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THE ACID DISSOCIATION OF THE HYDRATED CUPRIC ION

THE FORMATION OF DIMERIC CUPRIC COMPOUNDS

BY

KAI JULIUS PEDERSEN



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Printed in Denmark. Bianco Lunos Bogtrykkeri A/S. O ur knowledge of the hydrogen ion concentration of solutions of cupric salts, and the equilibria which determine it, is very incomplete. The hydrogen ion concentration of a cupric sulphate solution was first measured by CARRARA and VESPIG-NANI⁽¹⁾ using the method of inversion of sucrose. KULLGREN⁽²⁾ applied the same method to solutions of cupric chloride and nitrate. The quinhydrone electrode was used by O'SULLIVAN⁽³⁾ and by FRIEDMAN and STOKES⁽⁴⁾ in cupric sulphate solutions, by QUINTIN⁽⁵⁾ in cupric nitrate and sulphate solutions, and by ČUPR and ŠIRŮČEK⁽⁶⁾ in different cupric salt solutions. Finally, the glass electrode was employed for measuring the pH of cupric sulphate solutions by HUGHES⁽⁷⁾ and by CRANSTON and BROWN⁽⁸⁾. A selection of the best measurements is given in Fig. 1.

It is usually assumed that the hydrogen ion concentration is determined by the equilibrium

$$Cu^{++} + H_2O \rightleftharpoons CuOH^+ + H^+, \tag{1}$$

but the measurements are not sufficiently accurate to test this assumption. If anything, they rather disfavour it. Thus, QUINTIN as well as ČUPR and ŠIRŮČEK state that, when pH is plotted against the logarithm of the concentration of the copper salt, the points fall close to a straight line with the slope one (cf. Fig. 1), or, in other words, the hydrogen ion concentration is roughly proportional to the cupric ion concentration. If this holds, equilibrium 1 alone does not determine the hydrogen ion concentration. The relation found by these investigators rather indicates the equilibrium

$$2 \operatorname{Cu}^{++} + \operatorname{H}_2 O \rightleftharpoons \operatorname{Cu}_2 O \mathrm{H}^{+++} + \mathrm{H}^+.$$

1*

JANDER and MÖHR⁽⁹⁾ recently studied the diffusion of solutions of cupric perchlorate and cupric chloride in the presence of neutral salt or strong acid. For all their solutions, they found the diffusion coefficient to be the same within a few per cent, while pH varied from 5 to 0. However, their pH values were often several units lower than should be expected from measure-

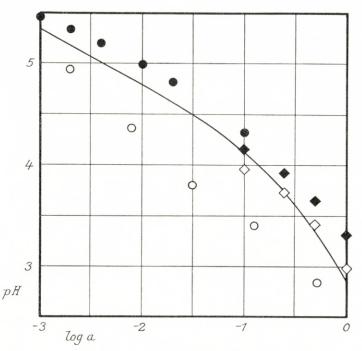


Fig. 1. Measurements of the pH of solutions of cupric nitrate and sulphate (*a* molar) by the investigators: QUINTIN (quinhydrone electrode, cupric nitrate, open circles), CRANSTON and BROWN (glass electrode at 15° C., cupric sulphate, solid circles), and ČUPR and ŠIRŮČEK (quinhydrone electrode at 20° C., cupric nitrate, open squares, and cupric sulphate, solid squares). The curve represents the measurements of the present author.

ments by other investigators, including the author of the present paper. It appears from their results that the degree of hydrolysis varies from 0 to a very considerable value (> $20 \, {}^0/_0$), while the diffusion coefficient is practically constant. Since a dimeric ion would have a smaller diffusion coefficient than the usual cupric ion, they conclude that only monomeric hydrolysis products are present. However, it seems safe to maintain that the degree of hydrolysis of their solutions is always less than $1 \, {}^0/_0$.

Therefore, their measurements cannot give any information concerning the nature of the hydrolysis products.

The object of the present paper is to throw light on the problem of acid-base equilibria in aqueous solutions of cupric ions. For this purpose, the hydrogen ion concentrations of solutions of cupric nitrate, varying in concentration from 1 to 0.001 molar, were measured by means of the glass electrode at 18.0° C. Cupric nitrate was chosen rather than sulphate, because the presence of sulphate ions would make the problem more complicated owing to their power of forming complexes with cupric ions and of combining with hydrogen ions. Furthermore, the greater electric charge of the sulphate ion would cause a greater salt effect. Measurements were carried out both on pure aqueous solutions of cupric nitrate and on solutions containing, in addition, various amounts of nitric acid or sodium hydroxide. It is well known that only very little sodium hydroxide can be added before a precipitate is formed. According to J. BJER-RUM⁽¹⁰⁾, the green crystalline substance which precipitates in weakly alkaline solutions of cupric nitrate has the composition Cu₂(OH)₂NO₂. It was often possible to carry out measurements on solutions containing so much sodium hydroxide that they were supersaturated with respect to the substance forming the precipitate. For such solutions, the e.m. f. remained constant for a period of time, but when they were kept standing sufficiently long, a gradual increase in the hydrogen ion concentration was measured. At the same time, the solutions turned turbid. The greatest concentration of sodium hydroxide which could be maintained without disturbing the measurements was: in 1 molar $Cu(NO_3)_2$ only $0.2^{0/0}$ of the concentration of cupric nitrate, in 0.1 molar $Cu(NO_3)_2$ it was 1.2 $^0/_0$, and in 0.005 molar $Cu(NO_3)_2$ it was more than $10^{0}/_{0}$. These examples show that the solutions measured were very weakly buffered. There is no sense in carrying out measurements on such systems if the amount of acid or basic impurity present is not known with great accuracy. We shall see later how these impurities may be estimated.

Cupric nitrate hexahydrate was used for the experiments. It was prepared by recrystallization of the best trihydrate available (Schering-Kahlbaum's *Cuprum nitr. puriss.*). 1000 g. of the

trihydrate were dissolved in 400 ml. of warm water. The solution was cooled down to 0°, seeded with the hexahydrate, and was left for crystallization at 0°. The crystals were separated from the mother liquor and were dried in a desiccator over anhydrous calcium chloride. When the crystals began to effloresce the drying agent was removed. The preparation contained an excess of base, or a deficiency of acid, which increased gradually during storage. At the beginning of the measurements it amounted to 3×10^{-5} , at the end to 18×10^{-5} equivalents per mole of cupric nitrate. This increase is probably due to the fact that the glass basin containing the preparation was inadvertently placed in the desiccator on a perforated iron plate. The nitric acid lost may have distilled from the slightly moist crystals to the iron plate.

Standard solutions of about 1 or 2 molar cupric nitrate were made from the preparation at different times. The solutions, which were kept in a silica flask, were analyzed for copper according to the modified iodometric titration method of HAGEN⁽¹¹⁾, which gave excellent results (accuracy about $0.1 \, {}^{0}/_{0}$). Metallic copper (electrolytically deposited, *pro analysi*) was used as a standard substance.

The barium nitrate used in some of the measurements was a Merck *pro analysi* preparation which was recrystallized and dried at 110° C. It contained no appreciable basic or acid impurities.

The water used for the solutions was redistilled in a still supplying water of a specific conductance of less than 3×10^{-7} ohmcm⁻¹. It was stored in bottles covered inside with a layer of paraffin wax.

The cells measured had the composition

where X is the solution under investigation. The glass electrode was of the plane type described by MACINNES and DOLE⁽¹²⁾. It was tested before use, and was found to follow practically completely the law holding for the hydrogen electrode through-

out the pH interval in which it was going to be used. As container for the solution under investigation served the vessel shown in Fig. 2. Its total volume was 100 ml., but only 70 ml. of solution were used for the measurement. For the solutions containing more than 0.1 molar cupric nitrate, a similar, but smaller vessel requiring only 10 ml. was applied. The opening D was closed with a rubber stopper carrying the glass electrode which dipped into the solution. The side tube E was completely

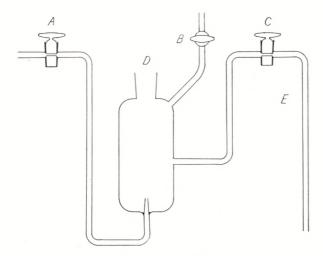


Fig. 2. Electrode vessel.

filled with the solution. Before the measurement, possible traces of carbon dioxide were removed by a vigorous current of pure, carbon dioxide free air which was passed through the solution for 10 minutes. The air entered through the stopcock A and escaped through B.

The ready cell was placed into a liquid paraffin thermostat at 18.0° C. The lower end of E dipped into a test tube containing 8 molar ammonium nitrate which provided connection with the quinhydrone electrode. The levels of the solutions were so arranged that the bridge solution rose a few centimetres into E when the stopcock C was opened. The e.m. f. was measured by means of a valve potentiometer (made by the firm "Radiometer", Copenhagen). The potential had become constant after a few minutes, and generally it did not change for hours. Only when the solution was supersaturated, a gradual decrease was

observed, as already mentioned, after a longer or a shorter period of constant potential.

At least once every day, a solution of the composition 0.01000 m HCl + 0.0900 m NaCl was measured instead of X. The small potential measured here is the asymmetry potential of the cell. When it is subtracted from the e.m. f. measured for X, the potential *E* of a cell of the composition

 $H_2 \mid X \mid 8 \text{ m } NH_4 NO_3 \mid 0.01000 \text{ m } HCl + 0.0900 \text{ m } NaCl \mid H_2$ (2)

is obtained.

To a series of measurements at constant cupric nitrate concentration but varying nitric acid or sodium hydroxide concentration, we may apply the equation

$$E = E_a - \frac{RT}{F} \ln \left(\mathrm{H}^+ \right),$$

where (H⁺) is the hydrogen ion *concentration*, and E_a includes the constant potential of the right half cell, the liquid-liquid potential, and the salt effect of the left half cell. When a sufficiently small amount of acid or base is added, E_a is constant. For the temperature 18° C. the equation may be written

$$-\log(\mathrm{H}^+) = A + 17.32 \, E,\tag{3}$$

where $A \equiv -17.32 E_a$. The value of A corresponding to an infinitely small concentration of acid or base added will be denoted by A_0 .

A was determined in the following way. E was measured for solutions of cupric nitrate containing so much nitric acid that the hydrogen ion concentration produced by the cupric ions was either negligible or could be estimated from preliminary values of the equilibrium constants. In no case, this correction exceeded 0.5 % of the concentration of nitric acid added. Also the small acid deficiency of the cupric nitrate, estimated from later measurements, was taken into account. (H⁺) and E were introduced into equation 3, and 4 was computed. A_0 , the value of A corresponding to the concentration 0 of nitric acid, was found for each series by extrapolation. The results are given in Table 1. The table includes two series of measurements in

Table 1.

Electromotive force E volts at 18.0° C. of the cell

H_2	a molar $Cu(NO_3)_2$ b molar $Ba(NO_3)_2$	8 molar NH ₄ NO ₃	0.01000 molar HCl 0.0900 molar NaCl	H_2
	y molar HNO ₃	1114103	0.0500 molar Naci	

measured by means of the glass electrode. $y = -x - a \delta$, where -x is the concentration of nitric acid added, and δ is the deficiency of nitric acid in the preparation of cupric nitrate expressed in equivalents per mole. A has been computed from equation 3. A_0 is the value of A corresponding to y = 0.

a	b	$\delta10^5$	<i>y</i>	E	A	A_0
1.000	0	18	$\left\{\begin{array}{c} 0.04975\\ 0.02468\\ 0.01002\end{array}\right.$	-0.0578 -0.0403 -0.0180	$\begin{array}{c} 2.304 \\ 2.305 \\ 2.309 \end{array} \right\}$	2.308
0.500	0	16	$\left\{ \begin{array}{l} 0.04098 \\ 0.02008 \\ 0.00801 \end{array} \right.$	$-0.0424 \\ -0.0247 \\ -0.0016$	$\left. \begin{array}{c} 2.122 \\ 2.125 \\ 2.123 \end{array} \right\}$	2.123
0 200	0	8	$\left\{\begin{array}{l} 0.01027\\ 0.004104\\ 0.002059\end{array}\right.$	$-0.0023 \\ 0.0205 \\ 0.0377$	$\left. \begin{array}{c} 2.028 \\ 2.031 \\ 2.032 \end{array} \right\}$	2.033
0.100	0	3	$\left\{ \begin{array}{l} 0 \ 0 \ 1000 \\ 0.005015 \\ 0.002525 \end{array} \right.$	$-0.0005 \\ 0.0166 \\ 0.0335$	$\left. \begin{array}{c} 2.009 \\ 2.012 \\ 2.017 \end{array} \right\}$	2.018
0	0.100		$\left\{\begin{array}{l} 0.01000\\ 0.005017\\ 0.002528\\ 0.001025\end{array}\right.$	-0.0005 0.0166 0.0335 0.0561	$\begin{array}{c} 2.009 \\ 2.012 \\ 2.017 \\ 2.017 \end{array}$	2.018
0.0500	0	8	$\left\{\begin{array}{l} 0.005134\\ 0.002055\\ 0.001033\end{array}\right.$	$0.0160 \\ 0.0387 \\ 0.0558$	$\left. \begin{array}{c} 2.012 \\ 2.017 \\ 2.019 \end{array} \right\}$	2.021
0.0200	0	3	$\left\{\begin{array}{l} 0.005138\\ 0.002059\\ 0.001033\end{array}\right.$	$0.0154 \\ 0.0378 \\ 0.0558$	$\left. \begin{array}{c} 2.022 \\ 2.032 \\ 2.033 \end{array} \right\}$	2.040
0.0100	0	3	$\left\{\begin{array}{c} 0.00504\\ 0.002021\\ 0.001018\end{array}\right.$	$0.0153 \\ 0.0373 \\ 0.0542$	$\left. \begin{array}{c} 2.033 \\ 2.048 \\ 2.054 \end{array} \right\}$	2.058
0.0020	0.0080	3	$\left\{\begin{array}{l} 0.002021\\ 0.001018\\ 0.000413\end{array}\right.$	$0.0374 \\ 0.0544 \\ 0.0767$	$\left. \begin{array}{c} 2.046 \\ 2.050 \\ 2.056 \end{array} \right\}$	2.058

а	A_0
0.0050	
0.0050 0	2.072
	2.086
0.0000	
0.0020	
	2.097
0.0010	
0.0010	2

Table 1 (continued)

solutions containing barium nitrate. It is seen that, both in 0.1 and 0.01 molar cupric nitrate, the cupric ions may be replaced by barium ions without altering the value of A_0 .

In the following, the hydrogen ion concentration of solutions of cupric nitrate containing small concentrations of sodium hydroxide or nitric acid will be computed from the measurements by means of the formula

$$-\log(\mathrm{H}^+) = A_0 + 17.32E, \qquad (4)$$

where A_0 has the values given in Table 1. Before analyzing the results given in Table 2, we must consider the theory to be applied.

We may explain the hydrogen ion concentrations of the solutions under the assumption that the following three equilibria take place

$$Cu^{++} \rightleftarrows CuOH^+ + H^+, \tag{5}$$

$$2\operatorname{Cu}^{++} \rightleftharpoons \operatorname{Cu}_2\operatorname{OH}^{+++} + \operatorname{H}^+, \tag{6}$$

and

$$2\operatorname{Cu}^{++} \rightleftharpoons \operatorname{Cu}_2 \operatorname{O}^{++} + 2\operatorname{H}^+. \tag{7}$$

In these schemes the water of hydration has been omitted. The mass action constants are

$$k_1 = \frac{(\text{CuOH}^+) (\text{H}^+)}{(\text{Cu}^{++})}, \qquad (8)$$

$$k_{2} = \frac{(\mathrm{Cu}_{2}\mathrm{OH}^{+++})(\mathrm{H}^{+})}{(\mathrm{Cu}^{++})^{2}},$$
(9)

and

$$k_{3} = \frac{(\mathrm{Cu}_{2}\mathrm{O}^{++}) (\mathrm{H}^{+})^{2}}{(\mathrm{Cu}^{++})^{2}}.$$
 (10)

We consider a series of measurements in a molar cupric nitrate. The cupric nitrate has an acid deficiency of δ equivalents per mole. x molar sodium hydroxide, or -x molar nitric acid, has been added. It follows from the theory that

$$\begin{split} (\mathrm{H}^+) + x + a\,\delta &= (\mathrm{CuOH}^+) + (\mathrm{Cu}_2\mathrm{OH}^{+++}) + 2\,(\mathrm{Cu}_2\mathrm{O}^{++}) \\ &= k_1 \frac{(\mathrm{Cu}^{++})}{(\mathrm{H}^+)} + k_2 \frac{(\mathrm{Cu}^{++})^2}{(\mathrm{H}^+)} + 2\,k_3 \frac{(\mathrm{Cu}^{++})^2}{(\mathrm{H}^+)^2}, \end{split}$$

which may also be written as follows

$$\frac{\left[(\mathrm{H}^+) + x\right](\mathrm{H}^+)}{(\mathrm{Cu}^{++})} + \frac{a\,\delta(\mathrm{H}^+)}{(\mathrm{Cu}^{++})} = k_1 + k_2\,(\mathrm{Cu}^{++}) + 2\,k_3\frac{(\mathrm{Cu}^{++})}{(\mathrm{H}^+)}\,.$$
 (11)

If the decrease in cupric ion concentration owing to the reactions 5 to 7 is denoted by \varDelta , we get

$$(Cu^{++}) = a - \varDelta \tag{12}$$

and

$$\begin{split} \mathcal{A} &= (\mathrm{CuOH^+}) + 2\,(\mathrm{Cu_2OH^{+++}}) + 2\,(\mathrm{Cu_2O^{++}}) \\ &= (\mathrm{H^+}) + x + a\,\delta + (\mathrm{Cu_2OH^{+++}}) \\ &= (\mathrm{H^+}) + x + a\,\delta + k_2\,\frac{(\mathrm{Cu^{++}})^2}{(\mathrm{H^+})}\,. \end{split}$$

An approximation leads to

$$\Delta = (\mathrm{H}^+) + x + k_2 \frac{a^2}{(\mathrm{H}^+)}.$$
 (13)

From the equations 11 and 12 we obtain

$$k = (k_1 + ak_2) + 2k_3 \frac{(\operatorname{Cu}^{++})}{(\operatorname{H}^{+})}, \qquad (14)$$

where

$$k \equiv \frac{\left[(\mathrm{H}^+) + x \right] (\mathrm{H}^+)}{(\mathrm{Cu}^{++})} + \frac{a \,\delta(\mathrm{H}^+)}{(\mathrm{Cu}^{++})} + k_2 \mathcal{A}. \tag{15}$$

In order to test the theory, we first compute \varDelta from equation 13 and a preliminary value of k_2 , and then (Cu⁺⁺) from equation 12. k of equation 15 is a sum of three terms, the first and most important of which may now be calculated. The third term is a very small correction and may be estimated from a preliminary value of k_2 . The relative importance of the second term, which contains the unknown δ , decreases when x increases within a series of measurements at constant a. If we as a first approximation put $\delta = 0$, preliminary values of k may be calculated and plotted against $\frac{(Cu^{++})}{(H^{+})}$ (the solid circles in Fig. 3). According to equation 14, the true value of k leads to a straight line when a is constant; actually, the points computed under the assumption that $\delta = 0$ fall on a curve which approaches a straight line with increasing x. Generally, the approach is the more rapid the smaller a. By the method of trial and error, we find the value of δ for which a rectilinear plot is obtained (open circles in Fig. 3). From the straight line, the numerical values of the two constants $(k_1 + ak_2)$ and $2k_3$ in equation 14 may be determined. The series chosen as an example in Fig. 3 is one in which the deviation from the straight line of the points computed for $\delta = 0$ is fairly great. Only for two of the 15 series, namely when a = 1 and a = 0.5, the deviations were greater.

The results of the measurements and computations are presented in Tables 2 and 3. The numbers of the series, given in the first column of Table 2, indicate the chronological order. It is seen that δ , which is given in the same column, increases with time. When the same standard solution of cupric nitrate was used for preparing the solutions in several series, the same δ was found. x, the concentration of sodium hydroxide added, is given in the second column. Nitric acid is reckoned as sodium hydroxide with negative sign. An asterisk denotes the addition of so much base that a precipitate was formed during the

measurement. In these cases, the potential decreased gradually after having been constant for a time. E, given in the next column, is the electromotive force in volts after subtraction of the asymmetry potential. Therefore, E may be considered the e. m. f. of the combination stated in scheme 2. $-\log(H^+)$, presented in the fourth column, has been computed by means of

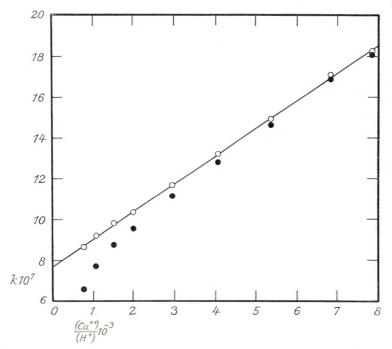


Fig. 3. Series 9: 0.200 m Cu(NO₃)₂. Solid circles: $\delta = 0$. Open circles: $\delta = 8 \times 10^{-5}$.

equation 4 and the values of A_0 in Table 1. The computations of $\frac{(Cu^{++})}{(H^+)}$ and k, given in the next two columns, were carried out as previously explained.

When k is plotted against $\frac{(Cu^{++})}{(H^{+})}$, the points for each series fall close to a straight line, in agreement with equation 14. The constants $(k_1 + ak_2)$ and k_3 determined from the plot are shown in the fourth and sixth column of Table 3. By means of these values and equation 14, the values of k given in the last column of Table 2 were calculated. The agreement between k found from the measurement and k calculated from formula 14 is satisfactory. Only in the least acid solutions of some of the

Table 2.

	1	abre				
	$x 10^4$	E –	$-\log(H^+)$	$\frac{(Cu^{++})}{(H^{+})}10^{-3}$	$k10^7$ found	$k10^7$ formula
	(-10.23)	0.0342	2.900	0.793	5.25	5.25
	-5.09	0.0424	3.042	1.100	5.27	5.28
	0.00	0.0509	3.190	1.546	5.35	5.32
Series 12	3.39	0.0567	3.290	1.945	5.31	5.36
$0.9994 \text{ m Cu}(\text{NO}_3)_2$	7.43	0.0626	3.392	2.459	5.42	5.41
$\delta = 18 imes 10^{-5}$	12.56	0.0693	3.508	3.208	5.46	5.48
	18.57	0.0754	3.614	4.093	5.59	5.57
	*25.50	0.0793	3.681	4.771	(6.18)	5.64
	10.00	010100				
	(-10.19)	0.0494	2.979	0.479	2.32	2.35
	-4.085	0.0658	3.263	0.921	2.36	2.40
	-2.034	0.0725	3.379	1.203	2.46	2.43
Series 11	0.000	0.0795	3.500	1.589	2.50	2.47
0.5031 m Cu (NO ₃) ₂	{ 1.390	0.0842	3.581	1.914	2.52	2.51
$\delta = 16 \times 10^{-5}$	3.768	0.0913	3.704	2.539	2.58	2.58
	7.038	0.0990	3.838	3.454	2.70	2.69
	11.70	0.1072	3.980	4.783	2.84	2.84
	*18.66	0.1155	4.123	6.636	3.06	3.06
	(-2.081)	0.0897	3.587	0.772	0.864	0.872
	-1.036	0.0980	3.730	1.073	0.920	0.913
	0.000	0.1066	3.879	1.512	0.981	0.973
	0.932	0.1138	4.004	2.015	1.034	1.041
Series 9	2.624	0.1235	4.172	2.963	1.169	1.170
$0.2000 \text{ m Cu}(\text{NO}_3)_2$	4.688	0.1314	4.309	4.057	1.320	1.319
$\sigma = 8 \times 10^{-5}$	7.465	0.1385	4.432	5.376	1.491	1.498
	11.19	0.1446	4.537	6.829	1.710	1.696
	13.87	0.1481	4.598	7.848	1.826	1.834
	*16.32					
	(-2.058)	0.0944	3.653	0.450	0.434	
	-1.033		3.870	0.741	0.468	0.477
	-0.520	0.1145	4.001	1.001	0.507	0.516
		0.1218	4.128	1.341	0.578	0.567
Series 5	0.704	0.1307	4.282	1.911	0.659	0.652
$0.1000 \text{ m Cu}(\text{NO}_3)_2$	1.641	0.1388	4.422	2.635	0.778	0.760
$\delta = 3 \times 10^{-5}$	2.797	0.1460	4.547	3.508	0.889	0.891
	4.243	0.1519	4.649	4.430	1.018	1.029
	5.854	0.1564	4.727	5.291	1.151	1.158
	9.305	0.1630	4.841	6.852	1.387	1.391
	*12.54	0.1672	4.914	8.077	1.578	

Table 2 (continued).

	rabie		ontinueu).			
	$x \ 10^4$	E	$-\log(H^+)$	$\frac{(Cu^{++})}{(H^{+})}10^{-3}$	$k10^7$ found	$k10^7$ formula
	(-1.028)	0.1113	3.946	0.442	0.270	0.268
	-0.518	0.1228	3 4.145	0.698	0.306	0.306
	0.000	0.1356		1.163	0.383	0.376
Series 6	0.479	0.1454		1.714	0.459	0.458
$0.05000 \text{ m Cu}(\text{NO}_3)_2$	1.180	0.1544		2.452	0.571	0.569
$+ 0.05000 \text{ m} \text{ Ba}(\text{NO}_3)_2$	2.122	0.1621		3.330	0.688	0.700
$artheta=3 imes10^{-5}$	3.293	0.1678		4.164	0.825	0.825
	4.663	0.1726		5.024	0.953	0.954
	7.014	0.1782		6.250	1.140	1.137
		012102		0.200		
	(-0.518)	0.1286	4.245	0.351	0.162	0.165
	-0.210	0.1423	4.483	0.608	0.206	0.203
Series 7	0.000	0.1522	4.654	0.900	0.253	0.247
0.02000 m Cu(NO ₃) ₂	0.421	0.1667	4.905	1.602	0.345	0.352
$+ 0.08000 \text{ m Ba}(\text{NO}_3)_2$	0.935	0.1756	5.059	2.277	0.452	0.453
$d = 3 \times 10^{-5}$	1.403	0.1807	5.148	2.788	0.531	0.529
	2.336	0.1876	5.267	3.650	0.658	0.658
	4.144	0.1954	5.402	4.933	0.851	0.850
						0.000
	(-1.036)	0.1115		0.447	0.271	0.280
	-0.518	0.1231		0.711	0.316	0.323
	-0.262	0.1292		0.907	0.363	0.355
	0.000	0.1353		1.155	0.409	0.395
Series 10	0.476	0.1448		1.687	0.482	0.481
0.05000 m Cu (NO ₃) ₂	{ 1.178	0.1537		2.401	0.595	0.596
$\delta = 8 imes 10^{-5}$	2.339	0.1623		3.376	0.749	0.754
	3.970	0.1694		4.465	0.925	0.930
	6.262	0.1756		5.684	1.126	1.127
	9.367	0.1811		7.043	1.349	1.346
	(*11.67	0.184	5.208	7.868	(1.500)	1.479
	0.590	0 1970	4.240	0.347	0.176	0.178
	(-0.520) (-0.262)	$0.1270 \\ 0.1377$		0.547	0.170 0.225	0.178 0.215
	$ - 0.202 \\ 0.000$	0.1377		0.864	0.225 0.274	0.213 0.271
				1.415	0.274 0.370	0.271 0.371
	0.377	0.1623			0.370 0.471	0.371
Series 8	0.836	0.1710		1.998		
0.02000 m Cu(NO ₃) ₂	1.401	0.1775		2.580		0.581
$\vartheta = 3 \times 10^{-5}$	2.102	0.1828		3.177	$\begin{array}{c} 0.684 \\ 0.807 \end{array}$	0.689
	3.040	0.1877		3.844		0.809
	4.204	0.1920		4.531	0.939	0.933
	5.599	0.1960		5.282	1.070	1.069 1.262
	8.178	0.2010		6.352	(1.296)	1.262
	*9.799	0.2036	5.566	6.985	(1.411)	1.376

Table 2 (continued).

	rabie	2 (00	munueu).			
	$x \ 10^4$	E	$-\log(H^+)$	$\frac{(Cu^{++})}{(H^{+})}10^{-3}$	$k 10^7$ found	$k10^7$ formula
	-0.414	0.1324	4.351	0.222	0.157	0.152
	-0.209	0.1450		0.367	0.174	0.181
	0.000	0.1593		0.649	0.240	0.236
Series 1	0.237	0.1716		1.058	0.315	0.316
$0.00991 \text{ m Cu}(\text{NO}_3)_2$	0.582	0.1815		1.565	0.414	0.414
$\delta=3 imes10^{-5}$	1.161	0.1906		2.236	0.540	0.545
	1.753	0.1959		2.747	0.653	0.645
l	2.322	0.2000		3.216	0.733	0.737
	-0.139	0.1553	4.748	0.279	0.147	0.139
	-0.0564	0.1649		0.408	0.164	0.164
Series 2	0.0001	0.1710		0.521	0.186	0.186
$0.004985 \text{ m } \text{Cu}(\text{NO}_3)_2$	0.158	0.1839		0.868	0.250	0.253
$+ 0.00493 \text{ m Ba}(\text{NO}_3)_2$	0.393	0.1940		1.294	0.334	0.336
$d = 3 \times 10^{-5}$	0.781	0.2034		1.868	0.433	0.447
0 = 0 × 10	1.171	0.2085		2.269	0.526	0.525
	1.550	0.2123		2.621	0.600	0.593
	1.000	0.2120	0.100	2.021	0.000	0.000
(-0.0423	0.1765	5.115	0.258	0.136	0.129
	0.0000	0.1850		0.361	0.153	0.150
	0.1331	0.2028		0.730	0.220	0.222
Series 3	0.2918	0.2128		1.080	0.287	0.290
$0.001980 \text{ m } \text{Cu}(\text{NO}_3)_2)$	0.5278	0.2211		1.484	0.364	0.367
$+ 0.00793 \text{ m Ba}(\text{NO}_3)_2$	0.8178	0.2273		1.874	0.442	0.446
$d = 3 \times 10^{-5}$	1.162	0.2322		2.238	0.523	0.517
0 0 10	1.779	0.2389		2.826	0.632	0.632
	2.330	0.2430			(0.725)	
	3.497	0.2498			(0.887)	
	0.107	0.2100	0.000	0.001	(0.001)	0.001
	-0.0521	0.1799	5.174	0.149	0.102	0.104
(-0.0211	0.1891		0.215	0.119	0.117
	0.0000	0.1954		0.276	0.132	0.129
	0.0884	0.2138		0.571	0.186	0.187
Series 4	0.1802	0.2231		0.820	0.235	0.235
$0.001001 \text{ m Cu} (\text{NO}_3)_2$	0.2918			1.060		0.282
$+ 0.00891 \text{ m Ba}(\text{NO}_3)_2$	0.5231	0.2384		1.458	0.363	0.360
$\delta = 3 \times 10^{-5}$	0.8161	0.2456		1.883	0.436	0.443
$\circ = 0 \land 10$	1.392	0.2530		2.371	(0.589)	
	2.334	0.2624		3.074	(0.761)	
	*3.488	0.264	6.63	2.782	(1.256)	
	*4.617	0.264	6.70	2.695	(1.200) (1.715)	
	1.017	0.400	0.70	2.000	(1.110)	0.004

	Table	e 2 (co	ontinued).			
	$x 10^4$	E	$-\log(H^+)$	$\frac{(Cu^{++})}{(H^{+})}$ 10 ⁻³	$k10^{7}$	$k 10^{7}$
				(H+) 10-3	Tound	formula
	(0.1600		0.349	0.143	0.153
		0.1711		0.542	0.187	0.192
		0.1823		0.845	0.253	0.254
		0.1924		1.258	0.345	0.337
Series 15		0.2001		1.702	0.437	0.428
$0.005007 \text{ m } \text{Cu}(\text{NO}_3)_2$	1.172	0.2071		2.227	0.540	0.534
$d = 18 \times 10^{-5}$	1.890	0.2139		2.880	0.666	0.667
	2.930	0.2203		3.638	0.812	0.821
	4.675	0.2272		4.605	1.020	1.017
4	7.026	0.2335		5.609	(1.258)	
	9.364 *11.71	0.2383		6.417 c.552	(1.464)	
		0.2403		6.553	(1.793)	
	-0.1034			0.166	0.127	0.136
	-0.0515			0.233	0.162	0.150
		0.1817		0.342	0.182	0.174
		0.1977		0.644	0.240	0.238
		0.2093		1.014	0.313	0.318
		0.2180		1.418	0.398	0.405
Series 14		0.2238		1.760	0.481	0.478
$0.002005 \text{ m Cu}(\text{NO}_3)_2$	1.167	0.2290		2.126	0.555	0.557
$\delta = 18 \times 10^{-5}$	1.708	0.2346		2.580	0.667	0.665
	2.335	0.2394		3.019	(0.777)	
	3.507	0.2464		3.727	(0.944)	
	4.679	0.2510		4.154	(1.129)	
	7.017 9.335	0.2527 0.2544		3.776	(1.862)	
	*13.62	0.2544 0.2537	$\begin{array}{c} 6.492 \\ 6.480 \end{array}$	$3.320 \\ 1.931$	(2.817)	
					(7.059)	
	-0.0515			0.146	0.132	0.134
	-0.0211			0.202	0.151	0.148
	0.0000			0.252	0.163	0.159
	0.1056			0.553	0.227	0.231
	0.2327			0.836	0.294	0.297
	0.4369			1.177	0.380	0.378
Series 13	0.7384	0.2387		$1.585 \\ 2.016$	0.471	0.474
$0.001007 \text{ m } \text{Cu}(\text{NO}_3)_2$	1.759	0.2439 0.2529		2.010 2.492	0.582 (0.709)	0.576
$d = 18 \times 10^{-5}$	2.334	0.2525		2.492 2.818	(0.709) (0.831)	
$0 - 10 \times 10^{-2}$	$2.554 \\ 3.258$	0.2578		2.818 2.625	(0.851) (1.244)	
	4.672	0.2604 0.2633		2.025 2.445	(1.244) (1.914)	
	5.850	0.2650		2.445 2.039	(1.914) (2.873)	
	7.001	0.2658		1.532	(4.574)	
	8.248	0.2653		1.002	(1.0/1)	0.104
	10.07	0.2657	6.699			
	*15.12	0.2725	6.817			
		.=0				

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series, values of k considerably greater than those corresponding to formula 14 are found. Such values are placed in parentheses. Especially in the most dilute solutions, which have the smallest hydrogen ion concentrations, the discrepancy may be due to an acid dissociation of the cupric ion which exceeds that expressed in the schemes 5 to 7. Thus, an ion of the composition $Cu_2O(OH)^+$ may possibly be formed. The discrepancy may also be due to a beginning precipitation. This question has not been examined more closely.

We now consider Table 3 where $(k_1 + ak_2)$ and k_3 found from the measurements are enumerated. With changing a, we may expect a variation of k_1 , k_2 , and k_3 owing to the change of the ionic strength μ . $(k_1 + ak_2)$ varies also as a direct consequence of the change of a. Within each of the two groups comprising series 5—7 and series 1—4, a varies, while μ is kept constant by replacing the cupric nitrate by barium nitrate. As may be seen from the table, k_3 is constant within each group, while $(k_1 + ak_2)$ varies linearly with a. From the latter relation we find

 $k_1 = 4.8 \times 10^{-9}$ and $k_2 = 3.16 \times 10^{-7}$, when $\mu = 0.300$, $k_1 = 7.6 \times 10^{-9}$ and $k_2 = 2.1 \times 10^{-7}$, when $\mu = 0.030$.

Table 3.

Comparison between the mass action constants found from the measurements and those calculated from formulae 16-18.

Series no.	$Cu(NO_3)_2$ molarity	$Ba (NO_3)_2$ molarity	$\frac{(k_1 + ak_2) 10^7}{\text{found}}$	$(k_1 + ak_2) 10^7$ formulae	$k_3 10^{12}$ found	$k_3 10^{12}$ formula
12	0.9994		5.17	5.17	4.9	5.9
11	0.5031		2.29	2.29	5.8	6.3
9	0.2000		0.767	0.774	6.8	6.9
5	0.1000		0.366	0.364	7.48	7.45
6	0.05000	0.05000	0.202	0.206	7.48	7.45
7	0.02000	0.08000	0.112	0.111	7.48	7.45
10	0.05000		0.208	0.193	8.08	8.09
8	0.02000		0.115	0.114	9.03	9.02
1	0.00991		0.109	0.097	9.76	9.73
2	0.004985	0.00493	0.085	0.086	9.70	9.73
3	0.001980	0.00793	0.079	0.080	9.78	9.73
4	0.001001	0.00891	0.075	0.078	9.78	9.73
15	0.005007		0.082	0.093	10.2	10.4
14	0.002005		0.100	0.095	10.8	11.1
13	0.001007		0.100	0.098	11.8	11.5

In the other cases, where μ changes from series to series, it is more difficult to estimate the separate values of k_1 and k_2 . However, it follows from the equilibrium

$$Cu^{++} + CuOH^+ \rightleftharpoons Cu_2OH^{+++}$$

that an increase in the concentration of cupric salt will favour the ion Cu_2OH^{+++} at the expense of $CuOH^+$. In more concentrated solutions (when a > 0.1), the concentration of the ion $CuOH^+$ is so small that k_2 may be calculated with sufficient accuracy from $(k_1 + ak_2)$, if k_1 can be estimated roughly. On the other hand, in the more dilute solutions (when a < 0.01), the ion Cu_2OH^{+++} is of so little importance that k_1 may be calculated from $(k_1 + ak_2)$ and a very rough value of k_2 .

We have now found k_1 , when $0.1 \ge a \ge 0.001$; k_2 , when $1 \ge a \ge 0.01$; and k_3 in the entire interval $1 \ge a \ge 0.001$. The values found for the three equilibrium constants may be expressed by means of the following formulae which, for very dilute solution, agree with the requirements of the DEBYE-HÜCKEL law:

$$-\log k_1 = 7.97 + \frac{0.996 \sqrt{\mu}}{1 + 1.0 \sqrt{\mu}}$$
(16)

$$-\log k_2 = 6.819 - \frac{0.996 \sqrt{\mu}}{1 + 1.3 \sqrt{\mu}}$$
(17)

$$-\log k_3 = 10.890 + \frac{0.996 \sqrt{\mu}}{1 + 2.36 \sqrt{\mu}}.$$
 (18)

The formulae contain two empirical constants each, while the third constant 0.996 is deduced from the Debye-Hückel theory. The values of $(k_1 + ak_2)$ calculated by means of formulae 16—17 are given in the fifth column of Table 3, while k_3 calculated from formula 18 is to be found in the last column of the table. A comparison of the calculated values with those obtained from the measurements shows good agreement. Only the values of k_3 for the two most concentrated solutions deviate appreciably from those calculated on the basis of formula 18.

If we extrapolate to the ionic strength 0 by means of the formulae 16-18, we obtain

 2^*

$$k_1^0 = 1.07 imes 10^{-8},$$

 $k_2^0 = 1.52 imes 10^{-7},$
d $k_3^0 = 1.29 imes 10^{-11}.$

and

We have seen that the assumption of the three equilibria 5-7 offers a satisfactory explanation for the hydrogen ion concentration of the cupric nitrate solutions. The existence of the three equilibria may be proved more directly in the following way: It has already been shown that the experimental values of k, defined by equation 15, satisfy equation 14, when the hydrogen and the cupric ion concentrations vary, if only the ionic strength is kept constant. k_1 , k_2 , and k_3 are constants, but they are not assumed to be equilibrium constants; it follows from this result that

$$(\mathbf{H^+}) + x + a \, \delta = k_1 \frac{(\mathbf{Cu^{++}})}{(\mathbf{H^+})} + k_2 \frac{(\mathbf{Cu^{++}})^2}{(\mathbf{H^+})} + 2 \, k_3 \frac{(\mathbf{Cu^{++}})^2}{(\mathbf{H^+})^2},$$

when (H⁺) and (Cu⁺⁺) vary, while μ is constant. The only possible explanation of this result is that hydrogen ions are produced by the three equilibria 5—7. The empirical constants k_1 , k_2 , and k_3 are the mass action constants of the equilibria.

The equilibrium constant k_1 is the acid strength of the hydrated cupric ion. At infinite dilution, it is 1.07×10^{-8} . According to J. BJERRUM⁽¹³⁾, the cupric ion in aqueous solution is a planar tetraquo ion which, through a more or less stable intermediate form, is in equilibrium with an octahedral hexaquo ion.

From the equilibria 6 and 7 we derive

$$Cu_2OH^{+++} \rightleftharpoons Cu_2O^{++} + H^+.$$

The mass action constant $\frac{k_3}{k_2}$ of this equilibrium is the acid strength of the ion Cu₂OH⁺⁺⁺. At infinite dilution, it is 8.5×10^{-5} . In the ion Cu₂OH⁺⁺⁺, the two copper atoms are bound together by the oxygen atom of the hydroxyl group. To the copper atoms are furthermore bound a certain number of water molecules which, as usual, have not been written. It is impossible to say whether the ion Cu₂OH⁺⁺⁺ splits off the hydrogen ion

from the hydroxyl group or from one of the water molecules. The question whether the ion Cu_2O^{++} contains a naked oxygen bridge, or a hydroxyl bridge and a hydroxyl group directly bound to one of the copper atoms, is therefore left open. The acid strength of the trivalent ion Cu_2OH^{+++} is about 8000 times that of the divalent ion Cu^{++} . A change in this direction might be expected on account of the electrostatic effect of the positive charges.

Owing to the fact that the maximum co-ordination number of oxygen is 3, the ion Cu_2OH^{+++} cannot take up another hydrogen ion. A dimeric cupric ion Cu_2^{++++} (omitting the water of hydration), therefore, does not exist.

From the equilibria 5 and 6 we derive

$$Cu^{++} + CuOH^+ \rightleftharpoons Cu_2OH^{+++}$$

with the equilibrium constant

$$\frac{k_2}{k_1} = \frac{(\text{Cu}_2\text{OH}^{+++})}{(\text{Cu}^{++})(\text{CuOH}^{+})}$$

which becomes equal to 14.2 at infinite dilution.

Table 4 contains the equilibrium constants k_1 , k_2 , and k_3 for cupric nitrate solutions, calculated from the formulae 16— 18, at round values of the concentration *a* molar. The fifth

Table 4.

The equilibrium constants, hydrogen ion concentrations, and degrees of "hydrolysis" α for pure aqueous solutions of cupric nitrate (a molar), calculated by means of the formulae 16–18.

			~			
а	$k_1 \ 1 \ 0^9$	$k_2 \; 10^7$	$k_3 \; 10^{12}$	$-\log(\mathrm{H}^+)$	$_{\rm pH}$	$\alpha \ 10^3$
1.000	2.51	5.15	5.90	3.137	2.85	1.43
0.500	3.02	4.49	6.25	3.456	3.36	1.34
0.200	3.98	3.67	6.87	3.856	3.85	1.22
0.100	4.79	3.16	7.45	4.125	4.13	1.17
0.0500	5.62	2.74	8.09	4.362	4.36	1.18
0.0200	6.76	2.32	9.02	4.634	4.62	1.36
0.0100	7.59	2.10	9.73	4.823	4.79	1.64
0.00500	8.32	1.93	10.35	5.004	4.95	2.08
0.00200	9.12	1.78	11.1	5.234	5.17	2.98
0.00100	9.55	1.71	11.5	5.405	5.33	3.99
0	10.7	1.52	12.9			

column shows $-\log(H^+)$ for a pure *a* molar cupric nitrate solution, calculated from the constants given in the table.

In their papers, most of the earlier investigators give the SØRENSEN pH of the cupric salt solution. Generally, the liquidliquid potential is partly eliminated by means of a bridge solution, such as concentrated potassium chloride or ammonium nitrate, and the remaining liquid-liquid potential is neglected. The results of the present measurements may be expressed in the same way if, instead of formula 4, the following formula is used

$$pH = 2.022 + 17.32 E, (19)$$

2.022 being the pH of the reference solution: 0.01000 m HCl + 0.0900 m NaCl. If E is eliminated from the equations 4 and 19, we obtain

$$pH = -\log(H^+) + 2.022 - A_0$$
.

The pH values given in the sixth column of Table 4 have been calculated from $-\log(H^+)$ by means of this formula. When these values are plotted against $\log a$, the curve of Fig. 1 is obtained.

The degrees of reaction ("hydrolysis") of the cupric ions according to each of the three equilibria 5-7 are

$$\alpha_{1} = \frac{(\text{CuOH}^{+})}{a} = \frac{k_{1}(\text{Cu}^{++})}{a(\text{H}^{+})},$$
$$\alpha_{2} = \frac{2(\text{Cu}_{2}\text{OH}^{+++})}{a} = \frac{2k_{2}(\text{Cu}^{++})^{2}}{a(\text{H}^{+})},$$
$$2(\text{Cu}_{2}\text{O}^{++}) = 2k_{3}(\text{Cu}^{++})^{2}$$

and

$$\alpha_3 = \frac{2 (\operatorname{Cu}_2 \operatorname{O}^{++})}{a} = \frac{2 k_3 (\operatorname{Cu}^{++})^2}{a (\operatorname{H}^{+})^2},$$

respectively. The total degree of reaction is $\alpha = \alpha_1 + \alpha_2 + \alpha_3$. The degrees of reaction have been calculated for pure cupric nitrate solutions by means of the equilibrium constants and the hydrogen ion concentrations given in Table 4. α is to be found in the last column of Table 4. When α_1 , α_2 , α_3 , and α are plotted against log α , the four curves of Fig. 4 are obtained. The figure shows how the relative importance of the three equilibria varies with the concentration of the cupric nitrate. Roughly speaking, the concentrations of the three ions are of

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the same order of magnitude, when 0.1 > a > 0.01. In more concentrated solutions, the ion Cu_2OH^{+++} gains more and more predominance, while the ion Cu_2O^{++} and, first and foremost, the ion $CuOH^+$ gradually disappear. In solutions more dilute

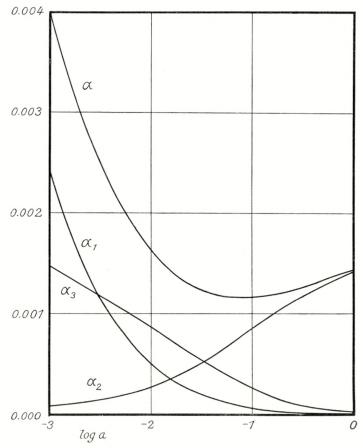


Fig. 4. The degrees of reaction ("hydrolysis") of the cupric ion in an *a* molar solution of cupric nitrate. $\alpha_1 = (CuOH^+)/a$ $\alpha_2 = 2(Cu_2OH^{+++})/a$ $\alpha_3 = 2(Cu_2O^{++})/a$

 $\alpha_{2} = \alpha_{1} + \alpha_{2} + \alpha_{3}.$

than 0.01 molar, the form Cu_2OH^{+++} gradually disappears. At sufficiently high dilution, only the monomeric ion $CuOH^+$ will remain.

I wish to express my thanks to the head of the laboratory, Professor NIELS BJERRUM, for advice and kind interest in my work.

Summary.

The hydrogen ion concentration of aqueous solutions of cupric nitrate was measured at 18.0° C. by means of the glass electrode. The range of concentrations examined extended from 0.001 to 1 molar cupric nitrate. The examination included both aqueous solutions and solutions to which small concentrations of nitric acid or sodium hydroxide had been added.

The measurements show that the following equilibria take place

and

$$Cu^{++} \rightleftharpoons CuOH^{+} + H^{+},$$

$$2Cu^{++} \rightleftharpoons Cu_{2}OH^{+++} + H^{+},$$

$$2Cu^{++} \rightleftharpoons Cu_{2}O^{++} + 2H^{+}.$$

The mass action constants of the equilibria have been computed.

From the Chemical Laboratory of the Royal Veterinary and Agricultural College. Copenhagen, Denmark.

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