

RÖNTGEN ANALYSIS OF COPPER AMALGAMS.

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In the author's previous paper⁽¹⁾ the following results were published:

- (1) The solid phase existing in high mercury amalgams has γ structure⁽²⁾ and the lattice dimension is $a=9.401\text{\AA}$.
- (2) The solid phase prepared by squeezing high mercury amalgams under high pressures has the dimension just described.
- (3) The upper copper limit of the homogeneity range of the γ phase lies somewhere between 27.1-29.9 weight per cent of copper.

(1) N. Katoh, *Z. physik. Chem.*, [B] **6** (1920), 27.

(2) See Westgren and Phragmen, *Phil. Mag.*, [6] **50** (1925), 311.

(4) High copper amalgams consist in copper and the γ phase at ordinary temperature.

The samples used in the present investigation were prepared at the Mineralogical Institute in Oslo.⁽¹⁾

For the powder photograms the Cr-K-radiation were used from a tube of the Siegbahn type, which was run with about 45 kilo-volts and 7.5ma. All cameras were designed by Dr. Phragmen.⁽²⁾ The powder photograms were taken with three cameras of the focussing type, covering whole diffraction range. The cameras were calibrated with sodium chloride. The exposure time varied 5 to 9 hours. All the dimension values based on the length 5.6280Å for the edge of the elementary cube of the sodium chloride.

The photograms were rather weak by want of the practice of handling cameras. The result of the measurement and calculation was shown in the Table 1 and confirmed the former view.

In the case of a good photogram on which every line can be measured accurately, the determination of the lattice dimension may be carried out by means of the most deviated interferences.⁽³⁾ The figures from each interference agree very well and it needs not farther calculation. But it is reasonable to take all the interferences when the photogram was weak.

The author used three photograms taken by the different cameras for determination of the γ phase. Taking $\text{tg}^2\theta$ as the weight of the each observation of the interference lines, the constant χ in the equation $\sin^2\theta = \chi (h^2 + k^2 + l^2)$ calculated for each photogram by the following formula.⁽⁴⁾

$$\chi = \frac{\sum \chi \text{tg}^2 \theta}{\sum \text{tg}^2 \theta}, \quad \Delta\chi = \pm \sqrt{\frac{\sum [(\chi - \chi_0) \text{tg}^2 \theta]^2}{(n-1) \sum \text{tg}^2 \theta}}$$

For the photogram taken by camera 5, $\chi = 0.01481 \pm 0.00001$

„ „ „ „ „ 6, $\chi = 0.01475 \pm 0.00003$

„ „ „ „ „ 8, $\chi = 0.01478 \pm 0.00001$

Then the most probable value of the constant was found by taking the weighted mean of the three figures as follows.

$$\chi = 0.01479 \quad (\text{for Cr-K}\alpha)$$

and the dimension of the γ phase is

(1) N. Katoh, loc. cit.

(2) Westgren and Phragmen, loc. cit.

(3) G. Hägg, Dissertation, 5, 1929 (Uppsala).

(4) E. Broch, *Z. physik. Chem.*, **127** (1927), 452.

$$a=9.399 \text{ \AA}$$

$$a=9.401 \text{ \AA} \quad (\text{by Debye cameras})^{(1)}$$

Tammann⁽²⁾ observed the existence of the α phase by the thermal analysis but the present author could not find it in the specimens tempered at 100°C. for 48 hours. The powder photograms of a 70.5% Cu amalgam quenched at 380°C, and pure copper tempered at 100°C. taken with the same camera were compared. All the copper lines agree very well, so that the existence of α phase at this temperature is doubtful one but it needs farther investigation to settle this question.

Table 1.

Cr. Radiation.

Cu—Hg (24.1% cu).

| Intensity | Radiation | $h k l$ | $\text{Sin}^2 \frac{\theta}{2} \cdot 10^4$ | $((h^2 + k^2 + l^2) \times 10^5)$ |
|-----------|------------------|---------------|--|-----------------------------------|
| w | β | 310 | 1231 | 10.1231 |
| s | α | 310 | 1482 | 10.1482 |
| w | β | 321 | 1722 | 14.1230 |
| $v w$ | α | 322 | 1777 | 12.1481 |
| s | α | 321 | 2072 | 14.1480 |
| m | β | 330, 411 | 2246 | 18.1248 |
| s | α | 330, 411 | 2660 | 18.1478 |
| $v w$ | β | 332 | 2730 | 22.1241 |
| $v w$ | α | 332 | 3238 | 22.1472 |
| w | α | 442, 600 | 5336 | 36.1482 |
| w | β | 622 | 5402 | 44.1228 |
| m | α | 532, 611 | 5616 | 38.1478 |
| m | $\alpha + \beta$ | 631 | 5669 | 46.1233* |
| $v w$ | β | 444 | 5870 | 48.1223 |
| $v w$ | β | 543, 550, 710 | 6099 | 50.1220 |
| $v w$ | α | 541 | 6187 | 42.1473 |
| m | α | 622 | 6504 | 44.1478 |
| m | α | 631 | 6787 | 46.1475 |
| s | α | 444 | 7081 | 48.1475 |
| m | α | 543, 550, 710 | 7361 | 50.1472 |
| w | α | | 7762 | * |
| s | α_1 | 552, 633, 721 | 7981 | 54.1478 |
| w | α_2 | 552, 633, 721 | 8010 | 54.1483 |
| $v w$ | β | 811, 741, 554 | 8123 | 66.1231 |
| w | α | 642 | 8326 | 56.1483 |
| w | α_1 | 651, 732 | 9126 | 62.1471 |
| w | α_2 | 651, 732 | 9160 | 62.1478 |

* Extra lines.

(1) W. Katoh, loc. cit.

(2) G. Tammann and Th. Stassfurth, *Z. anorg. Chem.*, **143** (1925), 357.

$$\begin{aligned}
 \text{Cr K}_{\alpha_1} &= 2.2848 \text{Å} & \sin^2 \vartheta &= 0.01479 (h^2 + k^2 + l^2) & \text{for CrK}_{\alpha} \\
 \text{Cr K}_{\alpha_2} &= 2.2890 \text{Å} & a &= 9.399 \text{Å} \\
 \text{Cr K}_{\beta} &= 2.0805 \text{Å} \\
 \text{Cr K}_{\alpha} &= 2.2862 \text{Å}
 \end{aligned}$$

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