

85. A High Pressure Hexagonal Form of MgSiO_3

By Naoto KAWAI,^{*)} Masaharu TACHIMORI,^{*)} and Eiji ITO^{**)}

(Comm. by Tei-ichi ITO, M. J. A., June 11, 1974)

The high-pressure and high-temperature state of MgSiO_3 has been studied in a spherical pressure vessel made up of two stages.¹⁾ Chemically pure clinoenstatite is synthesised and charged into the vessel together with a platinum heater and Pt-Pt13%Rh thermo-couple as shown in Fig. 1.

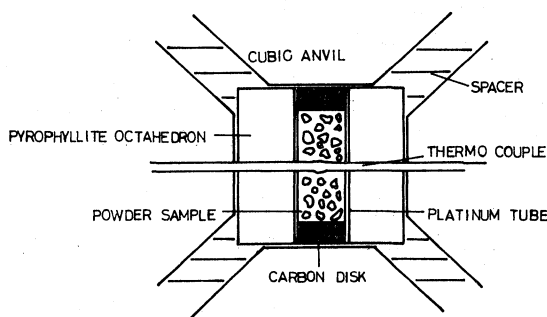


Fig. 1. Furnace assembly for quenching runs.

After the sample is compressed slowly up to the pressure higher than the metallic transition point of GaP (0.5 Mb), the electric current is driven through the heater, till the specimen temperature becomes 1000°C . Then, the P-T condition is kept constant for 10 hours. When the current is cut off, the sample is quenched, so that the specimen may be taken out of the vessel safely under the atmospheric pressure.

The quenched specimen is colourless aggregate of polycrystalline powder whose X-ray diffraction lines are given in Fig. 2. There appear more than 10 extra peaks in addition to those characteristic of $\gamma\text{-Mg}_2\text{SiO}_4$ ³⁾ and stishovite. They can be indexed on the basis of a hexagonal lattice as in the case of MgGeO_3 produced at 30 Kb by Ringwood and Seabrook⁴⁾ in 1962 (X-ray data are given in Table I).

Recently we have found⁵⁾ that clinoenstatite (MgSiO_3) breaks up at about 200 Kb into stishovite and $\beta\text{-Mg}_2\text{SiO}_4$, the latter being transformed further into $\gamma\text{-Mg}_2\text{SiO}_4$. The transformation is accompanied

^{*)} Department of Material Physics, Faculty of Engineering Science, Osaka University, Toyonaka, Osaka.

^{**)} Institute for Thermal Spring Research, Okayama University, Misasa, Tottori.

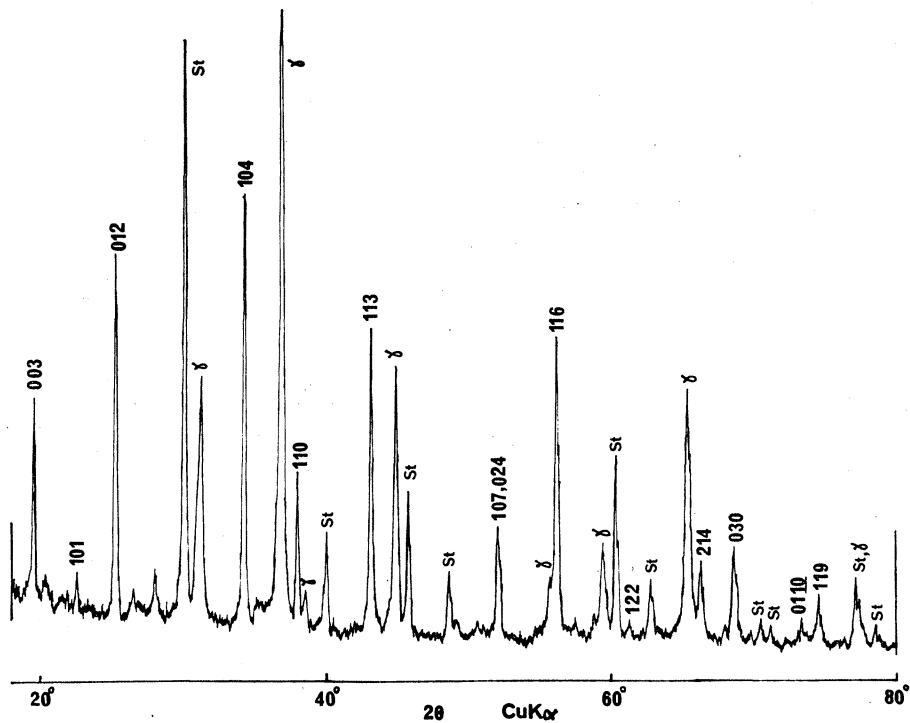


Fig. 2. Powder X-ray diffraction chart of the quenched sample.

Table I. Powder data for hexagonal MgSiO_3

h	k	l	d_{obs}	d_{calc}	Intensity
0	0	3	4.512 A	4.519 A	50
1	0	1	3.919	3.919	10
0	1	2	3.501	3.504	80
1	0	4	2.609	2.611	100
1	1	0	2.364	2.364	30
1	1	3	2.094	2.094	70
1	0	7	1.7521	1.7506	} 30
0	2	4	1.7521	1.7522	
1	1	6	1.6329	1.6332	70
1	2	2	1.5087	1.5086	<5
2	1	4	1.4076	1.4076	20
0	3	0	1.3647	1.3647	20
0	1	10	1.2867	1.2869	5
1	1	9	1.2703	1.2702	10

Hexagonal: $a = 4.727 \pm 0.001 \text{ \AA}$, $c = 13.556 \pm 0.002 \text{ \AA}$,
 $V = 262.3 \pm 0.5 \text{ \AA}^3$, $Z 6$, $\rho_{\text{calc}} = 3.81 \text{ g/cm}^3$

by the contraction of volume of about 15%. The hexagonal MgSiO_3 obtained by us is 2% greater in density than the mixture of $\gamma\text{-Mg}_2\text{SiO}_4$

and stishovite above mentioned.

γ - Mg_2SiO_4 and stishovite, as we have found in the X-ray chart, coexisting with the hexagonal MgSiO_3 show that the reaction in our experiment for the new phase has been incomplete. The hexagonal metasilicate has not been found in nature. However, that it has been produced and is stable under a pressure less than 0.5 Mb would indicate its possible existence in the earth's mantle.

References

- 1) Kawai, N., M. Togaya, and A. Onodera: Proc. Japan Acad., **49**, 623 (1973).
- 2) Onodera, A., N. Kawai, K. Ishizaki, and Ian L. Spain: Solid State Commun., **14**, 803 (1974).
- 3) Ito, E., Y. Matsui, K. Suito, and N. Kawai: Phys. Earth Planet. Inter. (in press).
- 4) Ringwood, A. E., and M. Seabrook: J. Geophys. Res., **67**, 1960 (1962).
- 5) Ito, E., T. Matsumoto, K. Suito, and N. Kawai: Proc. Japan Acad., **48**, 412 (1972).