Experimental study of plagioclase growth, nucleation rates, and shape evolution during cooling of an anhydrous basaltic andesite

1190 °C

1180 °C

1165 °

1140 °

1120 °(

(C)

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1. Introduction

Data on the nucleation and growth rates of plagioclase are underrepresented in the experimental dataset for basaltic andesites. Moreover, previous experiments were generally conducted on a **perfectly homogeneous glass** (without pre-existing crystals).

We aim at constraining the nucleation (J) and growth (G) rates of plagioclase in a natural basaltic andesite.

Experiments were performed at 1 atm, anhydrous and NNO conditions in a





24 hours ΔT = 10 °C Figure 1 : Multi-step cooling experiments quenched at a final

2. Experimental Method

	OS36	1190 °C
	Basaltic andesite	Run
SiO ₂	56,94	57,65
TiO ₂	1,30	1,28
Al ₂ O ₃	16,27	16,64
FeO	9,56	9,53
MnO	0,18	0,19
MgO	2,79	3,13
CaO	6,39	6,63
Na ₂ O	4,32	4,14
K ₂ O	0,91	0,89
P ₂ O ₅	0,26	0,3

3. Textural observations



Figure 2 : Plagioclase textural evolution. (a) 10 °C above the liquidus (end of the initial step; Fig. 1). (b-c) 15 °C and 60 °C below the liquidus

- \succ Close to the liquidus \rightarrow partial resorption of pre-existing plagioclase (Fig. 2a) favored nucleation around the Pt wire and bubbles (not shown here)
- Plagioclase texture: from equant (1165 °C) to tabular shape (1120 °C, Fig. 2b-c). \rightarrow High proportion of largest crystals (> 50 μ m) \rightarrow Tabular shape is dominant (Fig. 3)
- Large variation of 2D aspect ratio (from equant to tabular, Fig. 3)



- Starting composition : powdered basaltic andesite (Osorno volcano, Chile: Table 1) heated in a muffle furnace on platinum loops at 1000 °C to remove volatiles and a certain proportion of initial crystals.
- Experiments contained some pre-existing crystals of Plg (not homogeneous glass).
- Different cooling rates were used (Fig. 1). BSE images acquired for each experimental run were treated with GIMP and FIJI.
- 3D shape of plagioclase estimated using **ShapeCalc** spreadsheet (*Mangler et al. 2022*).



Table 1 : Composition of the starting material (OS36) analysed by XRF (Bechon et al., 2022) and by electron microprobe (experimental run held 10 °C above the liquidus for 24 hours). Platinum loops were pre-saturated in order to prevent FeO loss.

a)



Figure 3 : Evolution of the w/l 2D shape factor. w and l are the 2D minor and major axis of each crystal.



- Nucleation and growth rates are calculated by various methods (Imax, Batch, and CSD method; Equations).
- Strong variation between the different methods (Fig. 5a).
- Increase in growth rate with cooling rate (Fig. 5b) ➢ Great increase at 9 °C/h

Different cooling rates (Gmax values; Fig 5b)

➤ 1 °C/h : 2.7. 10⁻⁸ to 5.6. 10⁻⁸ cm/s ➤ 3 °C/h : 5.6. 10⁻⁸ to 7.8. 10⁻⁸ cm/s ➢ 9 °C/h : 1,1. 10⁻7 to 2.8. 10⁻7 cm/s

G values in agreement with other studies (cf. compilation of Moschini et al., 2023) : 10⁻⁹ cm/s (Gbatch) to 10⁻⁸-10⁻⁷ cm/s (Gmax)



b) Comparison between G values obtained by Imax, CSD, and Batch methods.

6. Nucleation rates



CSD : several crystals generations (with size variations)

Cooling history :



Figure 4 : CSDs obtained with the CSDcorrection plugin from Higgins (2000).

- a) CSDs evolution with increasing ΔT .
- b) CSDs at several cooling rates

➢ Flattened part : the largest

With increasing ΔT (Fig. 4a): Increase in the flattened part

• With increasing cooling rate (Fig. Increase in the initial density of

Nucleation rate increases with cooling rate (Fig. 7b)

Variable nucleation near the liquidus.

3 °C/h : high variability of the nucleation rate at **1165** °C (Δ T = 15°C).

Figure 6 \rightarrow : Variation of plagioclase nucleation rate with increasing ΔT a) Comparison between JCSD and JBatch methods. b) J_{Batch} at different cooling rates.



 1 °C/h (seeds) • 3 °C/h (seeds) 9 °C/h (seeds)

References

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