PREPARATION OF BARIUM TITANATE

| Expt. No. | Compos | sition of charge in mg | Pressure bomb | Temp. °C | Pressure atm. | Time h | Product | Largest crystal dimension mm |
|--------------|--|---------------------------|------------------|-------------|------------------|-----------|---------------------------|---------------------------------------|
| 11 | BaTiO ₃ 38 | 15 M NaOH 50 | D | 660 | 1120 | 92 | BaTiO ₃ | 0.1 |
| 12 | ${f BaTiO_3}\ 73$ | 15 M NaOH 270 | D | 460 | 1430 | 65 | BaTiO ₃ | 0.1 |
| 13 | $\begin{array}{c} \mathrm{BaTiO_3} \\ 500 \end{array}$ | 4 M NaOH 1000 | D | 605 | 3150 | 114 | BaTiO ₃ | 0.25 |
| 14 | ${f BaTiO_3}$ 789 | 4 M NaOH 1200 | D | 600 | 2920 | 500 | BaTiO_3 | 1 |
| 15 | BaTiO ₃ 1260 | 4 M NaOH 1200 | D | 620 | 3030 | 500 | BaTiO ₃ | 1 |
| 16 | $\operatorname{BaTiO_3}$ 1000 | 4 M LiOH 1100 | D | 625 | 1200 | 168 | BaTiO_3 | 0.01 |
| 17 | BaTiO_3 1000 | 4 M KOH 1225 | D | 625 | 2040 | 168 | BaTiO ₃ | 0.25 |
| 18 | $\frac{\mathrm{BaTiO}_3}{1000}$ | 4 M LiF 1100 | D | 605 | 1150 | 168 | no transport of $BaTiO_3$ | |
| 19 | BaTiO_3 1000 | 4 M NaF 1170 | D | 625 | 2720 | 168 | ${f BaTiO_3}\ {f BaF_2}$ | $0.5 \\ 0.5$ |
| 20 | ${f BaTiO_3}\ 1292$ | 2 M KF 1200 | D | 590 | 2800 | 500 | BaTiO ₃ | 1 |
| 21 | ${f BaTiO_3}\ 1072$ | 2 M KF 1200 | D | 620 | 3060 | 500 | BaTiO ₃ | 1 |

Table 2. Hydrothermal recrystallisation of barium titanate by transport reactions.

D: 6 ml pressure bomb, lined with pure gold. Dimensions inside: length 190 mm, diameter 6.25 mm. The temperature was measured using chromel-alumel thermocouples.

Hydrothermal synthesis of BaTiO₃

Expts. Nos. 1-3. Crystalline titanium oxide reacts with barium hydroxide to form $BaTiO_3$ (1), but with a barium chloride solution no reaction to $BaTiO_3$ is obtained under the same experimental conditions. In the hydrothermal synthesis of barium titanate the solution should be alkaline. In order to improve the crystal formation, freshly precipitated titanium oxide gel was used in the charge and the hydrothermal solution contained sodium hydroxide as well as barium hydroxide. The powder pattern of the product from (2) was diffuse but the product from (3), where the reaction temperature was higher than in (2), gave a sharp powder pattern and small crystals could be observed in the product.

Expts. Nos. 4-6. Titanium esters were used in the charge. In (4) and (6) the ester was $Ti(OC_2H_5)_4$, in (5) the ester was $Ti(OCH_2CH(CH_3)_2)_4$. The powder pattern of the product from (4) was diffuse, but the powder pattern of (6), where the hydrothermal solvent contained sodium hydroxide, was very sharp. Apparently it is important that the reaction temperature is over $400-500^{\circ}C$ and that the mole percent of sodium hydroxide is at least twice as much as that of the titanium compound. The main part of the product from (5) was barium carbonate, $BaCO_3$. The alcohol part of the ester was oxidised to form carbon dioxide. (When a solution of 3.1 g freshly recrystallised barium

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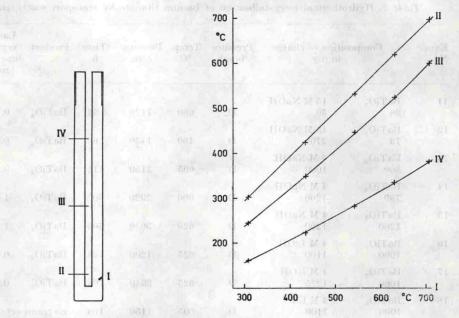


Fig. 1. Longitudinal slice of pressure vessel D. Inside diameter is 6.25 mm, inside length is 190 mm. When the pressure vessel is used the temperature is measured in the thermocouple well at I. A gold ampoule with the charge is inserted in the vessel. II is the position of the nutrients and III is the position of the crystals formed in the hydrothermal recrystallisation. Fig. 2. Plot of the temperatures at positions II, III, and IV (see Fig. 1) measured in air versus the temperature at position I, (the thermocouple well). In the temperature range from 300°C to 700°C the temperature difference between I and II does not exceed 10°C.

hydroxide, $Ba(OH)_28H_2O$, in 40 ml of carbon dioxide free water was treated with 3 ml of ethanol for 72 h at 425°C and at a pressure of 655 atm, barium carbonate crystals were formed).

Expts. Nos. 7-8. The products from (7) and (8) gave sharp powder patterns. The unit cell parameters of barium titanate obtained from the powder pattern of (8) are a=3.994(1) Å, c=4.035(2) Å at 25°C.

Expts. Nos. 9-10. Potassium fluoride was used in the solvent instead of sodium hydroxide. This resulted in the formation of barium titanate and barium fluoride, BaF₂. The product from (9) had a blue colour, and the powder lines of barium titanate were diffuse. A temperature rise (10) improved the crystal size considerably. However, the product was blue in contrast to the usual yellow colour of barium titanate.

Hydrothermal crystal growth of BaTiO₃ by transport reactions

Expts. Nos. 11-12. Barium titanate was prepared by hydrolysis of Ti(OCH₂CH₍CH₃)₂)₄ with a solution of barium hydroxide using the preparation procedure described by Flaschen.¹ Single crystals of barium titanate were grown from a melt of 3 g BaTiO₃

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