., and Robinson, A. L., unpublished measurements, quoted in Lange, E., and Monheim, J., Z. Elektrochem. **36,** 772–777 (1930).
- [328] Wallace, W. E., J. Am. Chem. Soc. 71, 2485-2487 (1949).

- [329] Hammerschmid, H., and Robinson, A. L., J. Am. Chem. Soc. 54, 3120-3125 (1932).
 [330] Pollok, J. H., Proc. Chem. Soc. 15, 8-9 (1899).
 [331] Bishop, F. L., Phys. Rev. [1] 26, 169-176 (1908).
 [332] Stahlberg, A., Öfvers. Finska Vetens.-Soc. Förh. 57, No. 17, 1-22 (1914).
 [333] Pollok, M. Angelsing at allow [41 20, 145 204 (1972)].
- [333] Berthelot, M., Ann. chim. et phys. [4] **30**, 145-204 (1873). [334] Partington, J. R., and Soper, W. E., Phil. Mag. [7] **7**, 209-247 (1929). [335] de Forcrand, R., Compt. rend. **152**, 1073-1077 (1911).
- Guntz, A., Compt. rend 98, 428-431 (1884).
- [337] Birnthaler, W., and Lange, E., Z. Elektrochem. 43, 643-659 (1937).
- [338] Lange, E., and Eichler, A., Z. physik. Chem. 129, 285-
- 300 (1927). [339] Messner, G., Dissertation, München, (1928), quoted in Lange, E., and Monheim, J., Z. Elektrochem. 33, 772-777 (1930).
- [340] Lange, E., and Messner, G., unpublished measurements, quoted in Lange, E., and Eichler, A., Z. physik. Chem. 129, 285-300 (1927).
- [341] Roth, W. A., and Zeumer, H., Z. Elektrochem. 38, 248 (1932).
- [342] Biermann, W. J., Can. J. Chem. 38, 57-60 (1960).
- [343] Stearn. A. E., and Smith, G. M., J. Am. Chem. Soc. 42, .18-32 (1920).
- [344] Nernst, W., and Orthmann, W., Sitzber. preuss. Akad. Wiss., Physik.-math. Kl. 1926, 51-56.
- [345] Lange, E., and Leighton, P. A., Z. Elektrochem. 34, 566-571 (1928).
- [346] Harrison, W. R., and Perman, E. P., Trans. Faraday Soc. 23, 1-22 (1927).
- [347] Andauer, M., and Lange, E., Z. physik. Chem. A165, 89-96 (1933).
 [348] Berthelot, M., Ann. chim. et phys. [5] 29, 201-231 (1883).
 [349] Berthelot, M., Ann. chim. et phys. [5] 13, 20-29 (1878).
 [350] Colson, A., Compt. rend. 161, 458-461 (1915).
 [351] Berenger-Calvet, Mme., J. chim. phys. 24, 325-345 (1927).
 [352] Craham, T. Phil Mos. [21, 22, 239, 250 (1942).

- [352] Graham, T., Phil. Mag. [3] **22**, 329–350 (1843). [353] Hunter, J. B., and Bliss, H., Ind. Eng. Chem. **36**, 945–953 (1944).

[354] de Forcrand, R., Compt. rend. 143, 98-101 (1906).
[355] Lange, E., and Kohler, M., unpublished measurements, quoted in Lange, E., and Messner, G., Z. Elektrochem. **33,** 431–440 (1927).

[356] Gillespie, L. J., Lambert, R. H., and Gibson, J., Jr., J. Am. Chem. Soc. 52, 3806–3813 (1930).
[357] Vanderzee, C. E., and Swanson, J. A., J. Phys. Chem. 67,

2608-2612 (1963).

[358] Papée, H. M., Canady, W. J., and Laidler, K. J., Can. J. Chem. 34, 1677–1682 (1956).

[359] Bidinosti, D. R., and Biermann, W. J., Can. J. Chem. 34, 1591–1595 (1956).

[360] Berthelot, M. and Louguinine, W., Compt. rend. 81, 909-915 (1875).

[361] Hess, H., Ann. Physik [2] **50**, 385–404 (1840).
[362] Mathews, J. H., J. Am. Chem. Soc. **33**, 1291–1309 (1911).
[363] Mathews, J. H., and Germann, A. F. O., J. Phys. Chem. **15**,

[364] Ellingson, E. O., J. Am. Chem. Soc. 37, 699-709 (1915).
 [365] Andrews, T., Brit. Assoc. Advance. Sci. Rept. 19, 63-78

(1849).

[366] Könnecke, H. G., Steinert, H., and Leibnitz, E., Z. physik. Chem. 208, 147-156 (1957).
[367] Eley, D. D., and Watts, H., J. Chem. Soc. 1954, 1319-

[368] Davies, T., Singer, S. S., and Stavely, L. A. K., J. Chem. Soc. 1954, 2304–2311.
 [369] Schultz, J. L., Ph. D. thesis, Univ. of Minnesota, (1958).

[370] Keyes, F. G., Gillespie, L. J., and Mitsukuri, S., J. Am. Chem. Soc. 44, 707-717 (1922).
[371] Douglas, T. B., J. Am. Chem. Soc. 68, 1072-1076 (1946).
[372] Muller, J. A., Ann. chim. et phys. [6] 15, 517-533 (1888).
[373] Lindner, J., Monatsh. 33, 613-672 (1912).
[374] Sendente J. P. and Statt F. Trage Foreday, Soc. 34.

[374] Speakman, J. B., and Stott, E., Trans. Faraday Soc. 34, 1203-1209 (1938).
[375] Olcott, J. S., Ph. D. thesis, Purdue University, (1957), taken from Dissertation Abstr. 18, 838-839 (1958).
[376] Muller, J. A., Bull. soc. chim. France [4] 23, 8-13 (1918).
[377] Berthelot, M., Ann. chim. et phys. [5] 4, 74-131 (1875).
[378] Berthelot, M., Ann. chim. et phys. [5] 29, 234-239 (1883).
[370] Perthelot, M. Ann. chim. et phys. [5] 29, 234-239 (1883).

[379] Berthelot, M., Ann. chim. et phys. [5] **4**, 500-512 (1875). [380] Hess, H., Ann. chim. et phys. [2] **75**, 80-103 (1840). [381] von Rechenberg, C., J. prakt. Chem. [2] **19**, 143-145

(1879).

[382] von Rechenberg, C., J. prakt. Chem. [2] 22, 1-45 (1880).

[383] Berthelot, M., and Ilosvay, L., Ann. chim. et phys. [5] 29, 295-342 (1883).

[384] Pickering, S. U., J. Chem. Soc. **51**, 290–356 (1887). [385] Brönsted, J. N., Z. physik. Chem. **56**, 645–685 (1906). [386] Cohen, E., Helderman, W. D., and Moesveld, A. L. T., Z. physik. Chem. **96**, 259–274 (1920).

[387] Mees, R., Unpublished measurements, quoted in Cohen, E., Helderman, W. D., and Moesveld, A. L. T., Z. physik. Chem. 96, 259-274 (1920).

[388] Haigh, F. L., J. Am. Chem. Soc. 34, 1137-1159 (1912). [389] Lange, E., and Dürr, F., Z. Elektrochem. 32, 85-87 (1926). [390] Holluta, J., and Werner, H., Z. physik. Chem. 129, 262–284

(1927)[391] Cohen, E., and Kooy, J., Z. physik. Chem. A139, 273-350

(1928). [392] Roth, W. A., and Eymann, C., Z. physik. Chem. A143, 321-353 (1929).

[393] Lange, E., and Rounsefell, E. O., Unpublished measurements, quoted in Mishchenko, K. P., Z. Elektrochem. 36, 777-783 (1930).

[394] Moles, E., and Perez-Vitoria, A., Anales soc. espan. fis. y quim. 30, 99-119 (1932).

[395] Smeets, C., Natuurw. Tijdschr. Belg. 15, 105-124 (1933). [396] Shibata, Z., and Terasaki, Y., Nippon Kagaku Zasshi 57, 1208-1211 (1936).

[397] Lange, E., and Martin, W., Z. physik. Chem. A180, 233-245 (1937).

245 (1937).
[398] Fontell, N., Soc. Sci. Fennica, Commentationes Phy. Math. 10, No. 5, 18 pp. (1938).
[399] Slansky, C. M., J. Am. Chem. Soc. 62, 2430-2434 (1940).
[400] Voskresenskaya, N. K., and Ponomareva, K. S., Zhur. Fiz. Khim. 20, 433-440 (1946).
[401] Tichelaar, G. W., Ph. D. thesis, Delft, Holland, (1946).

[402] Foz, O. R., and Colomina, M., Anales fis. y quim. (Madrid)

42, 129-161 (1946).
[403] Colomina, M., thesis, Univ. of Valencia, Valencia, Spain, (1947)

[404] Fineman, M. A., and Wallace, W. E., J. Am. Chem. Soc. 70.

4165–4169 (1948). [405] Mishchenko, K. P., and Kaganovich, Yu. Ya., Zhur. Priklad. Khim. **22**, 1078–1082 (1949).

[406] Barieau, R. E., and Giauque, W. F., J. Am. Chem. Soc. 72, 5676-5684 (1950).

[407] Samoilov, O. Ya., Doklady Akad. Nauk S.S.S.R. 81, 641-

644 (1951). [408] Samoilov, O. Ya., Izvest. Akad. Nauk S.S.S.R., Otdel. Khim. Nauk 1952, 398-405.

[409] Kaganovich, Yu. Ya., and Mishchenko, K. P., Doklady Akad. Nauk S.S.S.R. 87, 89-92 (1952).
[410] Evans, D. F., and Richards, R. E., J. Chem. Soc. 1952,

3932-3936. [411] Kapustinskii, A. F., and Drakin, S. I., Zhur. Fiz. Khim. 26,

581-585 (1952).

[412] Spedding, F. H., and Miller, C. F., J. Am. Chem. Soc. 74, 3158-3162 (1952).
[413] Voskresenskaya, N. K., and Patsukova, N. N., Doklady Akad. Nauk S.S.S.R. 87, 219-221 (1952).

[414] Westrum, E. F., Jr., and Eyring, L., J. Am. Chem. Soc. 74,

2045-2047 (1952) [415] Hutchinson, E., Manchester, K. E., and Winslow, L., J.

Phys. Chem. 58, 1124-1127 (1954).

[416] Chernyaev, I. I., Sokolov, V. A., and Palkin, V. A., Izvest. Sektora Platiny i Drug. Blagorod. Metal., Inst. Obschefii
Neorg. Khim., Akad. Nauk S.S.S.R. 28, 142–160 (1954).
[417] Hutchinson, W. P., and White, A. G., J. Sci. Instr. 32,
309–314 (1955).

[418] Higgins, T. L., Ph. D. thesis, Univ. of Michigan, (1955).

[419] Papadopoulos, M. N., and Giauque, W. F., J. Am. Chem. Soc. 77, 2740–2744 (1955).
[420] Coops, J., Balk, A. N., and Tolk, M. W., Rec. trav. chim. 75, 75–81 (1956).
[421] Gunn, S. R., Rev. Sci. Instr. 29, 377–380 (1958).
[422] P. J. B. B. C. G. L. Blerg, Chem. 62, 1009, 1012.

[422] Balk, P., and Benson, G. C., J. Phys. Chem. 63, 1009-1012 (1959)

[423] Sacconi, L., Paoletti, P., and Ciampolini, M., J. Am. Chem. Soc. **82**, 3828–3831 (1960).

[424] Sunner, S., Wadsö, I., Acta Chem. Scand. 13, 97-108 (1959).[425] Bills, J. L., and Cotton, F. A., J. Phys. Chem. 64, 1477-1479

(1960).[426] Hietala, J., , Ann. Acad. Sci. Fennicae Ser. AVI No. 63, 57

pp. (1960). [427] Gallagher, P. K., and King, E. L., J. Am. Chem. Soc. 82, 3510–3514 (1960).

[428] Dekker, H., Bull. Chem. Thermodynam. No. 4, 22 (1961). [429] Reshetnikov, N. A., Zhur. Neorg. Khim. 6, 682-686 (1961).
 [430] Ciampolini, M., and Paoletti, P., J. Phys. Chem. 65,

1224-1226 (1961). [431] Higgins, T. L., and Westrum, E. F., Jr., J. Phys. Chem. 65,

830-836 (1961).

[432] Argue, G. R., Mercer, E. E., and Cobble, J. W., J. Phys. Chem. 65, 2041–2048 (1961).
[433] Coops, J., Somsen, G., and Tolk, M. W., Private communication (Aug. 1961); Somsen, G., Coops, J., and Tolk, M. W., Rec. trav. chim. 82, 231–242 (1963).
[434] Busey, R. H., Dearman, H. H., and Bevan, R. B., Jr., J. Phys. Chem. 66, 23, 20 (1963).

[434] Busey, K. H., Dearman, H. H., and Bevan, K. B., Jr., J. Phys. Chem. 66, 82-89 (1962).
[435] Newman, E. S., J. Research Natl. Bur. Standards 66A, (Phys. and Chem.) No. 5, 381-388 (1962).
[436] Talakin, O. G., Akhanshchikova, L. A., Sosnovskii, E. N., Pankratov, A. V., and Zercheninov, A. N., Zhur. Fiz. Khim. 36, 1065-1067 (1962).
[437] Veteimisekii, K. B., and Carfona, Z. M., Zhur, Obelschoi.

[437] Yatsimirskii, K. B., and Grafova, Z. M., Zhur. Obshchei Khim. 23, 717-720 (1953).

[438] Gire, G., Ann. chim. (Paris) [10] 4, 183-221 (1925

[439] Kaganovich, Yu. Ya., and Mishchenko, K. P., Zhur. Obshcheĭ Khim. 21, 28-46 (1951).

[440] Lehtonen, L., Soc. Sci. Fennica, Commentationes Phy. Math. 1, No. 13, 1-9 (1922).
[441] Zemczuzny, S., and Rambach, F., Z. anorg. Chem. 65,

403-428 (1910).

- [442] Varali-Thevenet, A., Nuovo cimento [5] 4, 186-192 (1902).
 [443] Ostwald, W., J. prakt. Chem. [2] 25, 1-19 (1882).
 [444] Samoilov, O. Ya., and Buslaeva, M. N., Stroenie Vesh-chestva i Spektroskopiya, Akad. Nauk S.S.S.R. 1960, 102-110.
- [445] Kessler, Yu. M., and Gorbanev, A. I., Izvest. Vostoch. Filialov Akad. Nauk S.S.S.R. 1957, No. 4-5, 75-89.
 [446] Agostini, P., Atti accad. nazl. Lincei Rend., Classe sci. fis. mat. e nat. [6] 71, 1030-1032 (1928).
 [447] Richardson, L. T., and Wells, R. C., J. Wash. Acad. Sci. 21, 243-248 (1931).

- [448] Vrzhesnevskii, I. B., Z. anorg. Chem. 74, 95-121 (1912).
 [449] von Wartenberg, H., and Fitzner, O., Z. anorg. u. allgem. Chem. 151, 313-325 (1926).
- [450] Khaidukov, N., Linetskaya, Z., and Bognovarov, A., Zhur. Priklad. Khim. 9, 439–445 (1936).
 [451] Franck, E. U., and Meyer, F., Z. Elektrochem. 63, 571–582 (1959).

- [452] Smith, D. F., J. Chem. Phys. **28**, 1040–1056 (1958). [453] Smith, D. F., J. Mol. Spectroscopy **3**, 473–485 (1959). [454] Spalthoff, W., and Franck, E. U., Z. Elektrochem. **61**, 993–1000 (1957). [455] Long, R. W., Hildebrand, J. H., and Morrell, W. E., J. Am.
- Chem. Soc. 65, 182–187 (1943). [456] Hu, J. H., White, D., and Johnston, H. L., J. Am. Chem. Soc.
- 75, 1232-6 (1953).
 [457] Jarry, R. L., and Davis, W., Jr., J. Phys. Chem. 57, 600-
- 604 (1953). [458] Berthelot, M., and Louguinine, W., Ann. chim. et phys.
- [5] 6, 289-304 (1875). [459] Roth, W. A., and Richter, H., Z. physik. Chem. A170,
- 123-133 (1934).
- [460] Roth, W. A., and Bertram, A., Z. Elektrochem. 43, 376-378 (1937).
- [461] Roth, W. A., Börger, E., and Siemonsen, H., Atti congr. intern. chim., 10th Congr., Rome, 1938, 2, 775-784.
 [462] Gunn, S. R., and Green, L. G., J. Chem. Eng. Data 8, 180,
- [463] Abria, M., Compt. rend. 22, 372–373 (1846).
- [464] Favre, P. A., Compt. rend. 73, 971-979 (1871).
 [465] Rosolovskii, V. Ya., Krivstov, N. V., and Zinov'ev, A. A., Zhur. Neorg. Khim. 5, 778-781 (1960).
- [466] Arlman, E. J., Rec. trav. chim. 58, 871-874 (1939).
 [467] Ruff, O., and Menzel, W., Z. anorg. u. allgem. Chem. 198, 375-382 (1931).

- 198, 375-382 (1931).
 [468] Cottrell, F. G., J. Phys. Chem. 2, 492-495 (1898).
 [469] Ditte, A., Compt. rend. 70, 935-940 (1870).
 [470] Karve, D. D., J. Indian Chem. Soc. 1, 247-262 (1925).
 [471] Hess, H., Ann. Physik [2] 53, 535-547 (1841).
 [472] Leclerc, R., and Chédin, J., Mem. serv. chim. etat (Paris) 32, 87-91 (1946).
 [473] André, G., Compt. rend. 126, 1105-1107 (1898).
 [474] Pickering, S. U., J. Chem. Soc. 67, 664-684 (1895).
 [475] Pritchard, H. O., and Skinner, H. A., J. Chem. Soc. 1950, 272-276.

- 1950, 272-276.
- [476] Berthelot, M., Ann. chim. et phys. [4] **30**, 456–539 (1873). [477] Nacken, R., Zement **19**, 818–825 (1930). [478] Anderson, E., and Noyes, H. A., J. Phys. Chem. **17**, 249–251 (1913).
- [479] Varet, R., Ann. chim. et phys. [7] 8, 240-288 (1896).
- [480] André, G., Ann. chim. et phys. [6] **3**, 66–129 (1884). [481] Birky, M. M., and Hepler, L. G., J. Phys. Chem. **64**, 686–687 (1960).
- [482] Thomsen, J., J. prakt. Chem. [2] 16, 323-341 (1877).
 [483] Kapustinskii, A. F., Shidlovskii, A. A., and Shidlovskaya, Yu. S., Izvest. Akad. Nauk S.S.S.R., Otdel. Khim. Nauk 1958, 385-388.
- [484] Spencer, J. G., Jr., and Hepler, L. G., J. Phys. Chem. 64, 499-500 (1960).

- 499-300 (1900).

 [485] Berthelot, M., Ann. chim. et phys. [5] **6**, 145-178 (1875).

 [486] Berthelot, M., Compt. rend. **78**, 99-106 (1874).

 [487] Bureau, J., Ann. chim. (Paris) [11] **8**, 1-142 (1937).

 [488] Thomsen, J., J. prakt. Chem. [2] **17**, 165-183 (1878).

 [489] Becker, G., and Roth, W. A., Z. Elektrochem. **40**, 836-
- 843 (1934). [490] Cohen, E., and Helderman, W. D., Koninkl. Ned. Adad.
- Wetenschap. Proc. 28, 2–12 (1925). [491] Cohen, E., and Helderman, W. D., Z. physik. Chem. 113, 145–156 (1924).

- [492] Cohen, E., Moesveld, A. L. T., and Helderman, W. D., Z. physik. Chem. 112, 141-149 (1924).
- [493] Mondain-Monval, P., Compt. rend. 177, 175–178 (1923).
 [494] Berthelot, M., Compt. rend. 77, 24–32 (1873).
 [495] Berthelot, M., Compt. rend. 73, 448–459 (1871).
- [496] Joannis, A., Ann. chim. et phys. [5] 26, 482-548 (1882).
- [497] Felsing, W. A., and Thomas, A. R., Ind. Eng. Chem. 21, 1269-1272 (1929).
- [498] Berthelot, M., Compt. rend. 91, 139-145 (1880).
 [499] Askew, F. A., Bullock, E., Smith, H. T., Tinkler, R. K., Sacty, O., and Wolfenden, J. W., J. Chem. Soc. 1934, 1368–1376.

- [500] Walden, P. T., Z. physik. Chem. 58, 479–511 (1907).
 [501] Guntz, A., Compt. rend. 98, 819–821 (1884).
 [502] Guntz, A., and Guntz, A. A., Jr., Compt. rend. 157, 977–981 (1913).
- [503] Guntz, A., and Guntz, A. A., Jr., Ann. chim. (Paris) [9] 2, 101-132 (1914).
- [504] Smith, W. V., Brown, O. L. I., and Pitzer, K. S., J. Am. Chem. Soc. 59, 1213-1215 (1937).
 [505] Graham, R. L., Ph. D. thesis, Univ. of Virginia, (1958).
 [506] Graham, R. L., and Hepler, L. G., J. Am. Chem. Soc. 80, 1272, 1272, 1272.
- 3538-3540 (1958).

- 5338-3540 (1958).
 [507] de Forcrand, R., Compt. rend. 145, 702-705 (1907).
 [508] de Forcrand, R., Compt. rend. 146, 802-806 (1908).
 [509] Truchot, C., Compt. rend. 98, 1330-1333 (1884).
 [510] de Forcrand, R., Compt. rend. 142, 1252-1255 (1906).
 [511] Kapustinskii, A. F., and Stakhanova, M. S., Trudy Moskov. Khim.-Tekhnol. Inst. im. D. I. Mendeleeva, 1956, No. 22, 21-29, taken from Chem. Abstracts 51, 16071-22, 21-29, taken from Chem. Abstracts 51, 16071-16072 (1957).
- [512] Kolesov, V. P., and Skuratov, S. M., Zhur. Neorg. Khim. 6, 1741-1744 (1961).
- [513] de Forcrand, R., Ann. chim. et phys. [8] 24, 256-282 (1911).
- [514] Payne, J. H., J. Am. Chem. Soc. 59, 947 (1937).
- [515] Pickering, S. U., J. Chem. Soc. **53**, 865–878 (1888). [516] Bonnefoi, J., Compt. rend. **124**, 771–773 (1897).
- [516] Bonnefoi, J., Compt. rend. 124, 771-773 (1897).
 [517] Maier, O., Dissertation, Deutsche Universität, Prag, (1927), quoted in Slonim, C., and Hüttig, G. F., Z. anorg. u. allgem. Chem. 181, 55-64 (1929).
 [518] Bogorodskii, A. Ya., Zhur. Russ. Fiz.-Khim. Obshchestva 43, 1268-1274 (1911).
 [519] Markowitz, M. M., Harris, R. F., and Stewart, H., Jr., J. Phys. Chem. 63, 1325-1326 (1959).
 [520] Bodisko, A., Zhur. Russ. Fiz.-Khim. Obshchestva 21. 7-9 (1889)

- (1889).
- [521] Bonnefoi, J., Compt. rend. 130, 1394–1397 (1900).
 [522] Boyd, G. E., and Vaslow, F., J. Chem. Eng. Data 7, 237–238 (1962).
 [523] Bodisko, A., Zhur. Russ. Fiz.-Khim. Obshchestva 20, 500–503 (1900).
- 503 (1888).

- [524] Bodisko, A., Ber. 21, 586 (1888).
 [525] Bodisko, A., Z. physik. Chem. 4, 482 (1889).
 [526] Mosnier, A., Ann. chim. ct phys. [7] 12, 374-426 (1897).
 [527] Beketov, N., and Beketov, W., Z. anorg. Chem. 40, 355-373 (2004). 371 (1904)

- [528] Dodé, M., Bull. soc. chim. France [5] 4, 2093–2105 (1937).
 [529] de Forcrand, R., Compt. rend. 133, 223–226 (1901).
 [530] Bobtelsky, M., and Lairsch, R. D., J. Chem. Soc. 1950, 3612–3615.
- [531] Murch, L. E., and Giauque, W. F., J. Phys. Chem. 66, 2052-2059 (1962).
- [532] Pickering, S. U., Phil. Mag. [5] 39, 510-528 (1895).
 [533] Hepler, L. G., Jolly, W. L., and Latimer, W. M., J. Am. Chem. Soc. 75, 2809-2810 (1953).
 [534] Latimer, W. M., and Jolly, W. L., J. Am. Chem. Soc. 75, 1549 1559 (1957).
- 1548-1550 (1953)
- [535] Colson, A., Compt. rend. 161, 414-416 (1915).
- [536] Lipsett, S. G., Johnson, F. M. G., Maass, O., J. Am. Chem. Soc. 49, 1940–1949 (1927).
 [537] Seki, S., and Suzuki, K., Bull. Chem. Soc. Japan 26, 63–67

- [538] Benson, G. C., Goddard, E. D., and Hoeve, C. A., Rev. Sci. Instr. 27, 725 (1956).
 [539] Attree, R. W., Cushing, R. L., Ladd, J. A., and Pieroni, J. J., Rev. Sci. Instr. 29, 491-496 (1958).

- [540] Lister, M. W., and Meyers, N. F., J. Phys. Chem. 62, 145-150 (1958).
- [541] Criss, C. M., and Cobble, J. W., J. Am. Chem. Soc. 83, 3223-3228 (1961).
- [542] Staub, G., Dissertation, Zurich, (1890), quoted in Scholz,
 R., Ann. Physik [3] 45, 193-206 (1892).
- [543] Staub, G., Beibl. Ann. Physik 14, 493-494 (1890). [544] Berthelot, M., Ann. chim. et phys. [5] 4, 5-7 (1875). [545] Mishchenko, K. P., and Yakovlev, I. F., Zhur. Obshchei Khim. 29, 1761-1771 (1959).
- [546] Jones, E. D., Burgess, D. S., and Amis, E. S., Z. physik. Chem. [NF] 4, 220-232 (1955).
 [547] Allmand, A. J., and Polack, W. G., J. Chem. Soc. 115,
- 1020-1039 (1919).
- [548] Fontana, B. J., and Latimer, W. M., J. Am. Chem. Soc. 69, 2598-2599 (1947)
- [549] Levi, G. R., and Bisi, C., Gazz. chim. ital. 86, 392-398 (1956).
- [550] Ishi, G., Nippon Kagaku Zasshi 82, 1421–1422 (1961).
 [551] Nelson, T., Moss, C., and Hepler, L. G., J. Phys. Chem. 64, 376–377 (1960).
- [552] Foote, H. W., and Saxton, B., J. Am. Chem. Soc. 36, 1695-1704 (1914).
- [553] Brönsted, J. N., Z. physik. Chem. 80, 206-234 (1912).
 [554] Favre, P. A., and Valson, C. A., Compt. rend. 77, 577-584 (1873).
- [555] Mishchenko, K. P., and Sukhotin, A. M., Doklady Akad. Nauk S.S.S.R. 98, 103-106 (1954).
- [556] Shidlovskii, A. A., and Valkina, K. V., Zhur. Fiz. Khim. 35,
- [557] Kapustinskii, A. F., and Yatsimirskii, K. B., Zhur. Obshchei Khim. 19, 2191–2200 (1949).
 [558] Swietoslawski, W., Z. physik. Chem. 72, 49–83 (1910).
 [559] Matignon, C., and Marchal, G., Compt. rend. 170, 232–233
- [560] Tilden, W. A., Proc. Roy. Soc. (London) 38, 401–414 (1885).
 [561] Van Tassel, J. H., and Wendlandt, W. W., J. Am. Chem. Soc. **81**, 813–816 (1959).
- [562] Zawidzki, J., and Schagger, A., Kosmos **35**, 498–505 (1910). [563] Jolly, W. L., U.S. AEC UCRL **1688**, 11 pp. (April 1952). [564] Jolly, W. L., J. Am. Chem. Soc. **74**, 6199–6201 (1952). [565] Lord, G., and Woolf, A. A., J. Chem. Soc. **1954**, 2546–2551.
- [566] Lemoult, P., Ann. chim. et phys. [7] 16, 338-432 (1899).[567] de Forcrand, R., Compt. rend. 133, 157-159 (1901).
- [568] Noyes, A. A., and Sammet, G. V., Z. physik. Chem. 43, 513-538 (1903).
- [569] Latimer, W. M., and Ahlberg, J. E., J. Am. Chem. Soc. 52, 549-553 (1930).
- [570] Balarew, D., and Kolarow, N., Kolloid. Z. 88, 288-290
- [571] Berthelot, M., Ann. chim. et phys. [5] 13, 5–20 (1878).
 [572] Mel, H. C., U.S. AEC UCRL 2330, 108 pp. (Sept. 1953).
 [573] Mel, H. C., Jolly, W. L., and Latimer, W. M., J. Am. Chem. Soc. 75, 3827–3829 (1953). [574] Kantola, M., Ann. Acad. Sci. Fennicae, Ser. AI, No. 30,
- 3-7 (1945).
- [575] Hieber, W., and Mühlbauer, F., Z. anorg. u. allgem. Chem. 186, 97-118 (1930).
- [576] Roth, W. A., Z. physik. Chem. 130, 539-544 (1927).
 [577] Roth, W. A., and Müller, F., unpublished measurements, quoted in Roth, W. A., Z. physik. Chem. 130, 539-544
- [578] Roth, W. A., Z. Elektrochem. 49, 322-325 (1943).
 [579] Hieber, W., and Feder, E. E., Z. Elektrochem. 44, 881-887 (1938).
- [580] Hückel, W., Datow, J., and Simmershach, E., Z. physik. Chem. A186, 129-179 (1940).
 [581] Voskresenskaya, N. K., and Ponomareva, K. S., Compt. rend. acad. sci. U.R.S.S. 45, 188-190 (1944).
- [582] Lipilina, I. I., and Samoilov, O. Ya., Doklady Akad. Nauk S.S.S.R. 98, 99-102 (1954).
- [583] Turnbull, A. G., J. Phys. Chem. **65**, 1652–1654 (1961).
- [584] Person, C. C., Ann. chim. et phys. [3] 33, 448-464 (1851).
 [585] Leval't-Ezerskiï, M., Zhur. Russ. Fiz.-Khim. Obshchestva, Chast' Khim. 44, 665-677 (1912).
 [506] Berthelot, M., Compt. rend. 78, 1085-1092 (1874).
 [587] Roth, W. A., and Becker, G., Z. physik. Chem. A159, 23 (2012).
- -39 (1932)
- [588] de Forcrand, R., Compt. rend. 152, 27-31 (1911).

- [589] de Forcrand, R., Compt. rend. 152, 1208-1211 (1911).
 [590] Samoilov, O. Ya., Izvest. Akad. Nauk S.S.S.R., Otdel. Khim. Nauk 1956, 1415-1416.
- [591] Pitzer, K. S., J. Am. Chem. Soc. 60, 1828–1829 (1938).
 [592] Vorobiev, A. F., Skuratov, S. M., and Ebragen, N., Bull. Chem. Thermodynam. No. 4, 27 (1961).
- [593] Beketov, N., Bull. acad. imp. sci. (St. Petersbourg) 34, 171-173 (1892).
- [594] Rodnikova, M. N., Zhur. Neorg. Khim. 3, 2417-2419
- [595] Lemlich, R., Gottschlich, C., and Hoke, R., Chem. Eng. Data Ser. 2, 32-35 (1957).
 [596] Van Nuys, C. C., Trans. Am. Inst. Chem. Engrs. 39, 663-
- 678 (1943).
- [597] Chédin, J., J. chim. phys. 49, 109-125 (1952).[598] McCurdy, J. L., and McKinley, C., Ind. Eng. Chem. 34, 1002-1004 (1942)
- [599] Rozenfel'd, L. M., Doklady Akad. Nauk S.S.S.R. 82, 723-725 (1952).
- [600] Othmer, D. F., and Frohlich, G. J., A. I. Ch. E. Journal 6,
- 210-214 (1960). [601] Kangro, W., and Croeneveld, A., Z. physik. Chem. [NF] **32**, 110-126 (1962).
- [602] Löwer, H., Kältetechnik 13, 178–184 (1961).
- [603] Rozenfel'd, L. M., and Karnaukh, M. S., Zhur. Tekh. Fiz. 28, 655-660 (1958).
- [604] Rant, Z., Forsch. Gebiete Ingenieurw. 20B, 77-80 (1954).
 [605] Rant, Z., Forsch. Gebiete Ingenieurw. 26B, 1-7 (1960).
 [606] Ginzburg, D. M., Zhur. Fiz. Khim. 36, 747-751 (1962).
 [607] Haltenberger, W., Jr., Ind. Eng. Chem. 31, 783-786 (1939).
- [608] Fábry, G., Acta Tech. Acad. Sci. Hung. 14, 313-317
- [609] Shields, C. T., Refrig. Eng. 59, 880–884 (1951).
 [610] Ginzburg, D. M., Zhur. Fiz. Khim. 33, 1087–1092 (1959).
 [611] Franck, E. U., and Spalthoff, W., Z. Elektrochem. 61, 348-357 (1957)
- [612] Koehler, M. F., U.S. Bur. Mines, Rept. Invest. 5700, 10 pp. (1960).
- [613] Lange, E., and Monheim, J., Z. Elektrochem. 35, 29-33 (1929).
- [614] von Stackelberg, E., Z. physik. Chem. 20, 159-167 (1896).
 [615] Seidell, A., Solubilities of inorganic and metal organic compounds, Vol. I, 3d edition (D. Van Nostrand Co.,
- Inc., New York, 1940).
 [616] Seidell, A., and Linke, W. F., Solubilities of inorganic and organic compounds, 3d edition, supplement (D. Van
- Nostrand Co., Inc., New York, 1952).

 [617] Linke, W. F., Solubilities of inorganic and metal organic compounds, Vol. I, 4th edition (D. Van Nostrand Co., Inc., New York, 1958).
- 101., New 107K, 1938).

 [618] Rossini, F. D., Pitzer, K. S., Arnett, R. L., Braun, R. M., Pimentel, G. C., Li, K., Ries, H. J., Cowie, P. A., Mitchell, J., Parcella, L. A., Epstein, M. B., McClellan, A. L., Sarao, M. H., De Celles, C. R., and Flanagan, H. M., Selected values of physical and thermodynamic properties of hydrocybor and related properties of hydrocybor and hydrocybor ties of hydrocarbons and related compounds (Carnegie
- Press, Pittsburgh, Pennsylvania, 1953).
 [619] Kharbanda, O. P., Ind. Chemist 33, 136 (1957).
 [620] Bump, T. R., and Sibbitt, W. L., Ind. Eng. Chem. 47, 1665–1670 (1955).

- [621] Voskresenskaya, N. K., Yankovskaya, G. N., and Anosov, V. Ya., Zhur. Priklad. Khim. 21, 18-25 (1948).
 [622] Strelkov, I. I., Ukrain. Khim. Zhur. 21, 551-553 (1955).
 [623] Messer, C. E., and Ziegler, W. T., J. Am. Chem. Soc. 63, 2703-2708 (1941).
- [624] Mustajoki, A., Ann. Acad. Sci. Fennicae, AVI, No. 9, 16 pp. (1958).
- [625] Eigen, M., and Wicke, E., J. Phys. Chem. 58, 702-714 (1954).
- [626] Latimer, W. M., Chem. Revs. 18, 349-358 (1936).
 [627] Mishchenko, K. P., Zhur. Fiz. Khim. 26, 1736-1760 (1952).
 [628] Gucker, F. T., Jr., Ann. N.Y. Acad. Sci. 51, 680-704 (1949).
 [629] Wiist, J., and Lange, E., Z. Elektrochem. 30, 523-528
- (1924).
- [630] Vasil'ev, V. P., Zolotarev, E. K., Kapustinskii, A. F., Mishchenko, K. P., Podgornaya, F. A., and Yatsimirskii, K. B., Zhur. Fiz. Khim. 34, 1763-1767 (1960).
 [631] Robinson, A. L., and Frank, H. S., J. Am. Chem. Soc. 56,
- 2312-2313 (1934).

[632] Kapustinskii, A. F., Drakin, S. I., and Yakushevskii, B. M., Zhur. Fiz. Khim. 27, 433-442 (1953).
[633] Robinson, R. A., and Stokes, R. H., Electrolyte solutions, 2d edition (Academic Press, Inc., New York, 1959).

2d edution (Academic Press, Inc., New York, 1959).

[634] Hood, G. C., and Woyski, M. M., J. Am. Chem. Soc. 73, 2738–2741 (1951).

[635] Yatlov, V. S., and Polyakov, E. M., Zhur. Obshchei Khim. 8, 774–776 (1938).

[636] Appleby, M. P., Crawford, F. H., and Gordon, K., J. Chem. Soc. 1934, 1665–1671.

Soc. 1934, 1665–1671.
[637] Hüttig, G. F., and Reuscher, F., Z. anorg. u. allgem. Chem. 137, 155–180 (1924).
[638] Campbell, A. N., and Griffiths, J. E., Can. J. Chem. 34, 1647–1661 (1956).
[639] Hüttig, G. F., and Pohle, F., Z. anorg. u. allgem. Chem. 138, 1–16 (1924).

[640] Ray, J. D., and Ogg, R. A., Jr., J. Phys. Chem. **60**, 1599–1600 (1956).

[641] Riganti, V., Gazz. chim. ital. 89, 2275–2279 (1959). [642] Markowitz, M. E., and Boryta, D. A., J. Chem. Eng. Data 6,

[642] Markowitz, M. E., and Boryta, D. A., J. Chem. Eng. Data 6, 16-18 (1961).
[643] Dingemans, P., Rec. trav. chim. 57, 703-709 (1938).
[644] Bell, J., J. Chem. Soc. 1937, 459-461.
[645] Dingemans, P., Rec. trav. chim. 57, 144-151 (1938).
[646] Baxter, G. P., and Cooper, W. C., Jr., J. Am. Chem. Soc. 46, 923-933 (1924).
[647] Bell, J., J. Chem. Soc. 1940, 72-74.
[648] Lannung, A., Z. physik. Chem. A170, 134-138 (1934).
[649] Osborne, N. S., Stimson, H. F., and Ginnings, D. C., J. Research Natl. Bur. Standards 23, 197-260 (1939).
[650] Shibata, F. L. E., Oda, S., and Furukawa, S., J. Sci. Hiroshima Univ. Ser. A2, 85-102 (1932).
[651] Bolte, H., Z. physik. Chem. 80, 338-360 (1912).
[652] Vanderzee, C. E., and Nutter, J. D., J. Phys. Chem. 67, 2521-2525 (1963).

2521-2525 (1963).

X. Tables

TABLE 1

Φ°, cal/deg mole, of the ions at 25 °C

Ion	$\Phi_{\mathcal{C}}^{\circ}$
H+	0
NH ₄ +	19.1
Ag+	5.2
Li+	16.4
Na+	11.1
K ⁺	5.2
Rb+	*3.3
Cs+	1.9
OH	-35.5
F	- 25.5
Cl	-32.6
ClO ₃	- 19.7
ČlO ₄	-18.2
Br	-33.9
BrO ₃	-26.5
I	-34.0
IO_3	- 29.6
IO ₄	* 28.2
NO ₂ -	- 23.3
NO ₃	-20.7
CNS	-9.6
CHO ₂	-21.0
C ₂ H ₃ O ₂	-1.5
$C_{2}H_{3}O_{2}$ $C_{3}H_{5}O_{2}$	+26.7
MnO ₄	-19.6

^{*}Estimated.

TABLE II A. Heat capacity: aqueous HCl

Table II B. Heat capacity: aqueous NaOH Φ_c , cal/deg mole

 Φ_c , cal/deg mole

		TO Car	deg moie					Ψ _C , cai/deg	more	-	
n	m	15°	20°	25°	30°	n	m	15°	20°	25°	30°
0,000 50,000 20,000 10,000 7,000	0.00 .000555 .00111 .00278 .00555 .00793	-34.8 -34.6 -34.6 -34.5 -34.4 -34.3	-33.8 -33.6 -33.6 -33.5 -33.3 -33.2	-32.6 -32.4 -32.4 -32.3 -32.2 -32.1	-31.9 -31.7 -31.6 -31.5 -31.4 -31.3	100,000 50,000 20,000 10,000 7,000	0.00 .000555 .00111 .00278 .00555 .00793	-34.4 -34.3 -34.3 -34.2 -34.1 -34.0	-29.4 -29.3 -29.2 -29.1 -29.0 -28.9	-24.4 -24.3 -24.2 -24.1 -23.9 -23.8	-18.5 -18.6 -18.7 -18.6 -18.5 -18.4
5,000 4,000 3,000 2,000 1,500	.01110 .01388 .01850 .02775 .03700	-34.2 -34.2 -34.1 -33.9 -33.7	-33.1 -33.1 -33.0 -32.8 -32.6	-32.0 -31.9 -31.8 -31.6 -31.4	$\begin{array}{r} -31.2 \\ -31.1 \\ -31.0 \\ -30.8 \\ -30.6 \end{array}$	5,000 4,000 3,000 2,000 1,500	.01110 .01388 .01850 .02775 .03700	-33.8 -33.7 -33.5 -33.2 -32.8	-28.7 -28.6 -28.4 -28.1 -27.7	-23.7 -23.6 -23.4 -23.1 -22.8	-18.3 -18.2 -18.0 -17.8 -17.2
1,000 900 800 700 600	.05551 .0617 .0694 .0793 .0925	-33.4 -33.3 -33.2 -33.1 -32.9	-32.3 -32.2 -32.1 -31.9 -31.8	-31.2 -31.2 -31.1 -30.9 -30.8	-30.3 -30.3 -30.2 -30.0 -29.9	1,000 900 800 700 600	.05551 .0617 .0694 .0793 .0925	-32.1 -31.9 -31.6 -31.3 -30.8	-27.1 -26.9 -26.6 -26.3 -25.9	$\begin{array}{r} -22.2 \\ -22.0 \\ -21.8 \\ -21.5 \\ -21.2 \end{array}$	-17.2 -17.0 -16.9 -16.7 -16.4
500 400 300 200 150	.1110 .1388 .1850 .2775 .3700	-32.7 -32.4 -31.9 -31.1 -30.4	-31.6 -31.3 -30.9 -30.2 -29.6	-30.6 -30.3 -30.0 -29.3 -28.8	-29.7 -29.5 -29.1 -28.5 -27.9	500 400 300 200 150	.1110 .1388 .1850 .2775 .3700	-30.2 -29.4 -28.1 -25.8 -23.8	-25.4 -24.6 -23.5 -21.4 -19.7	-20.7 -20.0 -18.9 -17.1 -15.5	-16.1 -15.5 -14.6 -13.1 -11.8
100 75 50 40 30	.5551 .7401 1.1101 1.3877 1.8502	$ \begin{array}{r} -29.3 \\ -28.4 \\ -27.0 \\ -26.1 \\ -24.7 \end{array} $	-28.6 -27.8 -26.4 -25.5 -24.2	-27.8 -27.0 -25.6 -24.8 -23.6	$ \begin{array}{r} -27.1 \\ -26.3 \\ -25.1 \\ -24.3 \\ -23.2 \end{array} $	100 75 50 40 30	.5551 .7401 1.1101 1.3877 1.8502	-20.4 -17.3 -12.5 -9.8 -6.0	-16.6 -14.0 -9.7 -7.1 -3.5	-12.9 -10.7 -7.1 -4.8 -1.7	-9.7 -7.8 -4.7 -2.8 -0.1
25 20 15 12 10	2.2202 2.7753 3.7004 4.6255 5.5506	$ \begin{array}{r} -23.7 \\ -22.4 \\ -20.5 \\ -18.8 \\ -17.3 \end{array} $	-23.2 -21.9 -20.2 -18.5 -17.0	-22.7 -21.5 -19.8 -18.2 -16.8	-22.3 -21.2 -19.6 -18.0 -16.6	25 20 15 12 10	2.2202 2.7753 3.7004 4.6255 5.5506	$ \begin{array}{r} -3.5 \\ -0.3 \\ +4.1 \\ 7.5 \\ 10.3 \end{array} $	-1.2 +1.8 5.7 8.8 11.3	+0.4 3.2 6.8 9.8 12.2	+1.7 4.7 7.7 10.5 12.9
9.5 9.0 8.5 8.0 7.5	5.8427 6.1674 6.5301 6.9383 7.4008	-16.8 -16.4 -15.8 -15.3 -14.7	-16.6 -16.1 -15.6 -15.1 -14.5	- 16.4 - 15.8 - 15.4 - 14.8 - 14.2	-16.2 -15.7 -15.2 -14.6 -14.0	9.5 9.0 8.5 8.0 7.5	5.8427 6.1674 6.5301 6.9383 7.4008	11.0 11.8 12.6 13.4 14.2	12.0 12.7 13.5 14.3 15.1	12.8 13.5 14.3 15.1 15.8	13.6 14.3 15.1 15.8 16.6
7.0 6.5 6.0 5.5 5.0	7.9295 8.5394 9.2510 10.0920 11.1012	$\begin{array}{c} -14.0 \\ -13.2 \\ -12.3 \\ -11.3 \\ -10.1 \end{array}$	-13.8 -13.0 -12.1 -11.1 -9.8	-13.5 -12.7 -11.8 -10.8 -9.6	-13.2 -12.5 -11.6 -10.6 -9.4	7.0 6.5 6.0 5.5 5.0	7.9295 8.5394 9.2510 10.0920 11.1012	15.1 15.9 16.8 17.6 18.4	16.0 16.9 17.7 18.4 19.2	16.6 17.5 18.4 19.1 19.8	17.4 18.2 19.0 19.7 20.4
4.5 4.0 3.5	12.3346 13.8765 15.8589	-8.2 -7.1 -5.1	-8.5 -6.8 -4.9	-8.7 -6.6 -4.7	-8.0 -6.5 -4.6	4.5 4.0 3.5 3.25 3.0	12.3346 13.8765 15.8589 17.0788 18.5020	19.4 20.3 21.4 21.8 22.0	20.0 20.9 21.7 22.0 22.2	20.6 21.3 22.0 22.2 22.3	21.1 21.8 22.3 22.5 22.5
						2.5 2.0	22.2024 27.7530	22.0 21.2	22.1 21.2	$\frac{22.2}{21.2}$	22.3 21.2

TABLE II C. Heat capacity: aqueous NaCl

Φ_C, cal/deg mole

		Φ_C , cal/deg mo	le		
n	m	15°	20°	25°	30°
∞	0.00	-25.2	-23.3	-21.5	-20.1
100,000 50,000	.000555 .00111	$ \begin{array}{r} -25.1 \\ -25.0 \end{array} $	$-23.2 \\ -23.1$	$ \begin{array}{r r} -21.4 \\ -21.3 \end{array} $	-20.0 -19.9
20,000	.00278	-24.9	-23.0	-21.2	- 19.8
10,000	.00555	-24.8	- 22.9	-21.1	-19.6
7,000	.00793	-24.7	-22.8	-21.0	-19.5
5,000	.01110	-24.6	-22.7	-20.9	-19.4
4,000	.01388	-24.5	-22.6	-20.8	-19.3
3,000	.01850	-24.4	-22.5	-20.7	-19.2
2,000	.02775	-24.2	-22.3	-20.4	-19.0
1,500	.03700	-23.9	-22.0	-20.2	-18.8
1,000	.05551	-23.4	-21.6	-19.8	-18.4
900	.0617	-23.3	-21.4	-19.6	-18.2
800	.0694	-23.1	-21.3	- 19.5	-18.1
700	.0793	-22.8	-21.0	- 19.3	17.9
600	.0925	-22.5	-20.8	-19.0	-17.6
500 400	.1110	-22.0	-20.3 -19.8	-18.6 -18.2	-17.3
400	.1388	-21.4	- 19.8	- 18.2	16.9
300	.1850	-20.5	-19.0	-17.5	-16.2
200	.2775	- 19.0	-17.5	-16.3	-15.1
150	.3700	-17.7	-16.4	-15.2	-14.1
100 75	.5551 .7401	-15.5 -13.6	-14.8 -12.6	-13.4 -11.7	-12.5 -10.9
19	7401	-15.0	- 12.0	-11.7	- 10.9
50	1.1101	-10.4	-9.6	-8.8	-8.2
40	1.3877	-8.3	-7.6	-7.0	6.4
30	1.8502	-5.7	-5.1	-4.6	-4.1
25	2.2202	-3.5	-2.9	-2.4	-2.0
20	2.7753	-0.8	-0.4	0.0	+0.2
15	3.7004			+3.3	3.5
12	4.6255			6.3	
10	5.5506			9.0	
9.5	5.8427	•••••		9.7	
9.0	6.1674	•••••		10.5	

Table III. Heat capacity: acids

 $\Phi_{\it C},$ cal/deg mole, at 25 °C

				To, carries in	1010, at 20 0					
n	m	HF	HCl	HBr	ні	HIO ₃	HNO ₃	CH ₂ O ₂	C ₂ H ₄ O ₂	C ₃ H ₆ O ₂
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	-25.5 -23.0 -18.8 -16.6 -12.4 -8.6	-32.6 -32.4 -32.4 -32.3 -32.2	-33.9 -33.8 -33.7 -33.6 -33.4	-34.0 -33.9 -33.8 -33.7 -33.5	- 29.6 	$\begin{array}{c c} -20.7 \\ -20.6 \\ -20.5 \\ -20.4 \\ -20.3 \end{array}$	$\begin{array}{r} -21.0 \\ -9.8 \\ -1.2 \\ +3.1 \\ 10.2 \\ 12.7 \end{array}$	$ \begin{array}{r} -1.5 \\ +25.8 \\ 32.6 \\ 34.0 \\ 35.7 \\ 36.9 \end{array} $	+26.7 38.0 45.1 48.3 52.2 54.1
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	$ \begin{array}{r} -6.7 \\ -4.9 \\ -3.6 \\ -2.2 \\ -0.5 \end{array} $	$ \begin{array}{r} -32.1 \\ -32.0 \\ -31.9 \\ -31.8 \\ -31.6 \end{array} $	-33.4 -33.3 -33.2 -33.1 -32.9	-33.4 -33.3 -33.2 -33.1 -32.9	$\begin{array}{r} -28.4 \\ -28.1 \\ -27.7 \\ -27.2 \\ -25.8 \end{array}$	$\begin{array}{c} -20.2 \\ -20.1 \\ -20.1 \\ -20.0 \\ -19.8 \end{array}$	13.7 14.3 14.7 15.1 15.8	37.4 37.8 37.9 38.1 38.5	54.8 55.3 55.6 56.1 56.7
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	+0.4 1.6 1.8 2.0 2.2	$ \begin{array}{r} -31.4 \\ -31.2 \\ -31.2 \\ -31.1 \\ -30.9 \end{array} $	$ \begin{array}{r} -32.7 \\ -32.5 \\ -32.4 \\ -32.3 \\ -32.2 \end{array} $	$ \begin{array}{r} -32.7 \\ -32.5 \\ -32.4 \\ -32.3 \\ -32.1 \end{array} $	$\begin{array}{r} -24.9 \\ -23.0 \\ -22.3 \\ -21.5 \\ -20.2 \end{array}$	$ \begin{array}{r} -19.6 \\ -19.3 \\ -19.2 \\ -19.1 \\ -19.0 \end{array} $	16.4 16.8 17.0 17.2 17.3	38.7 39.0 39.0 39.1 39.2	57.1 57.7 57.8 57.9 58.0
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	2.4 2.7 2.8 3.2 3.3	$ \begin{array}{r} -30.8 \\ -30.6 \\ -30.3 \\ -30.0 \\ -29.3 \end{array} $	$ \begin{array}{r} -32.1 \\ -31.9 \\ -31.6 \\ -31.2 \\ -30.6 \end{array} $	-32.0 -31.7 -31.4 -30.9 -30.2	-18.7 -16.7 -13.9 -10.1 -4.7	$ \begin{array}{r} -18.9 \\ -18.7 \\ -18.4 \\ -17.9 \\ -17.2 \end{array} $	17.5 17.7 18.0 18.3 18.7	39.3 39.4 39.4 39.4 39.3	58.3 58.4 58.6 58.8 58.9
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877	3.5 3.8 4.1 4.5 4.9	$ \begin{array}{r} -28.8 \\ -27.8 \\ -27.0 \\ 25.6 \\ -24.8 \end{array} $	$ \begin{array}{r} -30.1 \\ -29.1 \\ -28.5 \\ -26.8 \\ -25.9 \end{array} $	-29.6 -28.4 -27.5 25.9 -24.8	$ \begin{array}{r} -0.7 \\ +4.8 \\ 9.2 \\ 15.7 \\ 19.0 \end{array} $	$\begin{array}{r} -16.4 \\ -15.0 \\ -13.7 \\ 11.5 \\ -9.9 \end{array}$	18.9 19.0 19.2 19.0 19.8	39.2 39.1 38.9 38.6 38.4	58.8 58.7 58.6 58.3 58.0
30 25 20 15 12	1.8502 2.2202 2.7758 3.7004 4.6255	5.3 5.6 5.7 6.0 6.2	$\begin{array}{r} -23.6 \\ -22.7 \\ -21.5 \\ -19.8 \\ -18.2 \end{array}$	-24.5 -23.6 -22.2 -20.3 -18.5	-23.3 -22.2 -20.8	23.1 25.7	$ \begin{array}{r} -7.3 \\ -5.4 \\ -2.7 \\ +1.4 \\ \hline 5.0 \end{array} $	20.0 20.1 20.2 20.4 20.4	38.0 37.7 37.1 36.3 35.4	57.1 56.2 55.0 53.1 51.3
10 9.5 9.0 8.5 8.0	5.5506 5.8427 6.1674 6.5301 6.9383	6.3 6.4 6.4 6.5 6.6	$\begin{array}{r} -16.8 \\ -16.4 \\ -15.8 \\ -15.4 \\ -14.8 \end{array}$	-16.8 -16.3 -15.7 -15.1 -14.4			8.5 9.2 10.3 11.4 12.5	20.6 20.6 20.7 20.7	34.8 34.7 34.5 34.4 34.2	49.7
7.5 7.0 6.5 6.0 5.5	7.4008 7.9295 8.5394 9.2510 10.0920	6.7 6.9 7.0 7.1 7.2	-14.2 -13.5 -12.7 -11.8 -10.8	- 13.2 - 12.7 - 11.9 - 10.8 - 9.6			13.7 14.9 16.1 17.1 18.3	20.8 20.8 20.8 20.9 20.9	34.0 33.8 33.5 33.3 33.0	
5.0 4.5 4.0 3.5 3.25	11.1012 12.3346 13.8765 15.8589 17.0788	7.3 7.4 7.5 7.6 7.6	-9.6 -8.7 -6.6 -4.7	-8.2 -6.8 -5.5 -4.0 -3.2			19.3 20.4 21.3 22.1 22.6	21.0 21.1 21.2 21.3 21.4	32.8 32.5 32.2 31.8 31.7	
3.0 2.5 2.0 1.5 1.0	18.5020 22.2024 27.7530 37.0040 55.506	7.7 7.8 7.9		-2.3 -0.4			23.0 23.8 24.6 25.2 25.7	21.5 21.6 21.7 21.8 22.0	31.5 31.2 30.8 30.4 30.1	

TABLE IV. Heat capacity: ammonium and alkyl ammonium compounds

 Φ_{c} , cal/deg mole, at 25 °C

Ψ _C , cal/deg mole, at 25 C										
n	m	NH₄OH	NH₄F	NH₄Cl	NH₄Br*	NH₄I*	NH ₄ NO ₃	CH₃NH₃Cl	(CH ₃) ₂ NH ₂ Cl	(CH ₃) ₃ NHCl
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	-16.4 +19.4 26.7 29.3 31.5 32.7	-6.4 -6.3 -6.2 -6.1 -6.0	-13.5 -13.4 -13.3 -13.2 -13.1	-14.8 (-14.7) (-14.6) (-14.5) (-14.4)	-14.9 (-14.8) (-14.7) (-14.6) (-14.5)	-1.6 -1.5 -1.4 -1.3 -1.1			
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	33.3 33.6 33.7 33.7 33.7	-5.9 -5.8 -5.7 -5.6 -5.4	-13.0 -12.9 -12.8 -12.7 -12.6	(-14.3) (-14.2) (-14.1) (-14.0) (-13.9)	(-14.4) (-14.3) (-14.2) (-14.1) (-14.0)	$ \begin{array}{r} -1.0 \\ -0.9 \\ -0.8 \\ -0.7 \\ -0.5 \end{array} $			
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	33.8 33.9 33.9 34.0 34.1	-5.2 -5.0 -4.9 -4.8 -4.7	-12.4 -12.1 -12.0 -11.9 -11.8	(-13.7) (-13.4) (-13.3) (-13.2) (-13.2)	(-13.8) (-13.5) (-13.4) (-13.3) (-13.2)	$ \begin{array}{r} -0.2 \\ +0.2 \\ 0.4 \\ 0.5 \\ 0.8 \end{array} $			
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	34.1 34.2 34.4 34.6 34.8	-4.5 -4.4 -4.1 -3.6 -2.9	-11.7 -11.6 -11.3 -11.0 -10.3	(-13.0) (-12.8) (-12.5) (-12.1) (-11.4)	(-13.0) (-12.8) (-12.4) (-11.9) (-10.9)	1.0 1.3 1.8 2.5 3.7	4.5 5.1 5.8 6.7 7.8	8,7 9.6 10.9 12.7 15.4	10.3 11.2 12.5 14.3 17.1
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877	34.9 35.2 35.4 35.6 35.7	$ \begin{array}{r} -2.3 \\ -1.4 \\ -0.6 \\ +0.7 \\ 1.7 \end{array} $	-9.7 -8.7 -7.8 -6.2 -5.3	(-10.9) (-9.8) (-9.0) (-7.5) (-6.5)	(-10.1) (-8.7) (-7.5) (-5.5) (-4.3)	4.8 6.6 8.0 10.4 11.9	8.2 8.1 7.5 5.6	17.2 18.8 19.4 19.4	19.3 22.4 24.5 26.8
30 25 20 15 12	1.8502 2.2202 2.7753 3.7004 4.6255	35.7 35.8 35.6 35.6 35.4	2.8 3.7 4.7 6.2 7.5	$ \begin{array}{r} -3.9 \\ -2.9 \\ -1.6 \\ +0.3 \\ 1.9 \end{array} $	(-5.0) (-4.0)	(-2.5) (-1.2)	14.0 15.4 17.3 19.7 21.6			
10 9.5 9.0 8.5 8.0	5.5506 5.8427 6.1674 6.5301 6.9383	35.3 35.3 35.2 35.1 35.1	8.3 8.5 8.8 9.1 9.5	3.4 3.8 4.3 4.8 5.3			22.9 23.3 23.7 24.2 24.7			
7.5 7.0 6.5 6.0 5.5	7.4008 7.9295 8.5394 9.2510 10.0920	35.1 35.0 34.9 34.8 34.7	9.9 10.2 10.6 11.2 11.8	6.0			25.1 25.6 26.1 26.6 27.2			
5.0 4.5 4.0 3.5 3.25	11.1012 12.3346 13.8765 15.8589 17.0788	34.6 34.5 34.5 34.6 34.7	12.5 13.3 14.1 14.9 15.3				27.8 28.5 29.1 29.8 30.2			
3.0 2.5 2.0 1.5 1.0	18.5020 22.2024 27.7530 37.0040 55.506	34.8 35.1 35.1 35.3 35.8	15.7 16.1 17.4 18.3 19.2				30.7 31.6 			

^{*}Estimated.

Table V. Heat capacity: silver and lithium compounds

 $\Phi_{\it C}$, cal/deg mole, at 25 °C

			Φ_{C} , cal/deg mo	le, at 25 °C				
n	m	AgF	$ m AgNO_3$	LiOH	LiCl	LiBr	LiI	LiNO ₃
100,000 50,000 20,000 10,000	0.00 .000555 .00111 .00278 .00555	-20.3 	- 15.5 - 15.4 - 15.3 - 15.2 - 15.0	-19.1 -13.9 -18.8 -18.7 -18.6	-16.2 -16.1 -16.0 -15.9 -15.8	-17.5 -17.4 -17.3 -17.2 -17.1	-17.6 -17.5 -17.4 -17.3 -17.2	-4.3 -4.2 -4.1 -4.0 -3.8
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775		- 14.9 - 14.8 - 14.7 - 14.5 - 14.2	-18.5 -18.3 -18.2 -18.1 -17.9	-15.7 -15.6 -15.5 -15.4 -15.2	-17.0 -16.9 -16.8 -16.7 -16.5	-17.1 -17.0 -16.9 -16.8 -16.5	-3.7 -3.6 -3.5 -3.4 -3.2
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793		-13.9 -13.2 -13.0 -12.8 -12.5	-17.7 -17.3 -17.2 -17.0 -16.8	-15.0 -14.8 -14.7 -14.6 -14.5	-16.3 -16.1 -16.0 -15.9 -15.8	-16.3 -16.0 -15.9 -15.7 -15.6	$ \begin{array}{r} -3.0 \\ -2.8 \\ -2.7 \\ -2.6 \\ -2.5 \end{array} $
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775		-12.1 -11.6 -10.8 -9.7 -7.8	-16.6 -16.2 -15.8 -15.1 -13.9	-14.3 -14.2 -13.9 -13.6 -13.0	-15.7 -15.6 -15.4 -15.0 -14.5	-15.4 -15.2 -14.9 -14.4 -13.6	$ \begin{array}{r} -2.4 \\ -2.2 \\ -1.9 \\ -1.5 \\ -0.8 \end{array} $
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877		$ \begin{array}{r} -6.1 \\ -3.3 \\ -1.0 \\ +2.9 \\ 5.5 \end{array} $	-12.8 -11.0 -9.5 -6.9 -5.3	-12.6 -11.7 -11.1 -10.0 -9.4	$\begin{array}{r} -14.0 \\ -13.2 \\ -12.5 \\ -11.5 \\ -10.8 \end{array}$	$ \begin{array}{r} -13.0 \\ -11.8 \\ -10.8 \\ -9.3 \\ -8.4 \end{array} $	$ \begin{array}{r} -0.1 \\ +1.1 \\ 2.1 \\ 3.8 \\ 4.9 \end{array} $
30 25 20 15 12	1.8502 2.2202 2.7753 3.7004 4.6255	-2.2 -2.1 -1.8 -1.5 -1.4	9.2 11.8	$ \begin{array}{r} -2.9 \\ -1.2 \\ +1.0 \\ \hline 4.6 \\ 7.5 \end{array} $	-8.5 -7.8 -6.9 -5.6 -4.5	-9.7 -9.0 -8.0 -6.8 -5.5	-7.2 -6.4 -5.3 *(-3.9) (-2.6)	6.5 7.7
10 9.5 9.0 8.5 8.0	5.5506 5.8427 6.1674 6.5301 6.9383	-1.2 -1.2 -1.2 -1.2 -1.1			-3.3 -2.9 -2.6 -2.2 -1.8	$ \begin{array}{r rrrr} -4.4 \\ -4.0 \\ -3.6 \\ -3.2 \\ -2.8 \end{array} $	(-1.5) (-1.2) (-0.9) (-0.5) (-0.1)	
7.5 7.0 6.5 6.0 5.5	7.4008 7.9295 8.5394 9.2510 10.0920	-1.1 -1.1 -1.1 -1.2 -1.3			$ \begin{array}{r} -1.2 \\ -0.6 \\ +0.1 \\ 0.9 \\ 1.7 \end{array} $	$\begin{array}{c} -2.2 \\ -1.6 \\ -0.8 \\ -0.1 \\ +0.8 \end{array}$	(+0.5) (1.1) (1.8) (2.6) (3.4)	
5.0 4.5 4.0 3.5 3.25	11.1012 12.3346 13.8765 15.8589 17.0788	-1.5 -1.6 -1.8			2.5 3.5 4.5 5.8 6.4	1.7 2.8 4.0 5.3 6.1	(4.4) (5.2) (6.2) (7.4)	
3.0	18.5020				6.9	6.8		•••••

^{*}Estimated.

TABLE VI. Heat capacity: sodium compounds

Φ_C, cal/deg mole, at 25 °C

				` `	·Ψ _C , ca	d/deg mole,	, at 25 C						
n	m	NaOH	NaCl	NaClO ₃	NaClO₄	NaBr	NaBrO ₃	NaI	NaNO ₂	NaNO ₃	NaCHO ₂	Na- C ₂ H ₃ O ₂	Na- C ₃ H ₃ O ₂
100,000 50,000 20,000 10,000	0.00 .000555 .00111 .00278 .00555	$\begin{array}{ c c c c c } -24.4 \\ -24.3 \\ -24.2 \\ -24.1 \\ -23.9 \end{array}$	-21.5 -21.4 -21.3 -21.2 -21.1	-8.6 -8.5 -8.4 -8.3 -8.2	-7.1 -7.0 -6.9 -6.8 -6.6	-22.8 -22.7 -22.6 -22.5 -22.4	-15.4 -15.3 -15.2 -15.1 -15.0	-22.9 -22.8 -22.7 -22.6 -22.5	-12.2 -12.1 -12.0 -11.9 -11.8	-9.6 -9.5 -9.4 -9.3 -9.1	-9.9 -9.8 -9.7 -9.6 -9.5	+9.6 9.7 9.8 9.9 10.0	+37.8 37.9 38.0 38.1 38.3
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	$ \begin{array}{c c} -23.8 \\ -23.7 \\ -23.6 \\ -23.4 \\ -23.1 \end{array} $	$\begin{array}{c} -21.0 \\ -20.9 \\ -20.8 \\ -20.7 \\ -20.4 \end{array}$	-8.1 -7.9 -7.8 -7.7 -7.4	-6.5 -6.3 -6.2 -5.9 -5.5	$\begin{array}{c} -22.3 \\ -22.2 \\ -22.1 \\ -22.0 \\ -21.8 \end{array}$	$\begin{array}{r} -14.8 \\ -14.7 \\ -14.6 \\ -14.4 \\ -14.1 \end{array}$	-22.4 -22.3 -22.2 -22.1 -21.9	-11.7 -11.5 -11.4 -11.3 -11.0	-9.0 -8.9 -8.8 -8.6 -8.3	-9.3 -9.2 -9.1 -8.9 -8.6	10.1 10.3 10.4 10.5 10.8	38.4 38.5 38.6 38.7 39.0
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	$\begin{array}{c c} -22.8 \\ -22.2 \\ -22.0 \\ -21.8 \\ -21.5 \end{array}$	-20.2 -19.8 -19.6 -19.5 -19.3	-7.1 -6.5 -6.3 -6.1 -5.7	-5.0 -4.1 -3.9 -3.5 -3.0	$\begin{array}{c} -21.6 \\ -21.2 \\ -21.0 \\ -20.9 \\ -20.7 \end{array}$	$\begin{array}{r} -13.7 \\ -13.1 \\ -12.9 \\ -12.6 \\ -12.3 \end{array}$	$ \begin{array}{r} -21.7 \\ -21.2 \\ -21.1 \\ -20.8 \\ -20.6 \end{array} $	$ \begin{array}{r r} -10.7 \\ -10.2 \\ -10.0 \\ -9.8 \\ -9.5 \end{array} $	-8.0 -7.4 -7.2 -7.0 -6.7	-8.3 -7.7 -7.6 -7.3 -7.0	11.1 11.7 11.8 12.0 12.3	39.2 39.7 39.8 39.9 40.2
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	$ \begin{array}{r} -21.2 \\ -20.7 \\ -20.0 \\ -18.9 \\ 17.1 \end{array} $	-19.0 -18.6 -18.2 -17.5 16.3	-5.4 -4.9 -4.1 -2.8 -0.7	$ \begin{array}{c c} -2.4 \\ -1.5 \\ -0.1 \\ +1.9 \\ 5.5 \end{array} $	-20.4 -20.0 -19.6 -18.8 -17.5	-11.8 -11.3 -10.4 -9.2 -7.1	-20.3 -19.9 -19.2 -18.3 -16.7	-9.2 -8.7 -7.9 -6.9 -5.1	$ \begin{vmatrix} -6.4 \\ -5.9 \\ -5.2 \\ -4.1 \\ -2.1 \end{vmatrix} $	-6.7 -6.2 -5.5 -4.4 -2.5	12.6 13.1 13.8 14.6 16.4	40.4 40.8 41.2 42.0 43.2
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877	-15.5 -12.9 -10.7 -7.1 -4.8	-15.2 -13.4 -11.7 -8.8 -7.0	+1.1 4.2 6.6 10.7 13.1	8.4 12.7 15.3 18.4 20.0	-16.5 -14.4 -12.7 -9.7 -7.7	$ \begin{array}{r} -5.2 \\ -2.1 \\ +0.5 \\ 4.9 \\ 7.4 \end{array} $	-15.3 -12.9 -11.0 -7.6 -5.4	$ \begin{array}{c c} -3.6 \\ -0.9 \\ +1.4 \\ 5.1 \\ 7.3 \end{array} $	$ \begin{array}{c c} -0.5 \\ +2.2 \\ 4.5 \\ 8.3 \\ 10.7 \end{array} $	$ \begin{array}{r} -1.0 \\ +1.6 \\ \hline 3.6 \\ \hline 6.8 \\ \hline 8.7 \end{array} $	17.9 20.5 22.7 26.4 28.5	44.2 45.8 46.9 48.6 49.6
30 25 20 15 12	1.8502 2.2202 2.7753 3.7004 4.6255	$ \begin{array}{r} -1.7 \\ +0.4 \\ 3.2 \\ 6.8 \\ 9.8 \end{array} $	$ \begin{array}{c c} -4.6 \\ -2.4 \\ 0.0 \\ +3.3 \\ 6.3 \end{array} $	16.4 18.4 20.7 23.4 25.3	22.3 23.8 25.9 28.4 30.1	$ \begin{array}{r} -4.9 \\ -3.0 \\ -0.4 \\ +2.9 \\ \hline 5.8 \end{array} $	10.8	$ \begin{array}{r} -2.4 \\ -0.4 \\ +2.1 \\ 5.6 \\ \dots \end{array} $	10.4 12.6 15.3 19.1	14.2 16.4	11.6 13.6 15.8 18.6	31.1 32.7 34.6 36.9 38.4	50.8 51.7
10 9.5 9.0 8.5 8.0	5.5506 5.8427 6.1674 6.5301 6.9383	12.2 12.8 13.5 14.3 15.1	9.0 9.7 10.5	26.5 26.9 27.2 27.5 27.8	31.2 31.4 31.7 32.0 32.3	8.2 9.0 9.8 10.7 11.6						39.5 39.9 40.2 40.6 40.9	
7.5 7.0 6.5 6.0 5.5	7.4008 7.9295 8.5394 9.2510 10.0920	15.8 16.6 17.5 18.4 19.1		28.6	32.6 32.9 33.3 33.6 34.0	12.7 13.8 15.0 16.3						41.4	
5.0 4.5 4.0 3.5 3.25	11.1012 12.3346 13.8765 15.8589 17.0788	19.8 20.6 21.3 22.0 22.2			34.4 34.8 35.2 35.5 35.6								
3.0 2.5 2.0	18.5020 22.2024 27.7530	22.3 22.2 21.2		•••••								······································	

TABLE VII. Heat capacity: potassium compounds

 $\Phi_{\it C}$, cal/deg mole, at 25 °C

				TC, Calluck I						
n	m ·	КОН	KCl	KBr	KI	KIO ₃	KNO ₃	KC ₂ H ₃ O ₂	KCNS	KMnO ₄
100,000 50,000 20,000 10,000	0.00 .000555 .00111 .00278 .00555	-30.3 -30.2 -30.1 -30.0 -29.9	-27.4 -27.3 -27.2 -27.1 -27.0	-28.7 -28.6 -28.5 -28.4 -28.3	-28.8 -28.7 -28.6 -28.5 -28.4	-24.4 -24.2 -24.1 -24.0 -23.6	-15.5 -15.3 -15.3 -15.1 -15.0	+3.7 3.8 3.9 4.0 4.1	$\begin{array}{r r} -4.4 \\ -4.2 \\ -4.2 \\ -4.1 \\ -4.0 \end{array}$	$\begin{array}{c c} -14.4 \\ -14.3 \\ -14.2 \\ -14.1 \\ -13.9 \end{array}$
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	-29.8 -29.7 -29.6 -29.5 -29.2	-26.9 -26.8 -26.7 -26.6 -26.4	-28.2 -28.1 -28.0 -27.9 -27.6	-28.3 -28.2 -28.1 -27.9 -27.7	$ \begin{array}{r} -23.5 \\ -23.3 \\ -23.1 \\ -22.7 \\ -21.7 \end{array} $	$\begin{array}{r} -14.9 \\ -14.8 \\ -14.7 \\ -14.5 \\ -14.2 \end{array}$	4.2 4.3 4.4 4.5 4.7	$ \begin{array}{r} -3.9 \\ -3.8 \\ -3.7 \\ -3.6 \\ -3.3 \end{array} $	-13.8 -13.6 -13.5 -13.3 -12.9
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	$ \begin{array}{r} -29.0 \\ -28.7 \\ -28.6 \\ -28.4 \\ -28.2 \end{array} $	$ \begin{array}{r} -26.2 \\ -25.9 \\ -25.7 \\ -25.6 \\ -25.4 \end{array} $	-27.4 -27.0 -26.9 -26.7 -26.6	-27.5 -27.0 -26.9 -26.7 -26.5	-21.3 -19.8 -19.6 -18.7 -17.9	-13.8 -13.3 -13.1 -12.8 -12.5	4.9 5.3 5.4 5.5 5.7	$ \begin{array}{r} -3.1 \\ -2.8 \\ -2.7 \\ -2.6 \\ -2.4 \end{array} $	-12.5 -11.6 -11.4 -11.0 -10.5
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	-28.0 -27.6 -27.2 -26.5 -25.3	-25.3 -25.0 -24.7 -24.1 -23.3	-26.4 -26.1 -25.7 -25.1 -24.0	-26.3 -26.0 -25.5 -24.8 -23.5	-16.8 -15.2 -12.9 -9.3 -3.6	-12.2 -11.7 -11.0 -9.9 -8.1	5.9 6.2 6.7 7.4 8.6	$ \begin{array}{r} -2.2 \\ -1.9 \\ -1.5 \\ -1.0 \\ 0.0 \end{array} $	-9.9 -9.2 -8.0 -6.2 -2.7
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877	-24.3 -22.6 -21.1 -18.6 -16.9	$\begin{array}{c} -22.6 \\ -21.2 \\ -20.0 \\ -17.8 \\ -16.4 \end{array}$	$ \begin{array}{r} -23.1 \\ -21.6 \\ -20.4 \\ -18.3 \\ -17.0 \end{array} $	-22.3 -20.4 -18.8 -16.3 -14.8	+1.4	$ \begin{array}{r} -6.4 \\ -3.8 \\ -1.6 \\ +2.1 \\ 4.4 \end{array} $	9.8 11.8 13.6 16.5 18.2	+0.8 2.0 3.2 4.9 6.0	+0.1 4.7
30 25 20 15 12	1.8502 2.2202 2.7753 3.7004 4.6255	-14.4 -12.5	$\begin{array}{r} -14.3 \\ -12.8 \\ -10.8 \\ -8.1 \\ -6.0 \end{array}$	-14.9 -13.4 -11.4 -8.7 -6.5	-12.7 -11.2 -9.3		7.9 10.3 13.5 17.7	20.4 21.9 23.5 25.3 26.5	7.6 8.7 10.2	
10 9.5 9.0 8.5 8.0	5.5506 5.8427 6.1674 6.5301 6.9383		-4.2 	-4.6 -4.2				27.5 27.8 28.1 28.5 28.8		
7.5 7.0 6.5 6.0 5.5	7.4008 7.9295 8.5394 9.2510 10.0920							29.2 29.6 30.0 30.4 30.9		
5.0	11.1012							31.5		

Table VIII. Heat capacity: rubidium compounds

 $\Phi_{\mathcal{C}}$, cal/deg mole, at 25 °C

n	m	RbOH*	RbCl*	RbBr*	RbI*	RbNO₃*
00,000 50,000 20,000 10,000	0.00 .000555 .00111 .00278 .00555	(-32.2) (-32.1) (-32.0) (-31.9) (-31.8)	(-29.3) (-29.2) (-29.1) (-29.0) (-28.9)	(-30.6) (-30.5) (-30.4) (-30.3) (-30.2)	(-30.7) (-30.6) (-30.5) (-30.4) (-30.3)	(-17.4) (-17.3) (-17.2) (-17.1) (-17.0)
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	(-31.7) (-31.6) (-31.5) (-31.4) (-31.1)	$\begin{array}{c} (-28.8) \\ (-28.7) \\ (-28.6) \\ (-28.5) \\ (-28.3) \end{array}$	(-30.1) (-30.0) (-29.9) (-29.8) (-29.6)	(-30.2) (-30.1) (-30.0) (-29.9) (-29.7)	(-16.9) (-16.8) (-16.7) (-16.5) (-16.2)
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	(-30.9) (-30.5) (-30.4) (-30.2) (-30.0)	(-28.2) (-28.0) (-27.9) (-27.8) (-27.6)	(-29.4) (-29.2) (-29.1) (-29.0) (-28.8)	(-29.5) (-29.1) (-29.0) (-28.9) (-28.7)	(-15.9) (-15.3) (-15.1) (-14.9) (-14.6)
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	(-29.8) (-29.5) (-29.1) (-28.4) (-27.3)	(-27.5) (-27.3) (-27.0) (-26.6) (-25.8)	(-28.7) (-28.5) (-28.2) (-27.7) (-26.9)	(-28.5) (-28.2) (-27.8) (-27.3) (-26.3)	(-14.3) (-13.8) (-13.0) (-12.0) (-10.1)
150 100 75 50	.3700 .5551 .7401 1.1101	(-26.2) (-24.4) (-22.9) (-20.2)	(-25.1) (-24.0) (-23.1) (-21.5)	(-26.2) (-25.0) (-24.0)	(-25.4) (-23.9) (-22.6) (-20.3)	(-8.6) (-5.8) (-3.6) (+0.2)

^{*}Estimated.

TABLE IX. Heat capacity: cesium compounds

 $\Phi_{\it C}$, cal/deg mole, at 25 °C

n	m	CsOH*	CsCl*	CsBr*	CsI	CsNO ₃ *
100,000 50,000 20,000 10,000	0.00 .000555 .00111 .00278 .00555	- 33.6 (-33.5) (-33.5) (-33.4) (-33.3)	-30.7 (-30.6) (-30.5) (-30.4) (-30.3)	-32.0 (-31.9) (-31.8) (-31.7) (-31.6)	-32.1 -32.0 -31.9 -31.8 -31.7	- 18.8 (- 18.7) (- 18.6) (- 18.5) (- 18.4)
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	(-33.2) (-33.1) (-33.0) (-32.9) (-32.7)	(-30.3) (-30.2) (-30.1) (-30.0) (-29.8)	(-31.5) (-31.4) (-31.3) (-31.2) (-31.0)	$ \begin{array}{r} -31.6 \\ -31.5 \\ -31.4 \\ -31.3 \\ -31.1 \end{array} $	(-18.3) (-18.2) (-18.1) (-17.9) (-17.6)
1,500 1,000 900 800 700	.03700 .05551 .0617 .0694 .0793	$ \begin{array}{c} (-32.4) \\ (-32.1) \\ (-31.9) \\ (-31.8) \\ (-31.6) \end{array} $	(-29.6) (-29.3) (-29.2) (-29.1) (-29.0)	(-30.8) (-30.5) (-30.4) (-30.3) (-30.2)	$ \begin{array}{r} -30.9 \\ -30.6 \\ -30.5 \\ -30.4 \\ -30.2 \end{array} $	(-17.3) (-16.7) (-16.5) (-16.3) (-16.0)
600 500 400 300 200	.0925 .1110 .1388 .1850 .2775	(-31.4) (-31.1) (-30.7) (-30.1) (-29.1)	(-28.8) (-28.6) (-28.3) (-27.9) (-27.2)	(-30.0) (-29.8) (-29.5) (-29.1) (-28.3)	-30.0 -29.7 -29.3 -28.7 -27.7	(-15.7) (-15.2) (-14.4) (-13.4) (-11.5)
150 100 75 50 40	.3700 .5551 .7401 1.1101 1.3877	(-28.0) (-26.4) (-25.0) (-22.3) (-20.6)	(-26.6) (-25.5) (-24.6) (-23.1) (-22.1)	(-27.6) (-26.5) (-25.5) (-23.8) (-22.7)	26.9 -25.5 -24.5 -22.6 -21.4	(-9.7) (-7.2) (-5.0) (-1.2) (+1.2)
30 25	1.8502 2.2202	(-17.5) (-15.7)			-19.7 -18.3	

^{*}Estimated.

TABLE X. Thermal properties of aqueous NaCl at 25 °C

n	m	$d\Phi_L/dm^{1/2}$	Φ_L	$ar{L}_2$
			cal/mole	cal/mole
∞	0.00	472	0	0
100,000	.000555	436	11	16
50,000	.00111	423	14	21
20,000	.00278	395	22	32
10,000	.00555	365	31	45
7,000	.00793	344	37	52
5,000	.01110	322	42	59
4,000	.01388	304	46	64
3,000	.01850	282	51	70
2,000	.02775	245	59	79
1,000	.05551	167	73	93
900	.0617	155	75	94
800	.0694	139	78	96
700	.0793	120	80	97
600	.0925	99	83	98
500	.1110	71	85	97
400	.1388	+35	87	94
300	.1850	-14	88	85
200	.2775	-87	83	+60
100	.5551	220	49	-33
75	.7401	-275	+20	-98
50	1.1101	-344	-41	-222
40	1.3877	-373	-85	-304
30	1.8502	-394	- 156	- 424
25	2.2202	-394	-207	-501
20	2.7753	-376	-274	-587
15	3.7004	-321	-367	-676
12	4.6255	-242	430	-690
10	5.5506	-135	-469	-628
9.5	5.8427	-93	-476	-588
			L	

TABLE XI. Heat of dilution: acids

					· · · · · · · · · · · · · · · · · · ·				
n	m	HF	HCl	HClO₄	HBr	ні	HNO ₃	CH ₂ O ₂	C ₂ H ₄ O ₂
∞	0.00	0	0	0	0	0	0	0	0
500,000	.000111	300	5	5	5	5	5	9	40
100,000	.000555	900	10	10	9	9	11	13	50
50,000	.00111	1,300	16	14	13	12	15	20	53
20,000	.00278	1,800	25	22	22	20	23	23	55
10,000	.00555	2,130	34	30	31	29	31	25	58
7,000		2,250	40	35	37	34	36	26	59
5,000	.01110	2,360	47	40	44	41	42	26	61
4,000	.01388	2,450	54	43	49	46	46	27	62
3,000	.01850	2,550	60	47	56	52	51	28	62
2,000	.02775	2,700	74	54	68	63	59	28	63
1,500	.03700	2,812	85	58	77	71	65	29	64
1,110	.05000	2,927	97	62	89	81	73	29	65
1,000	.05551	2,969	102	62	92	84	76	29	65
900	.0617	2,989	107	63	97	88	78	30	66
800	.0694	3,015	113	64	102	92	81	31	67
700	.0793	3,037	120	65	108	96	84	32	68
600	.0925	3,057	129	65	115	102	88	32	68
555.1	.1000	3,060	133	65	119	105	89	32	69
500	.1110	3,077	140	65	124	108	92	32	70
400	.1388	3,097	156	64	135	116	97	33	72
100		0,051	1.50	, J	100	1.10	1 ''	1 00	

TABLE XI. Heat of dilution: acids-Continued

-				ΨL, car/mor	s, at 25 G		·			
	n	m	HF	HCl	HClO₄	HBr	ні	HNO ₃	CH ₂ O ₂	C ₂ H ₄ O ₂
	300	.1850	3,126	176	61	150	125	103	34	76
	277.5	.2000	3,129	182	59	155	128	105	35	79
	200	.2775	3,142	212	50	176	140	117	36	82
	150	.3700	3,148	242	36	197	154	118	39	88
	111.0	.5000	3,156	280	18	225	170	119	42	97
	100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	3,160 3,167 3,179 3,184 3,192	295 343 405 431 493	+12 -14 -48 -61 -91	235 270 314 331 379	176 194 223 234 260	120 121 121 121 121 121	44 49 54 56 60	101 113 130 147 155
	37.00	1.5000	3,194	518	-103	398	269	121	62	162
	30	1.8502	3,200	595	-138	455	301	124	65	183
	27.75	2.0000	3,203	627	-149	477	315	126	66	192
	25	2.2202	3,208	674	-162	510	336	130	67	204
	22.20	2.5000	3,211	732	-173	550	365	139	68	218
	20	2.7753	3,214	792	-182	590	396	149	69	233
	18.50	3.0000	3,216	838	-187	624	427	159	69	245
	15.86	3.5000	3,221	946	-196	709	503	189	69	268
	15	3.7004	3,227	988	-195	743	536	203	69	277
	13.88	4.0000	3,234	1,052	-188	796	588	229	69	291
	12.33	4.5000	3,246	1,171	-175	887	676	265	69	313
	12	4.6255	3,249	1,190	-170	911	700	277	69	318
	11.10	5.0000	3,256	1,271	-150	983	764	313	69	333
	10	5.5506	3,265	1,396	-117	1,097	855	368	68	353
	9.5	5.8427	3,269	1,462	-97	1,156	920	400	68	363
	9.251	6.0000	3,272	1,498	-84	1,196	950	418	67	368
	9.0	6.1674	3,274	1,535	-72	1,230	980	437	67	373
	8.5	6.5301	3,278	1,618	-40	1,313	1,050	480	66	383
	8.0	6.9383	3,282	1,710	+4	1,401	1,115	530	65	392
	7.929	7.0000	3,283	1,725	11	1,416	1,130	538	65	394
	7.5	7.4008	3,286	1,820	61	1,497	1,210	595	63	402
	7.0	7.9295	3,290	1,942	135	1,600	1,325	661	61	411
	6.938	8.0000	3,291	1,960	146	1,622	1,340	667	61	412
	6.5	8.5394	3,296	2,090	229	1,738	1,450	745	58	420
	6.167	9.0000	3,302	2,202	306	1,845	1,570	805	55	426
	6.0	9.2510	3,305	2,265	348	1,903	1,630	840	53	429
	5.551	10.0000	3,316	2,447	481	2,078	1,820	940	49	436
	5.5	10.0920	3.317	2,472	499	2,102	1,850	950	49	437
	5.0	11.1012	3,335	2,721	730	2,344	2,100	1,098	43	445
	4.5	12.3346	3,362	3,025	1,144	2,655	2,460	1,270	37	453
	4.0	13.8765	3,400	3,404	1,574	3,089	2,960	1,495	29	462
	3.700	15.0000	3,428	3,680	1,893	3,415	3,350	1,645	26	469
	3.5	15.8589	3,450	3,882	2,150	3,668	3,660	1,770	21	473
	3.25	17.0788	3,483	4,160	2,460	4,005	4,110	1,920	17	481
	3.0	18.5020	3,520	4,460	2,880	4,370	4,630	2,101	13	488
	2.775 2.5 ·2.0 1.5 1.0	20.0000 22.2024 27.7530 37.0040 55.506	3,557 3,607 3,712	4,750 5,180 6,260 8,240 10,900	3,300 4,000 5,500	4,760 5,300 6,650 8,530 11,670	5,190 6,000 	2,270 2,520 3,060 3,770 4,715	9 +4 -5 -13 +11	496 506 528 532 518
	0.5 0.25	111.012 222.02							77 129	495

Table XII A. Heat of dilution: ammonium compounds $\Phi_L, \, cal/mole, \, at \, \, 25 \, \, ^{\circ}C$

			- 27 7					
n	m	NH₄OH	NH ₃	NH₄Cl	NH₄Br*	NH₄I*	NH ₄ NO ₃	NH ₄ C ₂ H ₃ O ₂
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 584 722 761 800 815	0 -584 -722 -761 -800 -815	0 5 10 14 22 29	0 (5) (10) (14) (22) (29)	0 (5) (10) (13) (21) (28)	0 5 10 14 22 28	0 82 86 95 105
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	- 824 - 829 - 833 - 838 - 841	824 829 833 838 841	34 40 44 48 57	(33) (39) (43) (46) (55)	(32) (38) (42) (44) (53)	32 36 39 43 48	112 119 125 134 147
1,500 1,110 1,000 900 800	.03700 .05000 .05551 .0617 .0694	- 846 846 846 846 846	-844 -846 -846 -846 -846	63 70 73 76 79	(61) (65) (70) (73) (74)	(59) (63) (68) (71) (72)	50 52 53 53 53	159 174 180 185 193
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	- 847 - 847 - 847 - 848 - 849	847 847 847 848 849	82 87 88 92 98	(76) (80) (82) (86) (90)	(73) (74) (76) (80) (82)	52 51 49 47 41	201 212 217 225 243
300 277.5 200 150 111.0	.1850 .2000 .2775 .3700 .5000	848 848 846 845 843	848 848 845 844 842	105 107 117 124 131	(95) (96) (100) (105) (106)	(85) (84) (80) (78) (77)	27 +23 -2 -36 -88	272 279 315 355 404
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	-843 -840 -835 -834 -830	-842 -838 -833 -831 -827	133 136 136 136 134	(106)	(76) 	-110 -182 -270 -308 -396	423 480 551 579 644
37.00 30 27.75 25 22.20	1.5000 1.8502 2.0000 2.2202 2.5000	-824 -818 -816 -814 -808	-821 -815 -813 -811 -805	133 128 125 120 114			442 535 575 636 704	699 740 769 810 850
20 18.50 15.86 15 13.88	2.7753 3.0000 3.5000 3.7004 4.0000	803 798 790 787 782	800 795 786 783 777	108 104 93 89 82			-770 -810 -910 -950 -1,000	895 930 1,000 1,040 1,069
12.33 12 11.10 10 9.5	4.5000 4.6255 5.0000 5.5506 5.8427	774 772 766 756 754	768 765 759 749 744	72 69 62 53 48			-1,085 -1,106 -1,165 -1,240 -1,285	1,130 1,145 1,190 1,250 1,290
9.251 9.0 8.5 8.0 7.929	6.0000 6.1674 6.5301 6.9383 7.0000	- 752 - 749 - 744 - 737 - 736	-741 -737 -730 -721 -720	45 42 36 30 29			-1,308 -1,330 -1,375 -1,430 -1,436	1,310 1,330 1,370 1,425 1,434

^{*}Footnote at end of table.

TABLE XII A. Heat of dilution: ammonium compounds - Continued

								· · · · · · · · · · · · · · · · · · ·
n	m	NH₄OH	NH ₃	NH₄Cl	NH₄Br*	NH₄I*	NH ₄ NO ₃	NH ₄ C ₂ H ₃ O ₂
7.5 7.0 6.938 6.5 6.167 6.0 5.551 5.5	7.4008 7.9295 8.0000 8.5394 9.0000 9.2510 10.0000 10.0920 11.1012	-730 -721 -720 -711 -703 -698 -686 -684 -667	- 711 - 698 - 696 - 684 - 672 - 667 - 648 - 646				-1,485 -1,545 -1,545 -1,605 -1,650 -1,675 -1,738 -1,745 -1,820	1,478 1,536 1,542 1,600 1,646 1,670 1,732 1,740 1,810
4.5 4.0 3.700	12.3346 13.8765 15.0000	-646 -620 -602	-588 -548 -520				-1,900 -1,988 -2,048	1,884 1,950 1,986
3.5 3.25 3.0	15.8589 17.0788 18.5020	- 588 - 570 - 548	-498 -471 -438				$ \begin{array}{r} -2,090 \\ -2,150 \\ -2,255 \end{array} $	2,008 2,034 2,060
2.775 2.5 2.0 1.5 1.0	20.0000 22.2024 27.7530 37.0040 55.506	530 498 438 358 235	-405 -358 -235 -39 +314				-2,320 	2,082 2,110 2,168

^{*}Estimated.

TABLE XII B. Heat of dilution: alkyl ammonium compounds

n	m	(CH ₃)NH ₃ Cl	(CH ₃) ₂ NH ₂ Cl	(CH ₃) ₃ NHCl	N(CH ₃) ₄ Cl	NH ₂ CH ₃	NH(CH ₃) ₂	N(CH ₃) ₃
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 5 10 14 23 30	0 5 10 14 23 30	0 5 10 13 21 26	0 5 10 13 20 23			
7,000 5,000 4,000 3,000 2,000 1,500 1,110 1,000 900 800	.00793 .01110 .01388 .01850 .02775 .03700 .05000 .05551 .0617	36 41 45 50 61 68 76 79 82 85	34 39 43 48 56 61 68 70 72 75	30 34 36 39 44 44 45 45 45	25 27 27 27 24 19 14 10 7 +1	0 20 50	 0	 0
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	89 93 96 99 105	77 81 82 84 88	44 43 42 41 38	-5 -13 -18 -24 -41	80 110 120 130 160	100	100 200
277.5 200 150 111.0	.1650 .2000 .2775 .3700 .5000	115 115 123 131 139	92 94 98 102 106	29 18 +2 -13	-07 -75 -110 -158 -211			
100 75 55.51 50	.5551 .7401 1.0000 1.1101	143 152 164 167	107 112 119 121	- 24 - 46 - 72 - 82	-232 	 		

TABLE XIII. Heat of dilution: silver and lithium compounds

				Φ_L , cal/mo						
n	m	AgF	AgNO ₃	LiOH	LiF	LiCl	LiClO₄	LiBr	LiI*	LiNO ₃
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555		0 4 9 12 17 19	0 5 11 16 25 35	0 5 11 16 25 35	0 5 10 14 23 32	0 5 10 14 23 31	0 5 10 13 21 29	0 (5) (10) (13) (20) (28)	0 5 10 13 20 28
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775		21 21 21 20 16	42 49 55 62 74	42 49 54 62 76	37 43 48 54 65	37 43 48 55 65	34 40 44 50 60	(32) (38) (42) (51) (55)	33 38 42 49 58
1,500 1,110 1,000 900 800	.03700 .05000 .05551 .0617 .0694		12 5 +2 -1 -6	84 97 102 106 112	87 101 105 110	73 82 86 89 94	73 82 85 88 92	67 76 80 83 86	(62) (71) (74) (77) (78)	65 74 76 80 83
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388		-13 -20 -25 -32 -49	120 127 132 139 153		99 105 108 113 124	97 103 106 110 119	92 97 100 105 115	(85) (89) (92) (97) (106)	88 93 96 100 107
300 277.5 200 150 111.0	.1850 .2000 .2775 .3700 .5000	1 1 2 3 5	-79 -88 -137 -196 -274	175 181 211 240 275		140 144 165 186 213	133 137 154 170 186	130 134 155 176 202	(120) (124) (145) (166) (192)	120 123 139 153 171
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	6 9 12 14 18	307 408 536 585	289 329 378 396 440		223 255 296 312 352	192 207 230 240 263	212 243 280 295 330	(202) (232) (267) (280)	177 194 212 218 234
37.00 30 27.75 25 22.20	1.5000 1.8502 2.0000 2.2202 2.5000	20 25 27 31 36		457 508 528 558 597		367 415 435 465 504	272 300 313 334 361	344 386 403 429 463		240 254 261 271 281
20 18.50 15.86 15 13.88	2.7753 3.0000 3.5000 3.7004 4.0000	41 45 54 58 64		635 667 738 766 810		542 575 645 674 720	390 414 469 491 523	496 524 588 614 652		*(293) (302) (329) (342) (363)
12.33 12 11.10 10 9.5	4.5000 4.6255 5.0000 5.5506 5.8427	72 76 87 112 128		882 900 		799 820 880 974 1,025		716 733 785 864 905		(403) (415) (450) (500) (530)
9.251 9.0 8.5 8.0 7.929	6.0000 6.1674 6.5301 6.9383 7.0000	138 144 172 200 204				1,054 1,083 1,148 1,225 1,236		930 953 1,009 1,069 1,079		(550) (570) (608) (660) (670)

^{*}Footnote at end of table.

TABLE XIII. Heat of dilution: silver and lithium compounds - Continued

n	m	AgF	$ m AgNO_3$	LiOH	LiF	LiCl	LiClO ₄	LiBr	LiI*	LiNO ₃
7.5 7.0 6.938 6.5 6.167 6.0 5.551 5.5	7.4008 7.9295 8.0000 8.5394 9.0000 9.2510 10.0000 10.0920	238 276 282 329 373 398 474 484				1,313 1,416 1,431 1,542 1,640 1,694 1,858 1,880		1,140 1,228 1,238 1,332 1,415 1,463 1,611 1,630		(720) (790) (800) (870) (920) (970) (1,070) (1,080)
5.0 4.5	11.1012 12.3346	600 748				2,115 2,411		1,848 2,147		(1,230)
4.0 3.700 3.5 3.25 3.0	13.8765 15.0000 15.8589 17.0788 18.5020	945				2,794 3,070 3,282 3,573 3,893		2,571 2,907 3,171 3,539		

^{*}Estimated.

Table XIV A. Heat of dilution: sodium compounds

	Φ _L , cal/mole, at 25 °C											
n	m	NaOH	NaF*	NaCl	NaClO ₃	NaClO ₄	NaBr .	NaBrO ₃				
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 5 11 15 23 32	0 (5) (11) (15) (23) (32)	0 5 11 14 22 31	0 5 10 14 21 27	0 5 10 14 21 27	0 5 11 14 21 30	0 5 10 14 20 26				
7,000	.00793	38	(38)	37	32	31	35	30				
5,000	.01110	44	(44)	42	36	35	40	33				
4,000	.01388	49	(49)	46	39	38	44	36				
3,000	.01850	55	(55)	51	42	41	49	38				
2,000	.02775	65	(65)	59	45	44	56	40				
1,500	.03700	72	(72)	64	47	46	60	39				
1,110	.05000	81	(81)	71	47	46	66	37				
1,000	.05551	84	(84)	73	47	44	68	36				
900	.0617	87	(87)	75	47	44	69	35				
800	.0694	90	(90)	78	46	42	71	34				
700	.0793	95	(95)	80	45	42	72	32				
600	.0925	99	(99)	83	42	38	74	28				
555.1	.1000	101	(101)	84	40	36	75	26				
500	.1110	104	(104)	85	37	32	75	22				
400	.1388	110	(110)	87	29	22	76	+12				
300	.1850	116	(115)	88	11	+4	74	*(-6)				
277.5	.2000	117	(116)	87	+7	-2	73	(-13)				
200	.2775	121	(119)	83	-22	-37	66	(-46)				
150	.3700	118	(115)	74	-58	-80	54	(-88)				
111.0	.5000	115	(110)	57	-109	-140	32	(-145)				
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	112 99 78 67 44	(107) (92) (69) (57)	49 + 20 - 23 - 41 - 85	-130 -200 -293 -333 -425	167 250 360 404 508	+23 -10 -59 -81 -134	(-170) (-247) (-350) (-388)				

^{*}Footnote at end of table.

TABLE XIV A. Heat of dilution: sodium compounds-Continued

Φ_L, cal/mole, 25 °C

Φ _L , cal/mole, 25 °C											
n	m	NaOH	NaF*	NaCl	NaClO ₃	NaClO ₄	NaBr	NaBrO ₃			
37.00 30 27.75 25 22.20	1.5000 1.8502 2.0000 2.2202 2.5000	36 13 +4 -4 -14		- 104 156 172 207 242	-462 -570 -613 -675 -747	- 551 - 672 - 719 - 787 - 866	155 219 245 282 326				
20 18.50 15.86 15 13.88	2.7753 3.0000 3.5000 3.7004 4.0000	- 20 - 22 - 16 - 10 0		-274 -299 -349 -367 -390	-814 -866 -973 -1,012 -1,068	-945 -1,004 -1,133 -1,179 -1,243	-368 -400 -465 -489 -523				
12.33 12 11.10 10 9.5	4.5000 4.6255 5.0000 5.5506 5.8427	+ 26 34 61 112 142		-423 -430 -449 -469 -476	$\begin{array}{c} -1,156 \\ -1,177 \\ -1,234 \\ -1,310 \\ -1,348 \end{array}$	-1,339 -1,360 -1,424 -1,508 -1,549	-574 -586 -618 -659 -677				
9.251 9.0 8.5 8.0 7.929	6.0000 6.1674 6.5301 6.9383 7.0000	161 181 229 289 299		-479 -480	-1,369 $-1,389$ $-1,430$ $-1,473$ $-1,479$	-1,568 -1,590 -1,632 -1,676 -1,683	-686 -694 -711 -725 -726				
7.5 7.0 6.938 6.5 6.167	7.4008 7.9295 8.0000 8.5394 9.0000	364 457 469 575 672			-1,518 -1,566 -1,574 -1,621 -1,658	-1,725 -1,777 -1,783 -1,833 -1,872	-735 -741 -741 -738 -730				
6.0 5.551 5.5 5.0 4.5	9.2510 10.0000 10.0920 11.1012 12.3346	727 903 926 1,178 1,513			-1,678 	-1,892 -1,945 -1,951 -2,006 -2,058					
4.0 3.700 3.5 3.25 3.0	13.8765 15.0000 15.8589 17.0788 18.5020	1,955 2,290 2,543 2,903 3,307				-2,109 -2,139 -2,159 -2,185					
2.775 2.5	20.0000 22.2024	3,722 4,260	•••••								

^{*}Estimated.

 $TABLE\ XIV\ B.\quad \textit{Heat of dilution: sodium compounds}$

	r	, 			-		
n	m	NaI	NaIO ₃	NaNO ₂	NaNO ₃	NaC ₂ H ₃ O ₂	NaCNS
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 5 11 14 21 30	0 5 10 12 17 21		0 5 11 15 22 28	0 5 12 16 25 35	0 5 11 14 22 29
7,000	.00793	34	22		32	41	33
5,000	.01110	39	22		35	48	38
4,000	.01388	43	21		37	54	41
3,000	.01850	48	20		40	62	46
2,000	.02775	54	13		44	75	51
1,500	.03700	58	+6		44	86	54
1,110	.05000	64	-5		44	91	58
1,000	.05551	66	-11		44	104	58
900	.0617	67	-15		43	109	59
800	.0694	69	-23		41	116	60
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	70 71 72 72 71	- 33 46 54 65		39 35 31 29 + 18	124 132 137 143 159	59 58 57 55 50
300	.1850	68		40	-2	180	43
277.5	.2000	66		+30	-8	187	40
200	.2775	55		-40	-43	215	+21
150	.3700	39		-100	-86	243	-3
111.0	.5000	12		-170	-147	274	-39
100	.5551	+1		-190	-173	286	-55
75	.7401	-39		-250	-255	318	-107
55.51	1.0000	-97		-310	-367	358	-179
50	1.1101	-122		-330	-412	372	-208
40	1.3877	-184		-375	-519	407	-277
37.00	1.5000	- 209	G.	-390	-560	421	-306
30	1.8502	- 283		-420	-678	459	-384
27.75	2.0000	- 313		-425	-724	474	-415
25	2.2202	- 358		-450	-791	494	-460
22.20	2.5000	- 410		-475	-869	522	-513
20	2.7753	-462		490	-939	556	- 564
18.50	3.0000	-501		510	-994	564	- 603
15.86	3.5000	-582		540	-1,134	620	- 686
15	3.7004	-613		545	-1,144	645	- 718
13.88	4.0000	-656		560	-1,200	675	- 763
12.33 12 11.10 10 9.5	4.5000 4.6255 5.0000 5.5506 5.8427	- 719 - 734 - 775 - 828 - 854		585 590 605 630 640	-1,284 $-1,303$ $-1,359$ $-1,431$ $-1,464$	732 750 794 870 914	- 834 - 852 - 901 - 966 - 997
9.251	6.0000	-866		-645	-1,483	936	-1,014
9.0	6.1674	-878		-650	-1,501	966	-1,029
8.5	6.5301	-902		-665	-1,536	1,016	-1,060
8.0	6.9383	-924		-675	-1,573	1,100	-1,092
7.929	7.0000	-927		-680	-1,578	1,108	-1,096

TABLE XIV B. Heat of dilution: sodium compounds - Continued

n	m	NaI	NaIO ₃	NaNO ₂	NaNO ₃	NaC ₂ H ₃ O ₂	NaCNS
7.5	7.4008	-943		-690	-1,608	1,180	-1,122
7.0	7.9295	-957		-700	-1,644	1,280	-1,149
6.938	8.0000	-959		-710	-1,648	1,290	-1,152
6.5	8.5394	-964		-720	-1,678	1,394	-1,173
6.167	9.0000	-963		- 735	-1,699	1,480	-1,187
6.0	9.2510	959		740	-1,708	1,530	-1,193
5.551	10.0000	941		-760		1,674	-1,201
5.5	10.0920	-938	l	-765		1,692	-1,202
5.0	11.1012	-891		 785		1,890	-1,197
4.5	12.3346	-805		-810		2,120	-1,175
4.0	13.8765					2,410	-1,144
3.700	15.0000					2,612	-1,113
3.5	15.8589					2,760	
3.25	17.0788					2,980	
3.0	18.5020				l	3,230	

TABLE XV A. Heat of dilution: potassium compounds

Φ_L, cal/mole, at 25 °C

Φ_{L_s} cal/mole, at 25 °C											
n	m	КОН	KF	KCl	KClO ₃	KClO ₄	KBr	KBrO ₃ *			
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 5 11 16 23 32	0 5 11 15 23 32	0 5 10 14 22 31	0 5 10 13 20 24	0 4 9 12 15	0 5 10 14 21 29	0 (5) (10) (13) (20) (24)			
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	38 44 50 56 67	38 44 49 56 67	36 41 45 50 57	27 28 29 30 27	16 14 11 +5 -9	34 38 42 46 53	(26) (26) (25) (24) (21)			
1,500 1,110 1,000 900 800	.03700 .05000 .05551 .0617 .0694	75 86 89 92 97	75 86 89 92 96	62 68 70 72 74	20 17 14 11 +5	-22 -40 -49 -58 -69	57 61 63 64 65	(17) (8) (4) (0) (-6)			
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	102 108 111 115 124	101 107 110 113 121	77 79 80 82 84	-2 -12 -14 -27 -50	-84 -103 -113	66 67 67 67 67	(-13) (-24) (-30) (-40)			
300 277.5 200 150 110.0	.1850 .2000 .2775 .3700 .5000	134 138 151 161 172	132 135 146 154 162	85 85 81 72 57	-94 -108 -193 -310 -480		64 62 53 38 13				
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	176 186 200 204 218	165 171 178 181 187	50 +23 -15 -32 -73	-546 		+2 -37 -96 -120 -179				
37.00 30 27.75 25 22.20	1.5000 1.8502 2.0000 2.2202 2.5000	224 240 246 256 268	189 195 198 203 208	-89 -139 -162 -190 -228			- 203 - 273 - 302 - 343 - 393				

^{*}Footnote at end of table.

TABLE XV A. Heat of dilution: potassium compound-Continued

			TE, can more					
n	m	КОН	KF	KCl	KClO ₃	KClO ₄	KBr	KBrO ₃ *
20	2.7753	283	213	-263			-440	
18.50	3.0000	299	218	-289			-476	
15.86	3.5000	347	233	-344			-555	
15	3.7004	370	241	-365			-584	
13.88	4.0000	407	253	-395			-628	
							0_0	
12.33	4.5000	476	276	-440			-696	
12	4.6255	494	284	-450			-714	1
11.10	5.0000	548	306	-480			-760	
10	5.5506	633	342				-825	
9.5	5.8427	679	364					
0.051	6 0000	706						
9.251 9.0	6.0000 6.1674	706 732	377 389					
9.0 8.5	6.5301	792	420			*********		
8.0	6.9383	860	420 458			•••••		
7.929	7.0000	870	458					
1.929	1.0000	870	404					
7.5	7.4008	940	504					
7.0	7.9295	1,032	559					
6.938	8.0000	1,045	568					
6.5	8.5394	1,138	629					1
6.167	9.0000	1,216	686			,,		
					1			İ
6.0	9.2510	1,262	719					
5.551	10.0000	1,391	817					
5.5	10.0920	1,407	830					
5.0	11.1012	1,592	970				·	
4.5	12.3346	1,881	1,150	•••••				1
4.0	13.8765	2,251	1,379					
3.700	15.0000	2,485	1,543	••••••				
3.700 3.5	15.8589	2,463	1,665	•••••			**********	
3.25	17.0788	2,860	1 '			••••••		
3.0	18.5020	3,086				*********		
0.0	10.0020	0,000	L					

^{*}Estimated.

TABLE XV B. Heat of dilution: potassium compounds

n	m	KI	KIO ₃ *	KNO₃	KC ₂ H ₃ O ₂	KCNS	KMnO ₄
500,000 100,000 50,000 20,000	0.00 .000111 .000555 .00111 .00278	0 5 10 14 21	0 (5) (10) (12) (16)	0 5 10 14 19	0 5 12 16 25	0 5 11 14 22	0 4 7 9 13
7,000 5,000 4,000 3,000 2,000	.00555 .00793 .01110 .01388 .01850 .02775	29 33 37 41 45 51	(20) (19) (18) (16) (13) (+6)	23 24 25 25 25 24 19	35 42 49 55 63 76	29 33 38 40 45 49	13 12 9 6 +1 -13
1,500 1,110 1,000 900 800	.03700 .05000 .05551 .0617 .0694	53 57 57 58 59	(-5) (-21) (-29) (-38) (-47)	14 4 +1 -4 -11	87 93 107 112 119	52 55 56 56 56	-24 -42 -50 -58 -67
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	59 58 57 57 53	(-62) (-82) (-95) (-122)	-20 -31 -38 -48 -73	127 136 141 148 165	56 55 54 52 47	-81 -96 -105 -117 -149

^{*}Footnote at end of table.

TABLE XV B. Heat of dilution: potassium compounds-Continued

Φ_L, cal/mole, at 25 °C

	· .	Φ_L	, cal/mole, at 2	5 ℃			
n	m	KI	KIO₃*	KNO ₃	KC ₂ H ₃ O ₂	KCNS	KMnO ₄
300 277.5 200 150 111.0	.1850 .2000 .2775 .3700 .5000	44 41 +22 -2 -39		-114 -127 -196 -274 -376	190 197 232 265 306	35 31 +7 -24 -70	195 209 277 352
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	-55 -110 -184 -214 -293		-418 -547 -715 -779 -931	322 372 435 460 516	-89 -154 -243 -280 -368	
37.00 30 27.75 25 22.20	1.5000 1.8502 2.0000 2.2202 2.5000	-324 -417 -455 -509 -575		-987 -1,150 -1,215 -1,307 -1,414	537 594 616 647 694	-403 -507 -550 -611 -684	
20 18.50 15.86 15 13.88	2.7753 3.0000 3.5000 3.7004 4.0000	-639 -688 -793 -832 -888		-1,513 -1,586 -1,741 -1,799	732 753 836 870 908	-755 -810 -923 -965 -1,029	
12.33 12 11.10 10 9.5	4.5000 4.6255 5.0000 5.5506 5.8427	-977 -999 -1,063 -1,148 -1,191			977 990 1,046 1,120 1,164	-1,120 $-1,145$ $-1,212$ $-1,310$ $-1,354$	
9.251 9.0 8.5 8.0 7.929	6.0000 6.1674 6.5301 6.9383 7.0000	-1,214 -1,237 -1,287 -1,338 -1,346			1,184 1,212 1,256 1,320 1,332	$\begin{array}{c} -1,382 \\ -1,408 \\ -1,460 \\ -1,520 \\ -1,528 \end{array}$	
7.5 7.0 6.938 6.5 6.167	7.4008 7.9295 8.0000 8.5394 9.0000	-1,392 -1,450 -1,457 -1,510			1,398 1,470 1,480 1,562 1,632	$\begin{array}{c} -1,580 \\ -1,638 \\ -1,652 \\ -1,713 \\ -1,764 \end{array}$	
6.0 5.551 5.5 5.0 4.5	9.2510 10.0000 10.0920 11.1012 12.3346				1,670 1,782 1,798 1,950 2,160	-1,790 -1,862 -1,870 -1,960 -2,062	
4.0 3.700 3.5 3.25 3.0	13.8765 15.0000 15.8589 17.0788 18.5020					-2,170 -2,236 -2,296 -2,330 -2,390	
2.775 2.5 2.0	20.0000 22.2024 27.7530					-2,442 -2,510 -2,640	

^{*}Estimated.

Table XVI. Heat of dilution: rubidium and cesium compounds $\Phi_{L},\, cal/mole,\, at\,\, 25\,\, ^{\circ}C$

n	m	RbF	RbCl*	RbBr*	RbI*	RbNO₃*	CsF*	CsCl	CsBr*	CsI*	CsNO ₃
500,000 100,000 50,000 20,000 10,000	0.00 .000111 .000555 .00111 .00278 .00555	0 5 10 13 21 29	0 (5) (10) (13) (21) (28)	0 (5) (10) (13) (20) (26)	0 (5) (10) (13) (20) (26)	0 (5) (10) (13) (17) (20)	0 (5) (10) (13) (20) (27)	0 5 10 13 21 28	0 (5) (10) (13) (20) (26)	0 (5) (10) (13) (20) (26)	0 4 9 12 15
7,000 5,000 4,000 3,000 2,000	.00793 .01110 .01388 .01850 .02775	34 39 43 48 57	(34) (39) (42) (47) (53)	(32) (36) (39) (43) (49)	(31) (35) (38) (42) (47)	(22) (22) (24) (19) (14)	(31) (35) (39) (42) (50)	32 36 39 43 49	(30) (33) (36) (39) (45)	(29) (32) (35) (38) (43)	17 16 14 11 +2
1,500 1,110 1,000 900 800	.03700 .05000 .05551 .0617 .0694	63 71 74 77 80	(57) (61) (63) (64) (66)	(52) (55) (56) (56) (57)	(48) (49) (50) (50) (51)	(+7) (-4) (-10) (-15) (-22)	(54) (60) (62) (65) (70)	52 55 56 56 57	(46) (48) (49) (48) (48)	(43) (43) (43) (42) (42)	-8 - 23 - 29 - 35 - 44
700 600 555.1 500 400	.0793 .0925 .1000 .1110 .1388	84 89 91 95 102	(67) (68) (68) (69) (69)	(56) (56) (55) (54)	(49) (47) (45) (44)	(-32) (-43) (-52) (-62) (-92)	(75) (80) (82) (85) (80)	56 56 55 54 50	(45) (44) (42) (39)	(38) (35) (32) (29)	-56 -70 -79 -90 -123
300 277.5 200 150 111.0	.1850 .2000 .2775 .3700 .5000	111 113 122 130 137	(65) (63) (56) (44) (24)			(-137) (-151) (-220) (-303) (-413)		42 38 22 +1 -30			- 173 - 189 - 266 - 354 - 463
100 75 55.51 50 40	.5551 .7401 1.0000 1.1101 1.3877	140 	(15) 			(-455) (-587) (-760)		-45 -91 -152 -178			-508 -652 -831 *(-900) (-1067)

^{*}Estimated.

Table XVII. ΔH_N° at 25 °C

Investigator	System	n, moles H ₂ O/mole solute			Original temp of	No. of	$-\Delta H_N(T)$	$-\Delta H_N$	$-\Delta H_N^{\circ}$
		Acid	Base	Salt	meas	meas		(25 °C)	(25 °C)
Richards and Rowe [224]	NaCl	100	100	201	°C 16.80 15.67	5 4	$cal/mole \\ 14,059 \pm 15 \\ 14,114 \pm 8$	cal/mole 13,687 13,689	cal/mole 13,364 13,366
	KCl	100	100	201	16.77 15.63	3 4	$14,157 \pm 10$ $14,218 \pm 10$	13,763 13,768	13,372 13,377
	NaNO ₃	100	100	201	16.76 15.56	4 7	$13,988 \pm 6$ $14,050 \pm 6$	13,604 13,608	13,330 13,334
	KNO ₃	100	100	201	16.77 15.63	3 6	$14,220 \pm 5$ $14,293 \pm 6$	13,808 13,821	13,317 13,330
Gillespie, Lambert and Gibson [356]	NaCl KCl	540 283 550 284 282.5 260 273.4	540 283 550 284 282.5 260 273.4	1081 567 1101 569 566 521 548	20 20 24,99 32,27 32,27 32,30 20	1 1 1 1 1	13,773 13,793 13,571 13,199 13,261 13,271 13,858	13,516 13,545 13,571 13,525 13,588 13,596 13,625	13,349 13,333 13,407 13,315 13,376 13,377 13,384
Bender and Biermann [213]	NaCl	18.42 17.14	18.42 17.14	37.84 35.30	25 25	2 4	$14,249 \pm 12$ $14,314 \pm 10$	14,249 14,314	13,332 13,331
Pitzer [206]	NaCl	46.5	1065.5	1112	25	3	13,828 ± 12	13,828	13,366
Vanderzee and Swanson [357]	NaClO ₄	5050 188.6	178.4 4060.4	5261 5160	25 25	2 6	13,465 ± 15 13,398 ± 5	13,465 13,398	13,332 13,337

Table XVIII. $\Delta H_{\infty}^{\circ}$ of KCl(c) in H₂O at 25 °C

		T	T =		
Investigator	Year reported	Temp of meas	No. of meas	Conc. range moles KCl/ 1000g H ₂ O	$\Delta H_{\omega}^{\circ}$
		°C			cal/mole
1. Winkelmann [94]	1873	26-31	17	5.59 to 0.42	4,104 ±50
2. Berthelot [377]	1875	21	?	0.28	$3,970 \pm 150$
3. von Rechenberg [381]	1879	16-20	. 7	.28	4,113 ±100
[382]	1880	17-20	7	.28	$4,200 \pm 100$
4. Berthelot and Ilosvay [383] 5. Thomsen [30]	1883 1883	18 18	?	.54 .28	4,000 ±150 4,107 ±50
6. Pickering [384]	1887	15-25	30	.28	4,143 ±50
7. von Stackelberg [275]	1898	18	3	4.44 to 0.55	4,110 ±100
8. Brönsted [385]	1906	25	4	0.46 to 0.09	$4,167 \pm 20$
9. Haigh [388]	1912	21	2	.14	$4,081 \pm 50$
10. Colson [350]	1915	18, 21	2	17	4,184
11. Cohen, Helderman, and Moesveld [386] 12. Mees [387]	1920 1920	18	$\frac{7}{2}$.28	4,070 ±30 4,062
13. Mondain-Monval [268]	1923	18	1 1	.37	3,954
14. Sandonnini and Gerosa [141]	1925	18.5	4	.75 to 0.17	4,100 ±100
15. Wüst and Lange [21]	1925	25	12	4.47 to 0.37	$4,157 \pm 40$
16. Lange and Dürr [389]	1926	25	10	0.56	$4,147 \pm 40$
17. Holluta and Werner [390]	1927	18.5, 20	2	.27	4,183 ±50
18. Cohen and Kooy [391]	1928	20 25	5 3	.28 .28	4,110 ±10 4,110 ±5
19. Chipman, Johnson, and Maass [133]	1929	15-27	16	.66 to 0.07	4,110 ±5 4,100 ±100
20. Partington and Soper [334]	1929	25	12	3.55 to 0.31	$4,160 \pm 20$
21. Roth and Eymann [392]	1929	21	8	0.37	$4,112 \pm 15$
22. Lange and Rounsefell [393]	1930	25	6	.28	$4,116 \pm 10$
23. Mishchenko [303]	1930	25	6	.25	$4,114 \pm 10$
24. Popov, Bundel, and Choller [165] 25. Moles and Perez-Vitoria [394]	1930 1932	20 20.5	10 1	.19 .34	$4,082 \pm 25$
26. Smeets [395]	1932	20.3	3	.34	$4,298$ $4,107 \pm 20$
27. Popov, Khomyakov, Feodos'ev, and Schirokich [166]	1933	20	10	.19	$4,077 \pm 20$
		21	8	.28	$4,115 \pm 10$
00 D 1 [47]	1005	23	19	.28	$4,114 \pm 15$
28. Drucker [45] 29. Shibata and Terasaki [396]	1935 1936	17-23 25	12 10	.56 to 0.11 .15 to 0.07	$4,095 \pm 35$
30. Fedorov and Sil'chenko [271]	1937	20	3	.15 to 0.07	$4,164 \pm 20$ $4,150 \pm 100$
31. Lange and Martin [397]	1937	25	lĭ	.13	4,130 ± 100 4,147
32. Fontell [398]	1938	22.5	4	.06 to 0.01	$4,124 \pm 10$
		25	6		$4,117 \pm 15$
33. Popov, Skuratov, and Strel'tsova [172]	1940	20	?	.28	$4,125 \pm 20$
34. Slansky [399] 35. Voskresenskaya and Ponomareva [400]	1940 1946	25 25	?	? 0.18	$^{a}4,075 \pm 40$
36. Tichelaar [401]	1946	25	6	.02?	$4,124 \pm 20$ $4,146 \pm 40$
37. Foz and Colomina [402]	1946	25	ĭ	.37	⁵ 4.115 ±10
[403]		1-0			$4,098 \pm 20$
38. Fineman and Wallace [404]	1948	25	?	?	^a 4,113
39. Mishchenko and Kaganovich [405]	1949	25	5 ? 6 3	0.28	4.115 ± 10
40. Barieau and Giauque [406] 41. Samoĭlov [407, 408]	1950 1951	25 25	3	.18	4,113 ±10
42. Kaganovich and Mishchenko [409]	1952	25 25	1 ?	.06 .26	4,169 $4,117 \pm 10$
43. Evans and Richards [410]	1952	25	4	.01	4,117 ±10 4,121 ±10
44. Kapustinskii and Drakin [411]	1952	25	4 ?	.001	4,077 ±25
45. Spedding and Miller [412]	1952	25	6	.19 to 0.03	$4,132 \pm 15$
46. Voskresenskaya and Patsukova [413]	1952	25	?	.12	$4,114 \pm 15$
47. Westrum and Eyring [414]	1952	25	? ? 5	.02	4,109 ±15
48. Davies, Singer, and Stavely [368] 49. Hutchinson, Manchester, and Winslow [415]	1954 1954	25 25	8	.32 .28, 0.03	$4,108 \pm 10$ $4,120 \pm 8$
50. Chernyaev, Sokolov, and Palkin [416]	1954	25	3	.02	4,120 ±8 4,129 ±15
51. Hutchinson and White [417]	1955	25	4	.01 to 0.007	4,120 ±25
52. Higgins [418]	1955	25	3	.11	4,133 ±10

TABLE XVIII. ∆H_∞ of KCl(c) in H₂O at 25 °C - Continued

Investigator	Year reported	Temp of meas	No. of meas	Conc. range moles KCl/ 1000g H ₂ O	$\Delta H_{\infty}^{\circ}$
53. Papadopoulos and Giauque [419] 54. Coops, Balk, and Tolk [420] 55. Gunn [421] 56. Balk and Benson [422] 57. Sacconi, Paoletti, and Ciampolini [205, 423] 58. Sunner and Wadsö [424] 59. Bills and Cotton [425] 60. Hietala [426] 61. Gallagher and King [427] 62. Dekker [428] 63. Reshetnikov [429] 64. Ciampolini and Paoletti [430] 65. Higgins and Westrum [431] 66. Argue, Mercer, and Cobble [432] 67. Coops, Somsen, and Tolk [433] 68. Busey, Dearman, and Bevan [434] 69. Newman [435] 70. Talakin et al., [436]	1955 1956 1958 1959 1959 1959 1960 1960 1961 1961 1961 1961 1961 1951 1957 1958 1957 1957 1962	°C 25 25 25 25 25 25 25 25 25 25 25 25 25	3 16 8 9 10 11 4 3 14 ? 6 8 3 4 10 10 4 5 6 ? 12 3	0.28 .01 .28 .20 .33 .12 .28 .28 .14-0.7 .28 .19 to 0.11 .33 .11 .005 .02 to 0.007	$\begin{array}{c} cal/mole\\ 4,107\pm6\\ 4,082\pm20\\ 4,122\pm2\\ 4,125\pm5\\ 4,119\pm10\\ 4,100\pm5\\ 4,121\pm10\\ 4,116\pm10\\ 4,120\pm10\\ 4,112\pm10\\ 4,112\pm10\\ 4,112\pm10\\ 4,113\pm10\\ 4,117\pm10\\ 4,109\pm5\\ 4,106\pm5\\ 4,107\pm10\\ 4,117\pm10\\ 4,105\pm10\\ 4,117\pm10\\ 4,105\pm10\\ 4,107\pm10\\ 4,107\pm10\\ 4,107\pm10\\ 4,107\pm10\\ 4,108\pm21\\ 4,088\pm25\\ 4,088$

Table XIX. $\Delta H_{\infty}^{\circ}$ of NaCl in H_2O at 25 $^{\circ}C$

Investigator	Year reported	Temp of meas	No. of meas	Conc. range moles NaCl/ 1000g H ₂ O	$\Delta H_{\infty}^{\circ}$
1. Winkelmann [94] 2. Ostwald [443] 3. Thomsen [30] 4. Pickering [384] 5. Colson [535] 6. Randall and Bisson [143] 7. Wüst and Lange [21] 8. Lange and Dürr [389] 9. Lipsett, Johnson, and Maass [323] [536] 10. Cohen and Kooy [391] 11. Askew, Bullock, Smith, Tinkler, Gatty, and Wolfenden [499] 12. Becker and Roth [228] 13. Lange and Martin [397] 14. Slansky [399] 15. Tichelaar [401] 16. Voskresenskaya and Ponomareva [400] 17. Fineman and Wallace [404] 18. Samoilov [407, 408] 19. Kapustinskii and Drakin [411] 20. Seki and Suzuki [537] 21. Benson and Benson [324] 22. Benson, Goddard, and Hoeve [538] 23. Attree, Cushing, Ladd, and Pieroni [539] 24. Lister and Myers [540] 25. Samoilov and Buslaeva [444] 26. Criss and Cobble [541]	1873 1882 1883 1887 1915 1920 1925 1926 1927 1927 1928 1934 1935 1937 1940 1946 1948 1951 1952 1953 1955 1956 1958 1958 1958	°C 16-30 16-17 16.8 15-25 21 25 25 20 20 20 29 27 25 25 25 25 25 25 25 25 25 25 25 25 25	20 4 2 35 1 7 14 10 20 15 4 5 9 5 1 ?? 16 3 ?? 21	5.50 to 0.53 0.19 .56 .28 .42 .54 to 0.13 5.62 to 0.52 0.56 5.50 to 0.07 5.92 to 0.04 0.56 .56 .03 .09 to 0.04 .25 ? 0.02? .37 ? 0.19 .0008 .14 1.3 to 0.05 0.50 .01 .18 .14 .02 to 0.001	cal/mole 914 ±30 970 897 ±20 935 ±20 894 929 ±5 931 ±10 841 923 ±5 922 ±8 915 ±5 921 ±8 904 ±30 918 ±10 934 4 915 ±10 950 ±30 970 2 925 ±5 932 932 ±3 972 ±3 927 ±10 914 ±10 874 ±10 914 ±10 914 ±10

^a Authors' extrapolation.

 ^a Authors' extrapolation.
 ^bFoz and Colomina report this as their best value.
 ^cMeasurements made by Tolk in apparatus of Coops, Balk and Tolk.

TABLE XX. ΔH_∞ of KNO₃ in H₂O at 25 °C

Investigator	Year reported	Temp of meas	No. of meas	Conc. range moles KNO ₃ / 1000g H ₂ O	$\Delta H_{_{\infty}}^{\circ}$
1. Winkelmann [94] 2. Thomsen [30] 3. Tilden [560] 4. Pickering [384] 5. von Stackelberg [275] 6. Haigh [388] 7. Mondain-Monval [267] 8. Berenger-Calvet [351] 9. Holluta and Werner [390] 10. Roth [576] 11. Roth and Müller [577] 12. Roth and Eymann [392]; Roth [578] 13. Cohen and Kooy [391] 14. Nacken [477] 15. Lange and Monheim [173] 16. Drucker [45] 17. Fedorov and Sil'chenko [271] 18. Hieber and Feder [579] 19. Hückel, Datow, and Simmersbach [580] 20. Voskresenskaya and Ponomareva [581] 21. Voskresenskaya and Ponomareva [400] 22. Westrum and Eyring [414] 23. Lipilina and Samoilov [582] 24. Bills and Cotton [425] 25. Turnbull [583]	1873 1883 1884 1887 1898 1912 1925 1927 1927 1927 1929 1928 1930 1930 1934 1937 1938 1940 1944 1946 1952 1954 1954 1960	°C 27-29 18 15, 34 15-25 15 21 16.9 14-16 18.5 20.5 20, 20.5 22; 25 18.1 22 0 18.5 25 25 25 25 25	14 3 4 27 10 3 1 30 1 6 3 4 6 2 5 1 3 1 3 1 4 6 2 1 1 3	2.41 to 0.31 0.28 .56 .28 2.44 to 0.22 0.14 .20 2.49 to 0.09 0.16 .16 .16 .16 .27 .07 to 0.05 .23 .56 to 0.14 .18 .18 .19 .11	$\begin{array}{c} cal/mole\\ 8,350\pm50\\ 8,300\pm50\\ 8,295\pm100\\ 8,430\pm75\\ 8,361\pm50\\ 8,240\pm75\\ 8,260\pm75\\ 8,200\\ 8,355\pm40\\ 8,592\\ 8,455\\ 8,304\\ 8,372\pm20\\ 8,585\\ 8,375\pm40\\ 8,285\\ 8,380\pm75\\ 8,340\\ 8,285\\ 8,380\pm75\\ 8,340\\ 8,407\pm40\\ 8,290\\ 8,290\\ 8,290\\ 8,400\pm40\\ 8,600\pm100\\ \end{array}$

Table XXI. Heat of Solution $\Delta H_{\infty}^o, \ 25 \ ^o\!C \ for \ uni-univalent \ electrolytes \ in \ H_2O$

Substance	State	$\Delta H_{\infty}^{\circ}$	Substance	State	$\Delta H_{\infty}^{\circ}$	Substance	State	$\Delta H_{\infty}^{\circ}$
		cal/mole			cal/mole			cal/mole
HF	g	-14,700	LiBr·2H ₂ O	c	-2,250	KCI	l c	4.11
HCl	g	-17,888	LiBrO ₃	c	340	KClO ₃	č	9,89
HClO ₄	Ĭ	-21,215	LiI	c	-15,130	KClO ₄	č	12,20
HClO ₄ ·H ₂ O	c	-7,875	LiI·H ₂ O	c	-7,090	KBr	c	4,75
HBr	g	-20,350	LiI·2H ₂ O	c	-3,530	KBrO ₃	c	9,83
HI	g	-19,520	LiI·3H ₂ O	c	140	KI	c	4.86
HIO_3	c	2,100	LiNO ₂	c	-2,630	KIO ₃	c	6,63
HNO_3	1	-7,954	LiNO ₂ ·H ₂ O	c	1,680	KNO ₂	c	3,19
НСООН	1 :	-205	LiNO ₃	c	-600	KNO ₃	c	8,34
CH₃COOH	1	-360			ļ	KC ₂ H ₃ O ₂	c	-3,66
			NaOH	c	-10,637	KCN	c	2,80
NH_3	g	-7,290	NaOH·H ₂ O	c	-5,118	KCNO	c	4,84
NH ₄ Cl	C	3,533	NaF	c	218	KCNS	c	5,79
NH ₄ ClO ₄	c	8,000	NaCl	c	928	KMnO ₄	c	10,41
NH ₄ Br	c	4,010	NaClO ₂	c	80			
NH ₄ I	c	3,280	NaClO ₂ ·3H ₂ O	c	6,830	RbOH	С	-14,90
NH ₄ IO ₃	c	7,600	NaClO ₃	c	5,191	RbOH⋅H₂O	c	-4,31
NII ₄ NO ₂	c	4,600	NaClO ₄	c	3,317	RbOH∙2H ₂ O	С	21
NH ₄ NO ₃	c	6,140	NaClO ₄ ·H ₂ O	c	5,380	RbF	c	-6,24
NH ₄ C ₂ H ₃ O ₂	c	-570	NaBr	c	-144	RbF·H₂O	c	-10
NH₄CN	c	4,200		c	4,454	RbF·1½H₂O	c	32
NH ₄ CNS	c	5,400	NaBrO ₃	c	6,430	RbCl	c	4,13
CH ₃ NH ₃ Cl	c	1,378		c	-1,800	RbClO ₃	c	11,41
(CH ₃) ₃ NHCl	c	350		С	3,855	RbClO ₄	c	13,56
N(CH ₃) ₄ Cl	c	975		С	4,850	RbBr	c	5,23
N(CH ₃) ₄ Br	c	5,800		c	3,320	$RbBrO_3$	c	11,70
N(CH ₃) ₄ I	c	10,055		c	4,900	RbI	c	6,00
	l		NaC ₂ H ₃ O ₂	С	-4,140	$RbNO_3$	c	8,72
AgClO ₄	c	1,760		c	4,700		1	ł
$AgNO_2$	C.	8,830	NaCN	c	290	CsOH	c	-17,10
$AgNO_3$	c	5,400		c	790	$CsOH \cdot H_2O$	c	-4,90
			NaCN·2H₂O	c	4,440	CsF	c	-8,81
LiOH	c	-5,632	NaCNO	c	4,590	CsF·H ₂ O	С	-2,50
LiOH·H₂O	С	-1,600	NaCNS	C	1,632	$CsF \cdot l \frac{1}{2}H_2O$	c	-1,30
LiF	c	1,130		-		CsCl	С	4,25
LiCl	c	-8,850		c	-13,769	CsClO ₄	c	13,250
LiCl·H ₂ O	c	-4,560	KOH·H ₂ O	c	-3,500	CsBr	c	6,210
LiClO ₄	c	-6,345		c	-2,500	$CsBrO_3$	c	12,06
LiClO₄·3H₂O	c	7,795	KF	С	-4,238	CsI	c	7,97
LiBr	c	-11,670	KF∙2H ₂ O	c	1,666	$CsNO_3$	c	9,56
LiBr∙H₂O	c	-5,560			'			1