

## H<sub>2</sub>O contents of Ca-rich plagioclase phenocrysts from arc volcanic front

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Ca-rich plagioclase is commonly found in island arc basalt at volcanic front. Experimental studies [1, 2, 3] have shown that composition of plagioclase becomes enriched in Ca with increasing H<sub>2</sub>O content in melt. Therefore, composition of plagioclase can be an indicator of H<sub>2</sub>O content of magma at the time of crystallization.

However, in Izu volcanic arc, H<sub>2</sub>O-rich nature in basaltic magmas has not proved yet. Analyses of basaltic melt inclusions hosted by Ca-rich plagioclase yield lower H<sub>2</sub>O content ( $\leq \sim 2$  wt.%) [3, 4]. Lower H<sub>2</sub>O content in melt inclusions may be explained by post-entrapment volatile leakage [3].

In order to confirm H<sub>2</sub>O content at the time of plagioclase crystallization, we analyzed trace quantities of H<sub>2</sub>O in Ca-rich plagioclase obtained from the 1986 eruption of Izu-Oshima volcano (34°N 44', 139°E 24'), using polarized infrared spectra [5]. Analytical results clearly demonstrate that H<sub>2</sub>O content becomes higher with increasing An content of plagioclase. Assuming partition coefficient of H<sub>2</sub>O between plagioclase and melt = 0.004 [6], H<sub>2</sub>O content in melt changes widely from 3 to  $\geq 6$  wt.%, which is consistent with melting experiments of hydrous basalt to crystallize Ca-rich plagioclase. Analyses of H<sub>2</sub>O in plagioclase phenocrysts suggest that H<sub>2</sub>O content of island arc basaltic magmas is higher than that of melt inclusions.

[1] Sisson & Grove (1993) *Contrib. Mineral. Petrol.* **113**, 143-166. [2] Takagi *et al.* (2005) *Contrib. Mineral. Petrol.* **149**, 527-540. [3] Hamada & Fujii (2007) *Geochem. J.* **41**, 437-461. [4] Saito *et al.* (2005) *Bull. Vol.* **67**, 268-280. [5] Johnson & Rossman (2003) *Am. Mineral.* **88**, 901-911. [6] Johnson (2005) *GCA* **69**, A743.

## Syn-eruptive magma crystallization

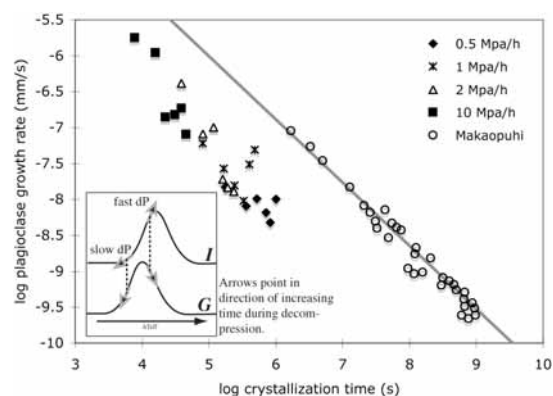
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The potential consequences of decompression-driven crystallization for modulating eruptive style are now widely recognized. The next step in interpreting natural crystal textures is to understand the crystallization response to decompression for a range of compositions and ascent rates. Experiments are underway on natural rhyodacite to understand how plagioclase nucleation rate (*I*), growth rate (*G*), crystal size distributions, morphologies, melt inclusions, and the effective undercooling ( $\Delta T_{\text{eff}}$ ) develop with time.

A family of experiments varying in quench pressure at each of four decompression rates (0.5, 1, 2, and 10 MPa h<sup>-1</sup>) reveals that *I* progressively increases with time by nearly 3 orders of magnitude at large  $dP/dt$ , yet decreases by a similar amount at low  $dP/dt$ . In contrast, *G* decreases with progressive decompression at all decompression rates. In combination with known functional dependence of *I* and *G* on  $\Delta T_{\text{eff}}$ , these observations suggest that slow ascending magmas maintain small  $\Delta T_{\text{eff}}$ , while magmas ascending  $\geq 0.06$  m s<sup>-1</sup> progressively lag behind equilibrium (Fig 1 inset).

*G* is a power law function of crystallization time (Fig.1), with time dependence similar to that of plagioclase in Makaopuhi basalt (Cashman 1993) and *G* values that are greater as a consequence of shorter crystallization times. The offset may reflect higher melt viscosity and nuclei density in the silicic magma.



[1] Cashman (1993) *Contrib. Mineral. Petrol.* **113**, 126-142.