

Title	Vaporization Behavior in the Lithium Oxide- Aluminum Oxide System
Author(s)	Iwamoto, Nobuya; Ikeda, Yasushi; Kamegashira, Naoki et al.
Citation	Transactions of JWRI. 1979, 8(2), p. 287-288
Version Type	VoR
URL	https://doi.org/10.18910/5518
rights	
Note	

Osaka University Knowledge Archive : OUKA

https://ir.library.osaka-u.ac.jp/

Osaka University

Vaporization Behavior in the Lithium Oxide-Aluminum Oxide System†

Nobuya IWAMOTO*, Yasushi IKEDA**, Naoki KAMEGASHIRA*** and Yukio MAKINO****

KEY WORDS: (Blanket Material) (Lithium Oxide) (Mass Analysis)

Lithium oxide has a potentiality for the blanket materials in nuclear fusion reactor 1 and high temperature behaviors of the systems with lithium oxide have been investigated. Besides pure Li_2O single solid, lithium aluminates are also considered to be useful for the blanket material. There exist several intermediate compounds, Li_5AlO_4 , LiAlO_2 and LiAl_5O_8 , in $\text{Li}_2\text{O-Al}_2\text{O}_3$ system, and several properties have been reported 2 2 2 . In this work Knudsen-effusion and mass spectrometric study of this system with various composition are undertaken, and some preliminary results are described.

The samples with each composition were prepared by usual solid state sintering method in vacuo at 1000°C of coprecipitation from aqueous solution of LiNO, and Al(NO₃)₃ and drying at 700°C⁸). Each sample was pre-heated in vacuo in Knudsen cell at a little higher temperature than the experimental temperature before the measurement. The samples with various composition were taken for chemical and X-ray analysis. The intermediate compounds such as LiAlO2 and LiAl5O8 were determined clearly, but the presence of Li₅ AlO₄ was not decisive by the interference of LiAlO₂. The Knudsen cell was made of platinum with an orifice of 0.5 mm diameter and is shown in Fig. 1. The platinum was found to be well compatible with this system at high temperature. The cell was heated by a tungsten resistance heater and the temperature of the Cell was measured by a W-WRe or a Pt-PtRh thermocouple.

Quadruppole type mass spectrometer, NEVA° II, was utilized for a preliminary measurement at a first step of this work, and 90°-sector magnetic type mass spectrometer, HITACHI RMU-K, was exclusively used in



^{*} Professor

**** Instructor

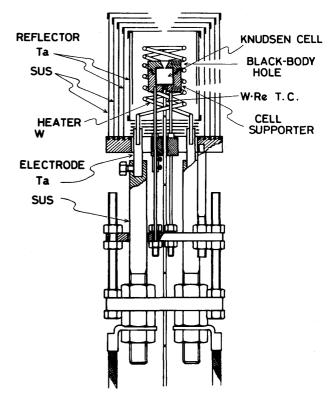


Fig. 1 Knudsen cell assembly

this experiment. The vapor pressure was calibrated with silver. The detailed procedure is described elsewhere 9). Two major species of vapor molecule, Li⁺ and Li₂O, were detected along with some minor species such as LiO, Li₃O and Li₂O₂ over 100% Li₂O in this experiment with shutter profile. The ionizing electron potential was selected to be 30 eV. The temperature variation of partial

Transactions of JWRI is published by Welding Research Institute of Osaka University, Suita, Osaka, Japan

^{**} Co-operative Researcher (Department of Nuclear Energy, Faculty of Engineering, Nagoya University)

^{***} Co-operative Researcher (School of Material Science, Toyohashi University of Technology)

pressures of these main species, plotted against the reciprocal temperature, is shown in Fig. 2, where the concentration of Li₂O was 64 mole %. Other vapor species were scarcely detected. The intensity of Li₂O and other minor species ions decreased with decreasing concentration of the lithium oxide in the sample and only Li⁺ ion was detectable for alumina rich composition. The vapor pressure of Li was plotted against the composition at 1600K, as is shown in Fig. 3. It is shown from this figure that the high temperature stability of lithium oxide increases with increasing aluminum oxide. Also it is noted

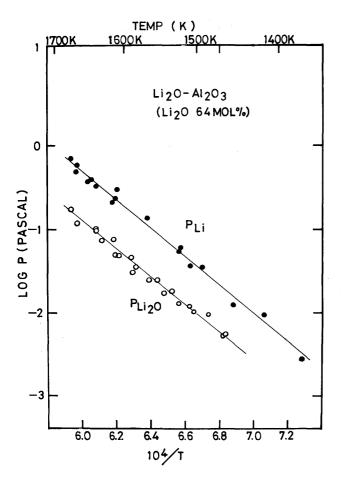


Fig. 2 Temperature dependence of Li and Li₂O pressures in the lithium aluminate containing 64 mol % Li₂O

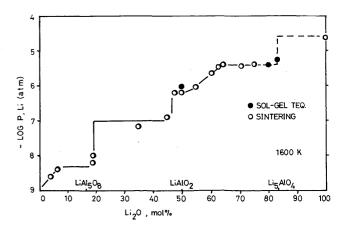


Fig. 3 Dependence of Li pressure upon Li_2O content in $\text{Li}_2\text{O-Al}_2\text{O}_3$ system

that the existence of some degree of solid solubility is suggested from the composition near 50 to 62 mole %. The possibility of solid solubility of lithium in LiAlO₂ at 1600K is inferred from this figure. The detailed discussion based on thermodynamics will be described.

Acknowledgement

This work supported by Science Research Grant of Ministry of Education.

References

- K. Sako, M. Ohta, Y. Seki, H. Tamato, T. Hiraoka, K. Tanaka, N. Asami and S. Mori, JAERI-M 5502. (1973).
- 2) M. Maryio, Acta. Cryst. 19 (1965) 396.
- C.H. Chang and J.L. Margrave, J.Am. Chem. Soc. 90 (1968) 2020
- 4) M. Marezio and J.P. Remeika, J. Chem. Phys. 44 (1966) 3143.
- 5) G.H. Stauss, J. Chem. Phys. 40 (1964) 1988.
- 6) R.K. Datta and Rustum Roy, J. Am. ceram. Soc. 46 (1963) 388.
- A.LA. Ginstra, M.Lo Jacono and P. Porta, J. Thermal Analysis.
 4 (1972) 5.
- 8) D.W. Strickler and Rustum Roy, J.Am. Ceram.Soc. 44 (1961) 225.
- 9) Mass Spectroscopy 26 (1978) 159.