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F.A.H. Schreinemakers & Thonus, J.C., The system HgCl2-CuCl2-H2O, in: KNAW, Proceedings, 15 I, 1912, 1912, pp. 472-474					
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If, for the sake of brevity, we call Ba $Cl_2.2H_2O = Ba_2$ and $Cu Cl_2.2H_2O = Cu_2$, the reaction is then:

$$\begin{array}{c|c} Ba_2 + Cu_2 + D_{1.2.2} \rightleftarrows D_{1.1} + Solution \\ Ba_2 + Cu_2 + D_{1.2.2} + Sol. \\ Ba_2 + Cu_2 + D_{1.2.2} + D_{1.1} & Ba_2 + Cu_2 + D_{1.1} + Sol. \\ Ba_2 + Cu_2 + D_{1.2.2} + D_{1.1} & Cu_2 + D_{1.2.2} + D_{1.1} + Sol. \end{array}$$

Hence, of the invariant (P) equilibrium two monovariant (P) proceed to lower and three to higher temperatures, or if we only consider the systems in which a solution occurs, one to lower and three to higher temperatures.

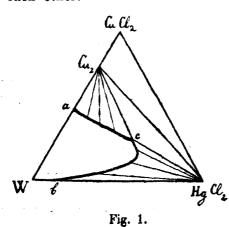
The system proceeding to lower temperatures: $Ba_2 + Cu_2 + D_{1.2.2} + sol.$ still exists at 40° and is represented in fig. 1 by the point h. The system proceeding to higher temperatures: $Cu_2 + D_{1.2.2} + D_{1.1} + solution$ terminates at 57° , when the solution only still contains the three components $CuCl_2$, KCl and water.

The other two systems proceeding to higher temperatures still exist at 60° ; the solution of the system $Ba_2 + Gu_2 + D_{1.1} + solution$ is represented in fig. 2 by the point l and that of the system: $BaCl_2 + D_{1.2.2} + D_{1.1}$ is indicated in fig. 2 by the point h.

Chemistry. — "The system $HgCl_1$ — $CuCl_2$ — H_2O ." By Prof. F. A. H. Schreinemakers and J. C. Thonus.

(Communicated in the meeting of September 28, 1912).

In order to ascertain whether or not the salts HgCl, and CuCl, form a double salt, the isotherm of 35° was determined; the result of this investigation is that, at 35° no double salt was found but that the salts HgCl, and CuCl, 2H,O can exist by the side of each other.



In fig. 1, the experimentally determined isotherm of 35° is represented schematically; the apexes W, HgCl, and CuCl, represent the three components, and the point Cu, the hydrate CuCl, 2 H,O.

The isotherm consists of the two branches ac and bc; ac indicates the solutions which are saturated with the hydrate CuCl₂. 2 H₂O, bc those saturated with

 $HgCl_1$; the point of intersection c of the two saturation lines represents the solution saturated with $CuCl_2 \cdot 2H_2O + HgCl_2$.

The solubility curve bc of the HgCl, has a peculiar form; for a tangent may be drawn to it parallel to the side W. CuCl₂. This means, in our case, that the solubility of HgCl₂ first increases and then decreases with an increased CuCl₂-content of the solution. From the Fig. 1 it is shown that the solubility of HgCl₂ is much increased by addition of CuCl₂; from the table we see that the solubility of HgCl₃, which in pure water amounts to 8.51 °/₀ can increase to fully 52 °/₀ by addition of CuCl₂.

The isotherm represented schematically in fig. 1 can be drawn with the aid of the determinations communicated in the table. As not only the compositions of the liquids, but also those of the corresponding "residues" have been determined, the composition of the solid substance may be deduced therefrom. We find that the solutions of branch ac are saturated with CuCl₂. 2 H₂O and those of bc with HgCl₂.

Compositions in % by weight at 35°.

solid phase	of the residue		of the solution	
	$^0\!/_0\mathrm{HgCl}_2$	⁰ / ₀ CuCl ₂	⁰ / ₀ HgCl ₂	% CuCl ₂
CuCl ₂ . 2H ₂ O	_		0	44.47
"	13.04	51.0	21.03	33.5
n.	16.97	55.82	37.3	26.07
,	19.70	54.77	44.47	23.31
CuCl ₂ .2H ₂ O + HgCl	36.63	43.60	50.45	21.49
, + ,	74.35	15.08	50.60	21.47
, + ,	_		50.37	21.54
$HgCl_2$	91.94	3.0	52.44	19.40
n	87.57	4.6	52.54	18.48
n	90.06	3.17	52.81	18.06
'n	_	_	51.03	14.73
n	<u></u> -	غست	49.5	5.94
#	_		23.87	2.64
n ,	. /		8.51	0

One of us has previously deduced the rule ') that the metastable continuations of the branches ac and bc must fall both together either within or without $\angle Cu$, $c \cdot HgCl_1$. Which case occurs here is difficult to prove experimentally as both branches, in the vicinity of point c practically coincide with the sides of the angle Cu, $c \cdot HgCl_1$. Moreover, the saturation line bc of the $HgCl_1$ exhibits a very peculiar form. The metastable continuation must, of course, terminate somewhere on the side $HgCl_1$ — $CuCl_1$ of the triangle; from the course of the stable part in the vicinity of c, it appears, however, that this will not be possible without a point of inflexion appearing somewhere on the metastable part or on the stable part situated in the vicinity of c.

Chemistry. — "The system Tin-Iodine". By Prof. W. Reinders and S. de Lange. (Communicated by Prof. Schreinemakers.)

(Cummunicated in the meeting of September 28, 1912).

1. Of tin and iodine two compounds are known, stannous and stannic iodide. As regards the preparation and properties of these compounds there exist in the literature different conflicting statements. By the older investigators 3), for instance, it is stated that on heating tin with iodine, stannous iodide is formed. Henry 3), however finds a mixture of SnI, and SnI, and Personne 4) SnI, only. The melting point of SnI, is given by Personne 4) as 145° (solidifying point 142°), by Emich 4) 143°. The boiling point according to Personne is at 295°, Emich finds 341°. Henry, however, states that it sublimes at 180°.

Of Snl₂ the melting point is given both at 246 ⁷) and at a dull red heat (Personne) and the boiling point both at 295° ⁷) and at the temperature of molten glass (Personne).

For the knowledge of the binary systems of a metal and a metalloid a renewed investigation was therefore desirable.

2. SnI_4 was prepared in two ways, a. by treating granulated tin for some days with a solution of iodine in carbon disulphide and

¹⁾ F. A. H. Schreinemakers, Die heter. Gleichg. von Bakhuis Roozeroom. III'. 268.

²⁾ I. a. Berzelius, Traité de chimie; Rammelsberg, Pogg. Ann. 48, 169.

³⁾ Phil. Trans. 135, 363 (1845).

⁴⁾ Compt. rendus. 54, 216 (1862).

⁵⁾ l. c.

⁶⁾ Sitzungsber, der K. Ak. v. W. Wien 118, IIb, 535 (1904) (Monatshefte 25, 907.

⁷⁾ Cohen, Abegg's Handbuch d. anorg. Ch. III. 2, 571.