

### THE EVAPORATION PECULIARITIES OF Ca–Al-INCLUSIONS OF CHONDRITES.

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Calcium and aluminium rich inclusions of chondrites (CAIs) are unique geochemistry objects. CAIs represent the most primitive matter of the Solar system with the age of 4.567 billion years.

During the high-temperature evaporation of CAIs melts (consisting mainly of  $\text{SiO}_2$ ,  $\text{MgO}$ ,  $\text{CaO}$  and  $\text{Al}_2\text{O}_3$  oxides), there are the inversion of the volatility of  $\text{SiO}_2$  and  $\text{MgO}$ . As a matter of fact the silicon oxide being more volatile than magnesium oxide are becoming less volatile under the evaporation conditions of melts enriched of  $\text{CaO}$  and  $\text{Al}_2\text{O}_3$  oxides. The volatility inversion is well explained in the framework of the Korginsky's theory of the acid-base interaction of components in silicate melts [1]. According to the theory, the increase of the  $\text{CaO}$  content in the melt of the  $\text{CaO-MgO-Al}_2\text{O}_3\text{-SiO}_2$  system leads to increasing of melt basicity, that in turn leads to increasing the  $\text{MgO}$  thermodynamic activity and volatility, and to decreasing activity and volatility of the  $\text{SiO}_2$ . In the first approximation the evaporation process of CAIs melts can be described in terms of the ternary CMAS oxide system. The oxides are divided into the next groups on acid-basic properties: acidic ( $\text{SiO}_2$ ), amphoteric ( $\text{Al}_2\text{O}_3$ ) and basic ( $\text{CaO}$  and  $\text{MgO}$ ). Moreover, the  $\text{CaO}$  as the main basic oxide of melts, is a major donor to oxygen ions, and its concentration indirectly specifies the basic index of the melts.

It follows: 1) the concentration increasing of calcium oxide in the melt should allow: a) to increase activity and the volatility of  $\text{MgO}$ , and b) to the drop of the activity and the volatility of  $\text{SiO}_2$ . The activity calculation carried out according to [2] and [3], illustrates the forecast of the theory and clearly shows the inversion in the activity values of  $\text{SiO}_2$  and  $\text{MgO}$ , depending on the  $\text{CaO}$  content in the melt (Fig. 1); 2) during the evaporation the increasing of calcium oxide concentration in the melt should lead to a reduction in the  $\text{MgO}$  content and relative increase of the  $\text{SiO}_2$  content. As a result of this process the ratio of  $\text{MgO} : \text{SiO}_2$  in the residual melt should decrease, as observed in the real Ca–Al-inclusions.

As shown by the analysis of the isotopic mass fractionation data of magnesium and silicon in the residue after evaporation of the melts [4], acid-basic factor has a strong influence on the efficiency of the separation isotopes. Fig. 2 clearly shows that the efficiency of isotope separation of magnesium compared to silicon isotopes in the CMAS melts increases with an increasing content of  $\text{CaO}$ .

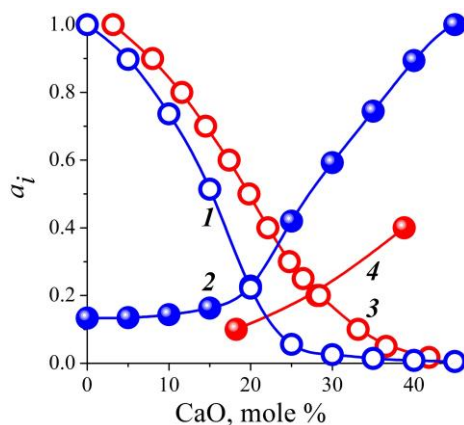


Fig. 1. The activities of  $\text{SiO}_2$  (1, 3) and  $\text{MgO}$  (2, 4) vs. the  $\text{CaO}$  content in the  $\text{CaO-MgO-SiO}_2$  system at  $1600^\circ\text{C}$  and  $\text{MgO} : \text{SiO}_2 = 2 : 3$  mole ratio according to the data: 1 and 2 – [3], 3 and 4 – [2].

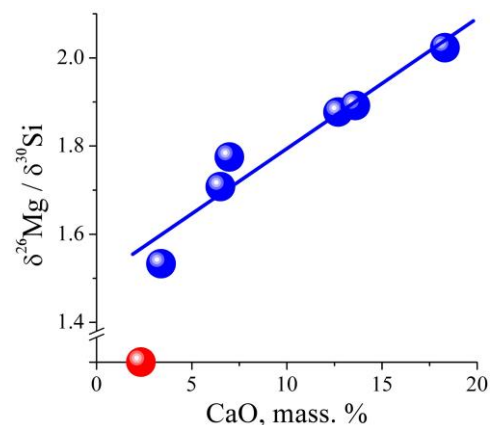


Fig. 2. The fractionation effectivity of the Mg and Si isotopes vs. the  $\text{CaO}$  content. The  $\text{CaO}$  initial content denoted the red symbol.

**References:** [1] Korginsky D. S. 1959. *Doklady AN SSSR* 128: 383–386. [2] Rein R. H. and Chipman J. 1965. *Trans. Met. Soc. AIME* 233: 415–425. [3] Shornikov S. I. 2008. *Experiment in Geosciences* 15: 147–149. [4] Mendybaev R. A. et al., 2013. *Geochim. Cosmochim. Acta* 123: 368–384.