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T. Hulsebosch**Sulfur, chlorine, and fluorine degassing and atmospheric loading by the 1783–1784 AD Laki (Skaftár Fires) eruption in Iceland**

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Abstract The 1783–1784 Laki tholeiitic basalt fissure eruption in Iceland was one of the greatest atmospheric pollution events of the past 250 years, with widespread effects in the northern hemisphere. The degassing history and volatile budget of this event are determined by measurements of pre-eruption and residual contents of sulfur, chlorine, and fluorine in the products of all phases of the eruption. In fissure eruptions such as Laki, degassing occurs in two stages: by explosive activity or lava fountaining at the vents, and from the lava as it flows away from the vents. Using the measured sulfur concentrations in glass inclusions in phenocrysts and in groundmass glasses of quenched eruption products, we calculate that the total accumulative atmospheric mass loading of sulfur dioxide was 122 Mt over a period of 8 months. This volatile release is sufficient to have generated ~250 Mt of H₂SO₄ aerosols, an amount which agrees with an independent estimate of the Laki aerosol yield based on atmospheric turbidity measurements. Most of this volatile mass (~60 wt.%) was released during the first 1.5 months of activity. The measured chlorine and fluorine concentrations in the samples indicate that the atmospheric loading of hydrochloric acid and hydrofluoric acid was ~7.0 and 15.0 Mt, respectively. Furthermore, ~75% of the volatile mass dissolved by the Laki magma was released at the vents

and carried by eruption columns to altitudes between 6 and 13 km. The high degree of degassing at the vents is attributed to development of a separated two-phase flow in the upper magma conduit, and implies that high-discharge basaltic eruptions such as Laki are able to loft huge quantities of gas to altitudes where the resulting aerosols can reside for months or even 1–2 years. The atmospheric volatile contribution due to subsequent degassing of the Laki lava flow is only 18 wt.% of the total dissolved in the magma, and these emissions were confined to the lowest regions of the troposphere and therefore important only over Iceland. This study indicates that determination of the amount of sulfur degassed from the Laki magma batch by measurements of sulfur in the volcanic products (the petrologic method) yields a result which is sufficient to account for the mass of aerosols estimated by other methods.

Key words Laki · Effusive basaltic volcanism · Fissure eruption · Eruption dynamics · Magma degassing · Volcanic gases · Volcanic glass chemistry · Volcanic aerosols

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Introduction

The growing interest over the past 20 years in the effects of volcanic eruptions on global climate has concentrated almost exclusively on large silicic explosive eruptions such as Tambora 1815, Krakatau 1883, and Pinatubo 1991 (e.g., Self et al. 1981; McCormick et al. 1995). However, large basaltic lava-producing eruptions, such as the Laki (Skaftár Fires) event in Iceland, 64°10'N, 17°15'W, in 1783–1784, are also known to have been linked to locally cooler weather and unusual atmospheric phenomena (Thorarinsson 1979, 1981; Sigurdsson 1982; Angell and Korshover 1985; Wood 1992; Stothers 1996; Thordarson 1995). Such eruptions have the potential to release great amounts of gas into the atmosphere, but studies have not been carried out on the details of the volatile emissions in terms of effi-

ciency of magma degassing, mechanisms and rates during eruption, heights of atmospheric volatile release, and accurate estimates of the masses of volatile species released.

Previous estimates of the amount of SO_2 released by the Laki eruption (Sigurdsson 1982; Óskarsson et al. 1984; Devine et al. 1984) range from 56 to 63 Mt when corrected for the new volume estimates for the Laki eruption products (Thordarson and Self 1993). Metrich et al. (1991) showed that three glass inclusions of Laki composition trapped in phenocrysts within phreatomagmatic clasts of the Laki tephra contained on average 1810 ± 100 ppm of sulfur, a considerably higher amount than the values of 800–960 ppm S used by Sigurdsson (1982), Óskarsson et al. (1984), and Devine et al. (1984). Although the phreatomagmatic tephra constitutes only approximately 0.5% of the total amount of magma erupted, the analyses of Metrich et al. (1991) are in good agreement with the results presented in this study (see below).

In this paper we address the degassing history and the volatile budget of the Laki eruption as revealed by the concentrations of S, Cl, and F in the products of all phases of the eruption. We present new major element and volatile analyses of 17 glass inclusions from Laki phenocrysts and of groundmass glass in quenched products. These data establish new and more accurate estimates of the atmospheric mass loading of gases by Laki and show that the volatile release by the eruption was considerably greater than has been estimated previously. The new fluorine and chlorine data indicate that Laki released large quantities of HF and HCl into the atmosphere, a previously unquantified, but not unexpected, occurrence during the Laki eruption (Thorarinnsson 1979, 1981). We also evaluate how degassing processes relate to the eruption mechanism and the rate of magma degassing during eruption.

The 1783–1784 Laki eruption

Nature of the eruption

The Laki fissure eruption lasted for 8 months, from 8 June 1783 to 7 February 1784, and produced one of the largest recorded basaltic lava flows, $14.7 \pm 1 \text{ km}^3$ (Fig. 1). In addition, it produced 0.4 km^3 (dense rock equivalent) of tephra, giving the total volume of 15.1 km^3 used in this paper. Over 60% of the volume was erupted during the first 1.5 months of activity, and

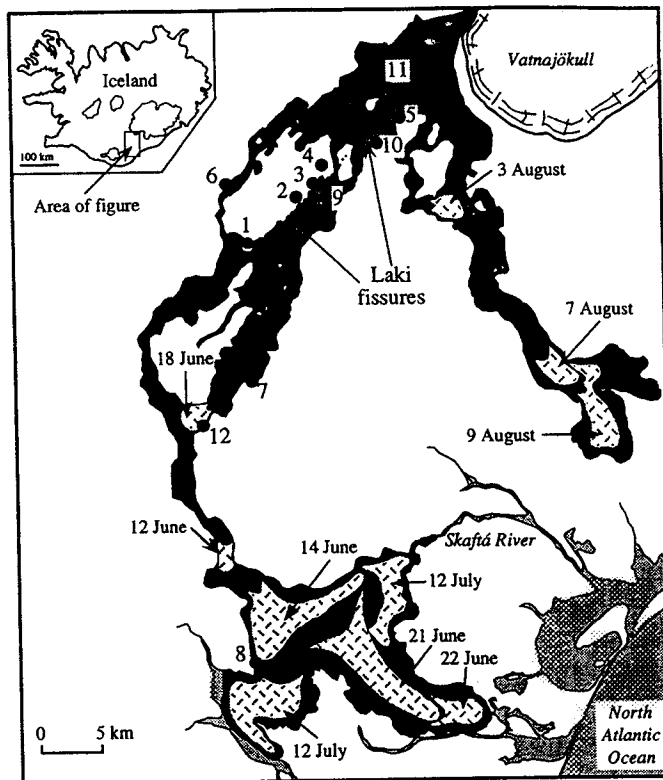


Fig. 1 Map of the Laki fissures and lava flow field, Iceland, showing locations (numbered 1–12) where the samples used in this study were collected (see text for discussion). Stippled areas denote locations of flow fronts on date shown and are reconstructed according to descriptions given by Reverend Jón Steingrímsson (1783, 1788)

90% by the end of the fifth month (Fig. 2). Ten eruption episodes occurred during the first 5 months of activity. Each began with a seismic swarm of increasing intensity and formation of a new fissure segment, followed by a short-lived explosive phase leading into lava

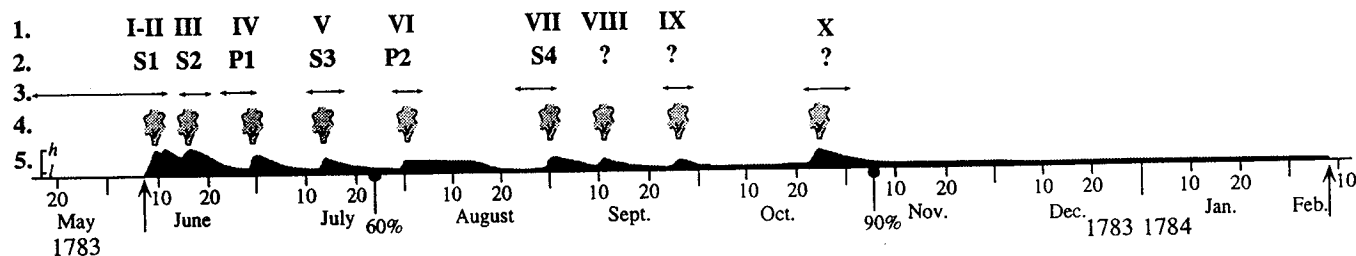


Fig. 2 Illustration of sequence of events during the Laki eruption up to eruption episode X. Eruption episodes are shown in row 1, labeled I–X; S, strombolian fall units; P, phreatomagmatic fall units. Row 2 shows the primary fall units produced by the eruption episodes. Row 3 designates earthquake swarms; the horizontal bars show the duration of each swarm. Timing of explosive activity is shown in row 4, where eruption clouds denote explosive activity at Laki fissures. Row 5 (stippled area) shows qualitatively the fluctuations in lava production during the eruption; *h* and *l* indicate high and low discharge, respectively (not to scale). Vertical arrows mark the beginning and end of Laki fissure eruption and the 60% and 90% refer to volume fraction erupted up to that time. (Modified from Thordarson and Self 1993)

Table 1 Laki sample locations and sample descriptions

Sample location	Label	Description	Unit ^a
Vent tephra			
Location 1; Hnúta, section 133 ^b	LS1-01	Strombolian scoria lapilli	S1
Location 12, Leidólfssfell	LS1-04	Strombolian ash	S1
Location 2, Vikradalur, section 94 ^b	LS2-06	Strombolian scoria lapilli	S2
Location 3, Hverfjall, section TC-1	LP1-U2	Phreatomagmatic tephra	P1
	LP1-U3	Phreatomagmatic tephra	P1
	LP1-U5	Phreatomagmatic tephra	P1
	LP1-U5	Phreatomagmatic tephra	P1
Location 4, Lambavatn, section 153 ^b	LS3-02	Strombolian scoria lapilli, base of unit	S3
	LS3-03	Strombolian scoria lapilli, top of unit	S3
	LP2-05	Phreatomagmatic tephra	P2
Location 5, Innrieyrar, section 75 ^b	LS4-01	Strombolian scoria lapilli	S4
	LS4-01	Strombolian scoria lapilli	S4
Lava flow			
Location 6, Kambar	L-20	Surface lava selvage	E1-FU4
	L-21	Basal lava selvage	E2-FU4
	L-23	Basal lava selvage	KAFU
Location 7, Kálfá	L-23	Basal lava selvage	KAFU
Location 12, Leidólfssfell	LEI-04	Lava core	LEIFU
	LEI-12	Lava crust	LEIFU
Location 8, Eldvatnsbrú	L-24	Basal lava selvage	SAL-9
	L-25	Surface lava selvage	SAL-9
Rootless cone tephra			
Location 9, Varmárdalur	VRC-9	Scoria lapilli	VRCTE
	VRC-11	Scoria lapilli	VRCTE
	BRC-9	Scoria lapilli	BRCTE
Location 10, Blängur	BRC-9	Scoria lapilli	BRCTE
Location 11, North of Innrieyrar	NIRC-7	Scoria lapilli	NIRCTE
Location 12, Leidólfssfell	LEIRC-03	Scoria lapilli	LEIRCU2
	LEIRC-03A	Scoria lapilli	LEIRCU2
	LEIRC-04	Ash	LEIRCU1
	LEIRC-12	Scoria lapilli	LEIRCU2
	LEIRC-20A	Ash	LEIRCU1
	LEIRC-20A	Ash	LEIRCU1

^a Primary tephra fall units are designated as in Thordarson and Self (1993); S and P denote strombolian and phreatomagmatic fall units, respectively. Lava flow units are described in Appendix I.

Unit designation for rootless cone tephra are based on unpublished tephra sections measured by Th. Thordarson

^b Section designation from Thordarson (1990)

fountaining and effusive activity characterized by voluminous lava surges. The magma volume erupted by each episode ranged from ~ 0.5 – 2.0 km^3 and the volume ratio of tephra to lava was $\sim 1/35$. Magma discharge varied considerably during the eruption with highest eruption rates in the range of 2 – $5 \times 10^3 \text{ m}^3 \text{ s}^{-1}$ occurring during each eruptive episode (Fig. 2). During the first 5 months of activity, the mean discharge was of the order of 1 – $2.5 \times 10^3 \text{ m}^3 \text{ s}^{-1}$. The episodic nature of the eruption cannot be attributed to separate inflation–deflation events within the Laki reservoir because of the high magma fluxes and the short time which elapsed between successive episodes. Evaluation of the timing and sequence of events suggest that unsteady or non-uniform conduit flow was responsible for the episodic behavior of the eruption (Thordarson and Self 1993).

Each explosive phase lasted for hours to a few days and was characterized by either magmatic or hydro-magmatic activity, producing a tephra deposit which consists of at least six mappable strombolian and phreatomagmatic fall units. The strombolian fall units (S1–S4; Table 1) account for $\sim 80\%$ of the total tephra volume, whereas the phreatomagmatic fall units (P1 and P2; Table 1) make up the remaining $\sim 20\%$. The Laki fire fountains extended to heights of 800–1450 m during periods of peak magma discharge (Thordarson 1990).

Historic evidence and calculations indicate that the Laki eruption column reached maximum altitudes of 11–13 km at times of vigorous activity, and that the eruption maintained 7- to 9-km-high columns during the first 5 months (Thordarson and Self 1993; Woods 1993). The tropopause over this region is at 9–10 km. As is demonstrated below, vent degassing, especially during periods of vigorous explosive activity and fountaining, is responsible for release of the bulk ($\sim 80\%$) of the volatiles during the Laki eruption, whereas degassing of lava during transport and cooling accounts for the remaining 20%.

Whole-rock compositions

The Laki products are glomeroporphyritic quartz-tholeiite basalt containing $\sim 5.5 \text{ vol.}\%$ of euhedral phenocrysts ranging in size from 0.2 to 3.0 mm. The glomerocrysts contain one, two, or more rarely, all of the following crystal phases: plagioclase (An_{90-55}), olivine (Fo_{86-68}) and clinopyroxene ($\text{Wo}_{40}\text{En}_{45}\text{Fs}_{14.5}$), and are in compositional equilibrium with the erupted magma (Grönvold 1984; Metrich et al. 1991). Laki tephra and lava (Fig. 3) show a small spatial and temporal compositional variance in major element concentrations (Bell and Humphries 1972; Grönvold 1984). Uniform U and

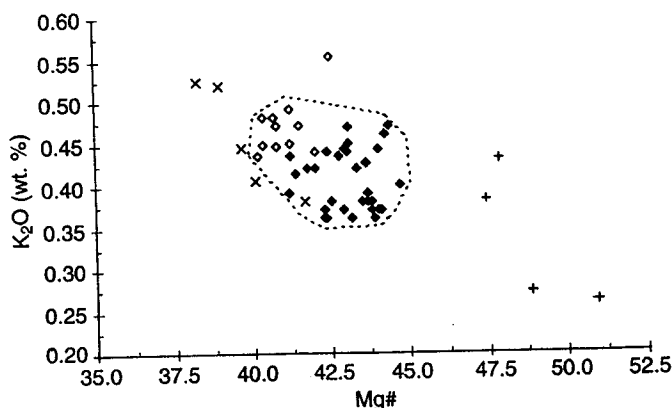


Fig. 3 Compositional variations of eruption products within the Grímsvötn volcanic system (GVS). K_2O (wt.%) variation as a function of Mg# (calculated as $[Mg/(Mg+Fe)] \times 100$). Laki samples are indicated by *open* (tephra) and *filled* (lava) diamonds. *Plusses* represent other subaerial lavas within the GVS and *crosses* denote tephra samples from the subglacial Grímsvötn central volcano. *Broken line* outlines the field defined by the Laki eruption products. Major element data are from this study and the sources given in Thordarson (1995)

Th abundances (0.344 ± 0.007 ppm and 1.12 ± 0.02 ppm, respectively) demonstrate the homogeneity of the Laki magma; differences in measured concentrations in samples from all main eruptive units are $<7\%$ (Sigmarsson et al. 1991). The apparent uniformity of the Laki magma is remarkable in view of the duration of the eruption and the large volume of magma involved.

Sample collection and analytical methods

Samples and their significance

Samples representing the main phases of the Laki eruption were collected at 12 different locations within the lava flow and along the fissures (Fig. 1; Table 1). The types of samples collected for analyses are shown schematically in Fig. 4.

Tephra samples, representing mappable fall units, were collected at locations 1–5. A complete proximal section through the Laki flow is not available within 10 km from source, whereas in the medial (10–30 km from source) and distal (>30 km from source) areas the best outcrops occur in river channels (Fig. 1). Locations 6, 7, 8, and 12 represent measured sections through the Laki lava flow where samples of the lava interior and basal glass selvage were collected (Fig. 4; Table 1). Samples were also collected from the rootless cone groups at locations 9, 10, 11, and 12. These samples together represent flow units emplaced in June and July 1783 (locations 7, 8, 9, and 12), and from August through October 1783 (locations 6, 10, and 11). The significance of the samples is discussed further in Appendix I.

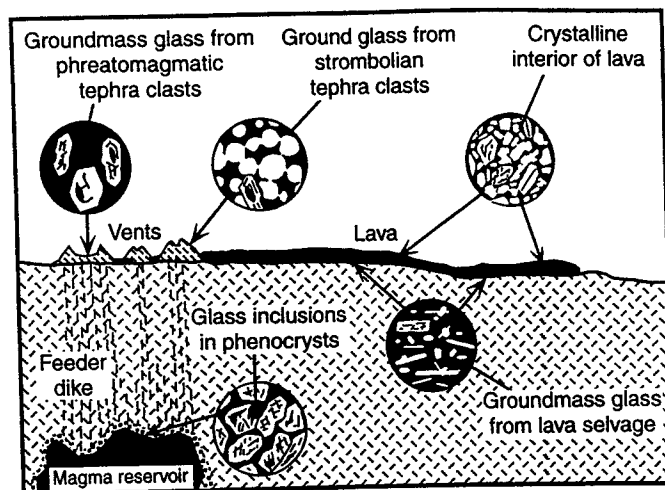


Fig. 4 Illustration (not to scale) of the Laki eruption viewed from the southwest showing the type of samples (arrows) collected for this study. *Glass inclusions in phenocrysts* represent the undegassed magma fraction residing in a reservoir at depth prior to eruption. *Groundmass glass from phreatomagmatic tephra clasts* represents partly degassed magma which froze in the feeder dike due to interaction with external water. *Groundmass glass of strombolian tephra clasts* represents magma which degassed at the vents. *Lava selvages* characterize magma which has gone through the stage of vent degassing and, in addition, has liberated a fraction of its remaining volatiles as the lava flowed away from the fissures. *Crystalline lava* denotes magma which has gone through the stages of vent and lava emplacement degassing and has lost an additional fraction of its volatiles during cooling and crystallization. It also contains the volatiles which did not escape from the lava into the atmosphere

Analytical procedure

Modes and vesicularity of representative tephra and lava samples were estimated by point counting of ~ 1000 points per thin section. Microprobe analyses used instrumental settings of 15 kV for the accelerating voltage, 15–20 nA in a beam-current-regulated mode, and a 5–20 μm focused beam in major element analyses of glass. Precision (2σ) for major elements is better than 1%. Analyses of S and Cl were conducted by the Cameca Trace routine, using the same general setup and counting time of 200 s. Based on counting statistics the precision for sulfur is 80 ppm and for Cl it is 60 ppm at the 2σ level. The same samples were also analyzed for S, Cl, and F by using the CSIRO-trace routine of Robinson and Graham (1992) with 15 kV accelerating voltage, 80 nA beam current, 10 μm focused beam, and counting time of 400 s. Precision (2σ) for S=35 ppm, Cl=30 ppm, and F=90 ppm. Raw data were corrected using the standard ZAF procedures. The Juan de Fuca glass standard VG2 and Makoupuhi glass standard A99 were analyzed as unknowns for S, Cl, and F during each analytical session and, as shown in Table 2a, our S values for both standards are in good agreement with Dixon et al. (1991). Precision and accuracy of the Cl and F microprobe analyses was tested by analyzing the MORB glass samples TR 154 21D-3 and TR 138 6D-1

Table 2a Representative microprobe analyses of international standards by Cameca and CSIRO trace routines

	VG-2	S	Cl	F	N	A-99	S	Cl	F	N
Cameca trace routine	Reported	1340 ^a	n.a.	n.a.	19		170 ^a	n.a.	n.a.	15
	SD	80	—	—			30	—	—	
	This study	1348	291	n.a.	134		135	229	n.a.	37
	SD	62	52	—			50	49	—	
CSIRO trace routine	This study ^b	1365	316	300	139		220	227	765	42
	SD	29	19	72			12	20	79	

SD, standard deviation; n.a., not analysed; N, number of analyses

^a Value reported by Dixon et al. (1991)^b Represents an average of analyses conducted over a period of 2 years by Th. Thordarson, S. B. Sherman, and L. Norby**Table 2b** New CSIRO trace routine analyses of basalt glass samples from Michael and Schilling (1989)

	TR 154 21D-3	S	Cl	F	N	TR 138 6D-1	S	Cl	F	N
CSIRO trace routine	Reported	n.a.	404 ^a	520 ^a			n.a.	44 ^a	280 ^a	
	This study ^b	977	488	546	11		1050	48	282	19
	SD	26	31	65			24	12	82	

SD, standard deviation; n.a., not analysed; N, number of analyses

^a Values reported by Michael and Schilling (1989)^b Analyst: L. Norby

from Michael and Schilling (1989), and our results are compatible with theirs (Table 2b).

The H₂O and CO₂ values used in this paper are from Óskarsson et al. (1984), which were obtained by Quadropol mass spectrometry on gasses released by vacuum fusion of crystalline lava, fibrous tephra, and crushed plagioclase phenocrysts rich in glass inclusions. The measured CO₂ values for glass inclusions may overestimate the concentration initially dissolved in the Laki magma, because it is likely that small CO₂ bubbles were present as a free volatile phase in Laki phenocrysts (Metrich et al. 1991). The method may also overestimate the CO₂ concentration dissolved in the groundmass glass of tephra, because the fibrous tephra clasts contain rare liquid-filled microbubbles (Ólafsson et al. 1984). Despite these uncertainties, we chose to use the CO₂ values of Óskarsson et al. (1984), because it is the only comprehensive data set which includes measurements of CO₂ inclusions, tephra, and crystalline lava.

Petrology and glass chemistry

In this section we outline the major element and volatile chemistry of glass inclusions and groundmass glass in quenched products of the Laki eruption. The chemistry of single samples is listed in Appendix II.

Glass inclusion

All of the Laki phenocryst phases contain clear, light-brown glass inclusions and sparse inclusions of dark-brown, transparent glass, ranging from ≤ 1 to 500 μm

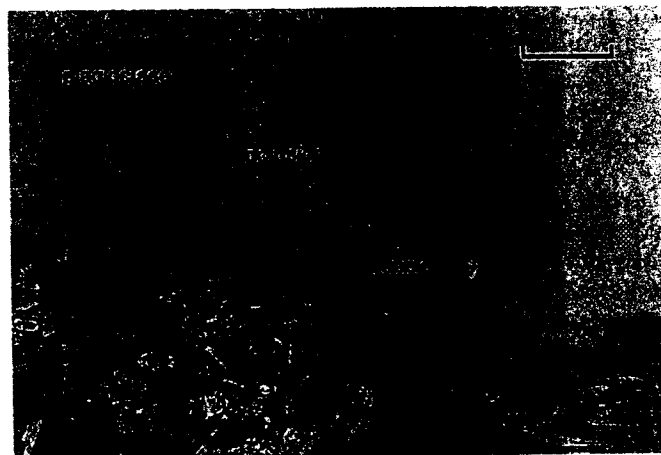


Fig. 5 Transmitted-light photomicrograph of a 60-×25- μm light-brown glass inclusion in plagioclase (sample L-21). The dark circular spot in the lower right corner is a bubble. Scale bar is 100 μm

in the longest dimensions (Fig. 5). Common inclusions are between 20 and 120 μm in size and of spherical or oblate shape (Thordarson 1995). Some inclusions contain vapor bubbles.

Major element microprobe analyses of more than 30 glass inclusions from Laki phenocrysts were performed to locate inclusions of composition similar to the bulk Laki magma. Several inclusions showed clear evidence of extensive postentrapment host crystallization (Table 3) and were therefore not considered further. Measured major element concentrations of inclusions L-i-7, L-i-8, and L-i-11 trapped in olivine phenocrysts (Table A-1, Appendix II), demonstrate that the glass composition has been slightly affected by host crystalliza-

Table 3 Major and volatile element composition of Laki glass inclusions

	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	N
Group 1	49.68	2.96	13.05	13.78	0.22	5.78	10.45	2.84	0.42	0.28	99.45	1677	310	665	42.77	21/23
SD	0.36	0.17	0.48	0.61	0.03	0.30	0.28	0.12	0.05	0.05	0.34	225	59	101		
Group 2	49.27	3.03	12.08	15.33	0.23	5.96	10.41	2.43	0.48	0.31	99.77	1632	246	537	40.92	6/6
SD	0.70	0.06	0.37	0.59	0.01	0.44	0.18	0.03	0.09	0.04	0.15	171	56	90		
Group 3	48.87	3.43	13.08	16.31	0.22	4.70	9.56	2.57	0.45	0.35	99.77	1622	298	660	33.92	6/6
SD	0.01	0.04	0.11	0.07	0.04	0.11	0.08	0.02	0.04	0.03	0.37	493	18	71		
Group 4	49.13	2.00	12.03	13.41	0.23	8.47	11.37	2.47	0.39	0.17	99.67	1675	198	562	50.44	4/5
SD	0.04	0.02	0.01	0.11	0.03	0.14	0.02	0.05	0.01	0.01	0.06	50	45	70		
Unaffected by host crystallization																
L-i-3	49.61	2.85	13.07	13.88	0.24	5.77	10.39	3.01	0.42	0.25	99.50	1520	300	705	42.56	5
SD	0.16	0.03	0.07	0.15	0.03	0.15	0.09	0.05	0.02	0.04	0.40	25	55	10		
Affected by host crystallization ^a																
L-i-18	48.42	2.67	7.37	16.48	0.22	10.44	11.30	1.80	0.31	0.23	99.25	1555	260	460	53.03	2
SD	0.01	0.12	0.63	0.08	0.04	0.30	0.08	0.30	0.07	0.01	0.65	90	85	130		
Laki whole-rock composition																
L-lava	49.63	2.68	13.87	13.45	0.22	5.73	10.40	2.69	0.41	0.33	99.28				43.13	32
SD	0.62	0.13	0.31	0.34	0.01	0.22	0.20	0.15	0.04	0.06	1.03				0.97	

Mg# = 100 Mg/(Mg + Fe*), Fe* = total Fe; N, number of analyses – 1/1 = major/volatile element analysis; SD, standard deviation

^a Host is plagioclase

tion. These inclusions were restored to their original composition using the method of Metrich et al. (1991).

The Laki inclusion data are ordered into four groups on the basis of major element abundances (Table 3). Group 1 glass inclusions exhibited major element abundances (Mg# = 42.77 ± 1.37) that closely match the averaged bulk composition of the Laki magma (Mg# = 43.34 ± 0.66) and are considered to be representative of the pre-eruption magma composition. For most inclusions, S concentrations range from 1500 to 1700 ppm (mean 1677 ± 225 ppm), chlorine 250–350 ppm (mean 310 ± 59 ppm), and fluorine 600–800 ppm (mean 665 ± 100 ppm). Group 2 inclusions demonstrate higher Fe and lower Ca and Na contents than those of group 1. However, their mean abundances of S (1632 ± 171 ppm), Cl (246 ± 56 ppm), and F (537 ± 90 ppm) are strikingly similar. The difference in the major element composition cannot be accounted for entirely by host crystallization after entrapment. Group 3 inclusions are strongly enriched in Fe and Ti, depleted in Mg, Ca, and Na, but with volatile concentrations in the same range as the two former groups. Inclusions of groups 2 and 3 may represent liquids which have evolved by fractionation in the Laki reservoir. Group 4 inclusions have an olivine tholeiite composition and may represent a parental liquid to the Laki magma.

The S, Cl, and F values of Laki inclusions have some degree of variance (Fig. 6), which cannot be attributed to analytical error alone. This variance could be caused by several factors: (a) compositional heterogeneity in the magma reservoir; (b) a fraction of the volatiles may have separated from the liquid phase into a bubble within the inclusions as the system was decompressed;

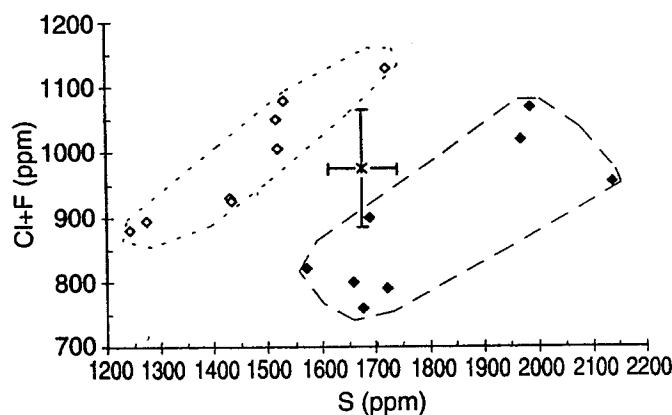


Fig. 6 Halogen-sulfur covariation diagram for Laki glass inclusions. Inclusions with low (Cl+F)/S values (filled diamonds) are outlined by the heavy broken line, whereas the light broken line outlines inclusions with high (Cl+F)/S values (open diamonds). The average inclusion value is shown by the asterisk and the bars indicate the estimated analytical error (2σ)

and (c) S-, Cl-, and/or F-bearing mineral phases may have formed in some inclusions after entrapment, although the major element concentrations and petrographic examination reveal no evidence of such minerals. Despite the relatively random distribution of data points in Fig. 6, the data appear to form two arrays, one with low (0.45–0.54) and another with high (0.64–0.71) halogen-to-sulfur ratios. Vapor bubbles are not present in inclusions with low (Cl+F)/S values, whereas many, but not all, of the inclusions with high (Cl+F)/S values contain vapor bubbles. The apparent association of vapor bubbles and low S values in some Laki inclusions may suggest that S (along with CO₂ and H₂O?) was exsolved from the inclusion melt to form bubbles contain-

Table 4 Composition of groundmass glass in Laki tephra clasts

Fall Unit	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Ves.%	Cr.%	N
Strombolian tephra																		
S1	50.06	3.06	13.22	14.31	0.21	5.43	9.87	2.80	0.45	0.32	99.74	453	205	480	40.35	59	2	29/23
SD	0.19	0.06	0.09	0.23	0.04	0.07	0.15	0.09	0.02	0.05	0.23	78	48	85	0.50	19	1	
S2	49.95	3.05	13.29	14.34	0.25	5.38	10.01	2.81	0.48	0.35	99.71	513	148	480	40.05	45	4	6/5
SD	0.16	0.06	0.11	0.33	0.02	0.08	0.16	0.05	0.02	0.01	0.41	21	11	53	0.83	11	1	
S3	49.80	3.01	13.19	14.03	0.21	5.41	9.87	2.72	0.45	0.30	98.98	515	256	472	40.51	77	1	30/36
SD	0.34	0.04	0.11	0.25	0.03	0.05	0.10	0.15	0.02	0.04	0.44	88	77	99	0.41	10	0	
S4	50.07	3.14	13.08	14.42	0.24	5.21	10.10	2.83	0.49	0.36	99.93	473	149	491	39.16	27	4	6/6
SD	0.11	0.04	0.06	0.15	0.03	0.04	0.05	0.05	0.02	0.01	0.26	26	20	64	0.12	2	2	
S1-S4																		
Average	49.94	3.05	13.20	14.20	0.22	5.40	9.90	2.77	0.45	0.32	99.43	490	223	476	40.39	64	2	71/70
SD	0.28	0.06	0.11	0.28	0.03	0.09	0.15	0.12	0.03	0.05	0.52	82	74	89	0.64	20	1	
Phreatomagmatic tephra																		
P1	49.66	2.84	13.34	13.80	0.22	5.74	10.71	2.40	0.56	0.26	99.53	937	218	421	42.56	33	4	32/54
SD	0.25	0.14	0.14	0.35	0.03	0.26	0.55	0.21	0.10	0.04	0.38	217	79	106	1.57	31	4	
P2	49.84	2.78	13.48	13.60	0.22	5.70	10.68	2.77	0.42	0.31	99.81	903	133	418	42.74	36	1	9/7
SD	0.27	0.30	0.20	0.78	0.03	0.57	0.76	0.15	0.07	0.05	0.33	176	24	64	3.77	35	1	
P1-P2																		
Average	49.70	2.83	13.37	13.75*	0.22	5.73	10.70	2.48	0.53	0.27	99.59	933	208	420	42.60	34	3	41/61
SD	0.26	0.18	0.16	0.47	0.03	0.34	0.59	0.25	0.11	0.04	0.38	212	79	102	2.18	31	4	

Individual fall units are designated as in Thordarson and Self (1993). S, strombolian tephra units; P, phreatomagmatic tephra units; Mg# = 100 Mg/(Mg + Fe*), Fe* = total Fe; Ves% and

Cr.%, vesicle and crystal content of sample in vol. %, respectively, N, number of analyses - 1/1 = major/volatile element analyses; SD, standard deviation (2 σ)

ing a free volatile phase as the magma rose to the surface, although more work is necessary to confirm this relationship. Consequently, the mean S concentration of group 1 inclusions (Table 3) is likely to be a minimum estimate for dissolved sulfur in the Laki magma prior to eruption. The variance in volatile concentrations demonstrates clearly the importance of analyzing multiple inclusions to obtain a reasonable estimate of pre-eruption volatile concentrations.

Vent tephra

The Laki fall deposits consist of holohyaline tephra clasts with sparse phenocrysts (1–10 vol.%) of olivine, plagioclase, and clinopyroxene. Although the analyzed tephra represents fall units produced by episodic explosive activity over a period of 3 months (Fig. 2), the major element composition of their groundmass glass show no systematic variations with stratigraphic position (Table 4). However, the major element data indicate a slight compositional difference between groundmass glass of the two types of tephra produced by the eruption, identified as two partly overlapping fields on bivariate plots (Fig. 7). The strombolian tephra has higher FeO and TiO₂, but lower MgO than the bulk magma, and is slightly more evolved. The dense clustering of data points confirms the uniform composition of all strombolian fall units. The groundmass glass of the phreatomagmatic clasts shows more variable major element concentrations, forming an array spanning the

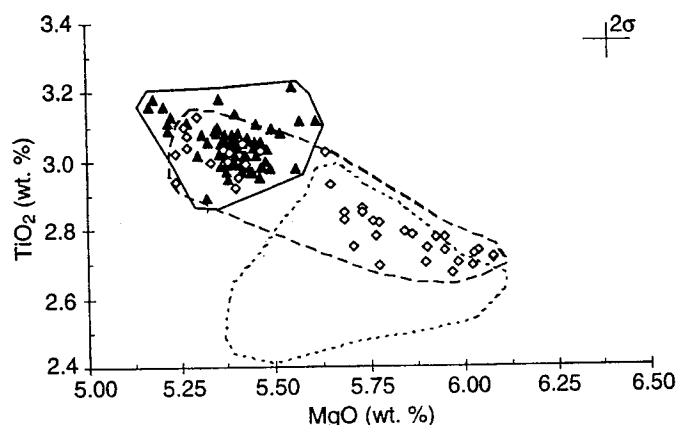


Fig. 7 Compositional variations in Laki tephra samples. TiO₂ (wt.%) as a function of MgO (wt.%). Note the strong clustering exhibited by the strombolian tephra (filled triangles) and the rough linear trend defined by the phreatomagmatic tephra (open diamonds). Solid and broken lines outline the compositional fields of strombolian and phreatomagmatic tephra, respectively. Dotted line defines the whole-rock compositional field of Laki eruption products. Estimated analytical error (2 σ) is shown by the cross in the upper right corner

compositional range of the bulk magma and the strombolian tephra. Nevertheless, a majority of the clasts resemble the bulk magma composition (Fig. 7).

The variable, but high, S concentration in groundmass glass of phreatomagmatic clasts (mean 933 ± 212 ppm) contrasts with very uniform S abundances in the strombolian clasts (mean 490 ± 82 ppm).

The S values of the phreatomagmatic tephra range from approximately 1260 to 490 ppm, but the corresponding compositional range for the strombolian tephra is 418–640 ppm S (Table A-1, Appendix II). The groundmass glass of both tephra types have variable, but similar, Cl and F concentrations. However, the compositional range of Cl and F is greater in phreatomagmatic tephra (Cl=122–450 ppm; F=220–560 ppm) than in the strombolian tephra (Cl=134–320 ppm; F=310–530 ppm).

The compositional differences and similarities between the two tephra types are of interest because the tephra represent different styles of explosive activity. The strombolian tephra deposits formed as the melt was disintegrated by rapidly expanding magmatic gases, whereas the phreatomagmatic tephra was produced by interaction of external water and magma (Thordarson and Self 1993). The strombolian tephra units are composed of well-sorted scoria lapilli, consisting primarily of highly vesicular clasts (60–90 vol.% voids) featuring a single population of 2- to 10-mm spherical vesicles. The strombolian clasts are either partly or wholly coated by a thin fluidal skin (<0.01 mm), suggesting that the outer surfaces of the clasts were fused (Fig. 8a and b), presumably by a stream of hot gas. The strombolian tephra also contains reticulite and Pele's hair. The phreatomagmatic tephra units are composed of poorly sorted ash to scoria lapilli consisting of non-vesicular to highly vesicular angular clasts (Fig. 8c) in approximately equal proportions. The phreatomagmatic clasts lack the glassy fluidal skin that characterizes the strombolian tephra clasts. The highly vesicular clasts in both tephra types represent true magmatic foam, and the reticulite-like strombolian clasts indicate that a small fraction of the melt had evolved to mature polyhedral foam (Cashman and Mangan 1994).

Major element abundances in both tephra types show poor correlation with clast vesicularity, although higher vesicle abundance (>45 vol.%) appears to accompany a more evolved (iron-rich) composition (Fig. 9a). S abundances in the groundmass glass of phreatomagmatic clasts show a weak negative correlation with degree of vesicularity, whereas the S concentrations in strombolian clasts are independent of vesicle abundance (Fig. 9b). Furthermore, the fluorine and chlorine concentrations show no obvious correlation with vesicularity (Fig. 9c). Considering the nature and timing of eruptive events and the relatively small tephra volume, the more evolved tephra compositions suggest that a portion of the magma fractionated chemically during its rise to the surface.

Lava selvages and rootless-cone tephra

The basal-quenched lava selvages and the rootless-cone tephra exhibit hyalophitic (intersertal) texture, in which glomerocrysts (5–6 vol.%) and the seriate-textured groundmass mineral assemblage of plagioclase, olivine,

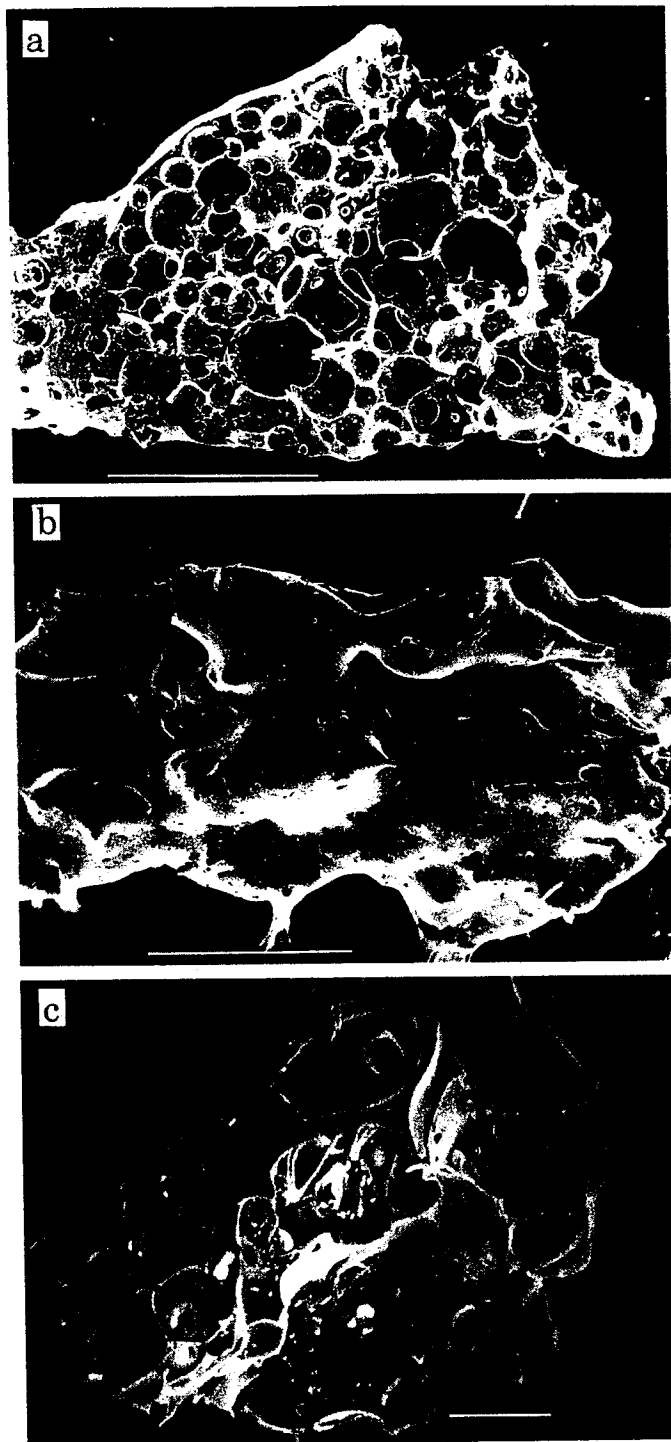


Fig. 8a–c Scanning electron photomicrographs of highly vesicular (foam-like) tephra particles from Laki. **a** Interior of a clast from the strombolian tephra deposit. Scale bar is 1000 μm (white line). **b** Strombolian clast completely covered by a very thin fluidal skin. The interior of the clast is highly vesicular (~80 vol.% voids). Scale bar is 1000 μm . **c** Clast from the phreatomagmatic tephra deposit. Scale bar is 1000 μm .

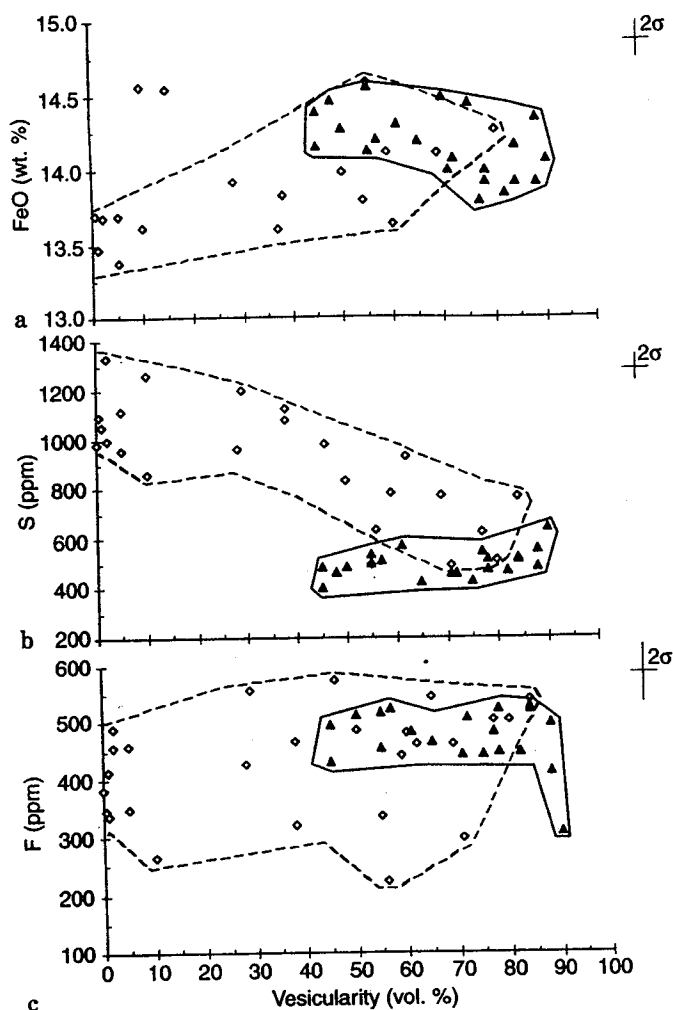


Fig. 9a-c Variation in volatile abundances as function of vesicularity in Laki tephra samples. **a** FeO vs vesicularity. **b** S vs vesicularity. **c** F vs vesicularity. Phreatomagmatic tephra is indicated by open diamonds, and strombolian tephra by filled triangles. Estimated analytical error (2σ) is shown by the cross in the upper right corner

and clinopyroxene (~ 10 – 40 vol.%) are completely embedded in light- to dark-brown sideromelane glass (Fig. 10). Fe–Ti oxides are not present in the groundmass mineral assemblages of lava selvages and rootless-cone tephra.

The groundmass glass of the lava selvages and the rootless-cone tephra clasts shows a more evolved composition than the bulk lava and tephra (Table 5a). Groundmass glass of proximal lava samples shows the smallest variation from the bulk lava composition, with an Mg# of 35.4 ± 2.2 . The groundmass glass of the medial and distal samples, however, shows a considerable degree of differentiation with strong enrichment in TiO_2 and FeO, and Mg# between 32.7 ± 1.8 and 34.0 ± 0.2 (Fig. 11a; Table 5a). The S, Cl, and F concentrations of the groundmass glass in the proximal lava are in the same range as that of the strombolian tephra (Tables 4 and 5a). Furthermore, the measured abundances of S (334 ± 29 to 489 ± 5 ppm), Cl (188 ± 49 to



Fig. 10 Transmitted-light photomicrograph of Laki lava selva (sample L-21) demonstrating the holohyaline nature of the groundmass and the seriate texture of the mineral assemblage. Note also that no Fe–Ti oxide mineral phases are present. Scale bar is $500 \mu\text{m}$

309 ± 98 ppm), and F (495 ± 109 to 613 ± 93) in groundmass glass of lava selvages and rootless-cone tephra from the medial and distal locations (Fig. 11b) are surprisingly uniform and on average similar or only slightly lower than those found in the proximal lava and strombolian tephra.

The crystallinity of lava selvages and rootless-cone tephra ranges from 12 to 50 vol.%, with the least crystalline samples being closest to source (Table AII-1, Appendix II). The crystal content of the lava glass selvages falls between 25 and 38 vol.% and, in general, increases with distance from the source (Fig. 11c). The rootless-cone tephra at Leidólfssell shows a greater range of crystallinity (30–50 vol.%). Despite the evolved groundmass glass composition, the bulk (whole-rock) composition of the rootless-cone tephra and lava (Table 5b) is essentially the same as the original magma composition. A general negative correlation between Mg# and degree of crystallinity (Fig. 11a and c) implies that the observed differentiation of the groundmass glass is caused by in situ crystallization (or equilibrium crystallization at a constant total composition) of the liquid lava during emplacement. If we take the whole-rock composition of the rootless-cone tephra as the starting liquid composition and allow $\sim 30\%$ crystallization of the observed groundmass mineral assemblage in lava selvages and rootless-cone tephra (plagioclase, augite, and olivine in the fractions of 0.4, 0.4, and 0.2, respectively), the calculated liquid composition agrees with the mean composition of the rootless-cone groundmass glass (Table 5b). The absence of Fe–Ti oxides from the groundmass mineral assemblage of these samples is consistent with the strong FeO and TiO_2 enrichment observed in the groundmass glass and implies that the emplacement temperatures of the lava were above the Fe–Ti oxides liquidus temperatures. The concentrations of S, Cl, and F gradually increase in the remaining liquid as crystallization progresses be-

Table 5a Composition of groundmass glass in Laki lava selvages and rootless cone tephra (RCT) at proximal, medial and distal locations in the Laki lava field. Also shown are average whole-rock composition of rootless cone tephra and crystalline lava (Cr. lava)

	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Cr.%	N
Proximal																	
RCT-gm	49.61	3.61	12.37	15.62	0.24	4.80	9.89	2.70	0.57	0.44	99.84	480	167	416	35.40	20	23/24
SD	0.53	0.36	0.56	0.83	0.03	0.25	0.19	0.19	0.14	0.15	0.24	201	58	55	2.16	11	
Cr. lava-wr ^a	50.15	2.84	13.72	13.65	0.21	5.61	10.40	2.51	0.43	0.32	99.84				42.26		7
SD	0.21	0.09	0.17	0.22	0.00	0.09	0.04	0.11	0.01	0.04	0.20				0.76		
Medial																	
Lava selvage	49.99	3.84	11.82	16.11	0.25	4.64	9.44	2.77	0.58	0.37	99.82	356	240	494	33.92	25	94/53
SD	0.27	0.10	0.21	0.34	0.04	0.14	0.21	0.14	0.11	0.07	0.46	40	73	108	0.93	5	
RCT-gm	49.69	3.97	11.66	16.69	0.24	4.54	9.30	2.73	0.58	0.47	99.85	275	168	390	32.67	38	33/27
SD	0.16	0.18	0.14	0.52	0.02	0.24	0.17	0.07	0.04	0.08	0.22	37	31	20	1.80	9	
RCT-wr	49.42	2.73	13.40	13.88	0.23	5.95	10.64	2.69	0.51	0.41	99.86				43.30	35	3
SD	0.30	0.07	0.11	0.20	0.01	0.12	0.21	0.05	0.02	0.01	0.01				0.81	10	
Cr. lava-wr ^a	49.81	2.65	13.62	13.48	0.23	5.88	10.50	2.77	0.43	0.37	99.73				43.73		3
SD	0.24	0.07	0.43	0.45	0.01	0.07	0.16	0.15	0.05	0.10	0.15				0.56		
Distal																	
Lava selvage	49.78	3.81	11.81	16.19	0.28	4.69	9.37	2.70	0.55	0.37	99.54	340	159	438	34.03	37	9/21
SD	0.19	0.05	0.12	0.09	0.02	0.04	0.08	0.05	0.02	0.03	0.33	14	21	55	0.28	1	
Cr. lava-wr ^a	49.74	2.76	13.84	13.57	0.23	5.83	10.44	2.67	0.44	0.32	99.84	195	150	320	43.35		6/2
SD	0.66	0.10	0.14	0.34	0.01	0.19	0.15	0.13	0.02	0.06	0.30				1.10		

gm, groundmass; wr, whole-rock; Mg# = 100 mg/(Mg + Fe*), Fe* = total Fe; N, number of analyses - 1/1 = major/volatile element analyses; SD, standard deviation (2σ); Cr.%, crystal content of sample in vol. %

^a Average composition of crystalline lava derived from published whole-rock analyses compiled by Thordarson (1955)

Table 5b Crystal fractionation calculations using whole-rock composition of rootless cone tephra as starting liquid composition (see text for further details)

	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	Mg#
RCT-wr	49.42	2.73	13.40	13.88	0.23	5.95	10.64	2.69	0.51	0.41	99.86	43.30
Calc comp ^a	50.60	3.72	10.86	16.73	0.31	4.55	8.85	3.04	0.53	0.56	99.75	32.65
RCT-gm	49.69	3.97	11.66	16.69	0.24	4.54	9.30	2.73	0.58	0.47	99.85	32.67

^a Crystallization required: 29%. gm, groundmass; wr, whole-rock

cause these elements are incompatible with the crystallizing phases. This simple chemical process is complicated by the fact that the lava loses a fraction of these elements (especially S) through degassing, but at the same time the strong FeO enrichment is likely to increase the solubility of sulfur in the liquid lava (Haughton et al. 1974). Consequently, to find the representative concentration of these elements in the bulk lava (the original liquid after vent degassing) it is necessary to correct for the amount of crystallization which occurred during lava transport. When corrected, the S, Cl, and F values of samples from the medial and distal part of the lava still show remarkable uniformity (Fig. 11b; Table A-2, Appendix II).

Geothermometry

Because of the temperature dependence of sulfur solubility and magma rheology, and the long duration of

the Laki eruption, it is important to evaluate the variations in melt temperatures during the eruption. One-atmosphere melting experiments performed on Laki lava samples suggest that magma-reservoir temperatures were at least 1150 ± 10°C (Bell and Humphries 1972), and that lava-emplacement temperatures were above 1085°C (Thordarson 1995). From the empirical glass geothermometer of Helz and Thornber (1987), inclusion chemistry indicates magmatic temperatures near 1140 ± 8°C (Fig. 12).

We estimate the melt temperature of Laki tephra and lava on the basis of MgO and CaO concentrations of groundmass glasses (Helz and Thornber 1987), which is justified by the linear co-variations of these elements (Thordarson 1995). The calculated MgO- and CaO temperatures agree and are internally consistent. The CaO-derived temperatures give, on average, slightly higher values which may be attributed to the difference in the crystallization sequence between the Laki magma (plagioclase-olivine-clinopyroxene) and the

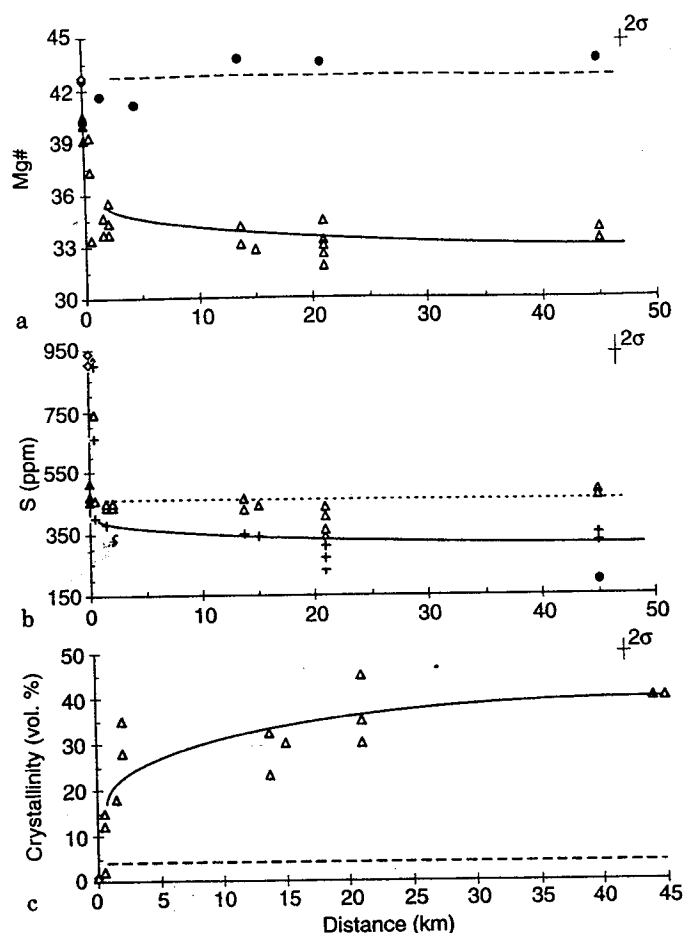


Fig. 11a-c Variation in Mg#, S, and crystallinity with distance from source. Lava selvages and rootless-cone tephra are indicated by *open triangles*, strombolian tephra by *filled triangles*, phreatomagmatic tephra by *open diamonds*, and crystalline lava by *filled circles*. **a** Mg# vs distance. *Solid line* indicates the inferred change in Mg# of lava selvege with distance and *broken line* changes in Mg# of crystalline lava. **b** S vs distance. *Broken and solid line* indicate inferred changes in measured and corrected S abundances (*plusses*), respectively, of lava selvages with distance. **c** Crystallinity of lava selvages and rootless-cone tephra with distance from source. *Solid line* indicates the inferred changes in crystal content of lava selvege with distance; *broken line* indicates the pre-eruption crystal content of the Laki magma. Estimated analytical error (2σ) is shown by the cross in the upper right corner

Kilauea Iki lava lake magma (olivine-clinopyroxene-plagioclase) used for geothermometer calibration. The geothermometer calculations indicate that the melt temperature of the phreatomagmatic tephra was $1138 \pm 9^\circ\text{C}$ at the time of explosive interaction of magma and external water (Fig. 12), whereas the temperature of strombolian tephra was $\sim 1127 \pm 3^\circ\text{C}$ upon melt disintegration. The results suggest that quenching and fragmentation of the phreatomagmatic tephra occurred at greater depth in the feeder dikes. Furthermore, the high melt temperatures of the phreatomagmatic tephra suggest that cooling during ascent had insignificant, if any, effects on the flow regime within the feeder dike. The calculated emplacement temperature for the proxi-

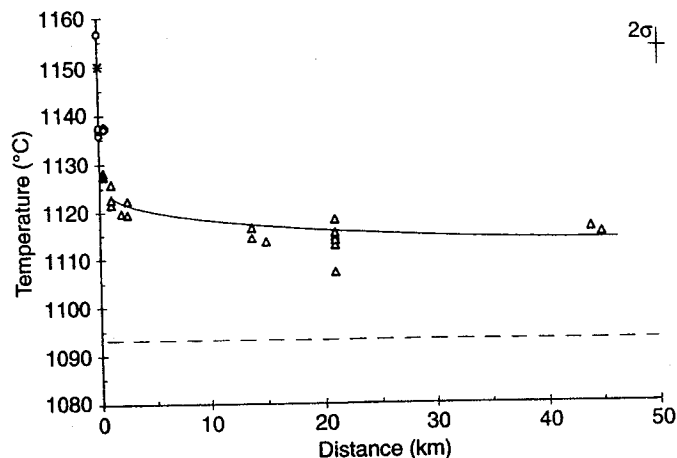


Fig. 12 Laki magma and liquid lava temperatures as a function with distance from source. Temperatures were calculated according to the empirical geothermometer of Helz and Thornber (1987). *Cross* indicates liquidus temperature of the Laki magma as determined by 1-atm melting experiments (Bell and Humphries 1972). Laki inclusions are indicated by *open circles*, phreatomagmatic tephra by *open diamonds*, and strombolian tephra and lava selvages by *filled and open triangles*, respectively. Inferred change in lava temperature with distance from source is indicated by the *solid line*. *Broken line* shows the liquidus temperature of FeTi-oxides in the Laki magma. Estimated analytical error (2σ) is shown by the cross in the upper right corner

mal lava ranges from 1120 to 1126°C and is similar to that of the strombolian tephra. Temperatures obtained from samples of the medial and distal lavas are 10 – 20°C lower (1107 – 1117°C) and show no obvious variation with distance from source (Fig. 12), indicating that lava transport was thermally efficient.

Estimates of volatile release by the Laki eruption

Methodology

Several studies (Rose 1977; Óskarsson et al. 1984; Devine et al. 1984) have demonstrated clearly the validity of a petrologic approach to evaluate the potential atmospheric sulfur yield from eruptions, although the applicability of the method may depend on the pre-eruption condition in the magma source region. For example, Wallace and Gerlach (1994) and Gerlach et al. (1994) argue that the petrologic method underestimates the SO_2 flux from andesite-dacite eruptions at convergent margin volcanoes, because the magma carries the bulk of the sulfur in an accumulated vapor phase. It should be noted, however, that these convergent margin magmas are hybrids residing in shallow crustal magma chambers, and that their pre-eruption evolution is complicated by incomplete magma mixing, crystallization, and fractionation (e.g., Nye et al. 1994).

The large volume and homogeneous composition of the Laki products imply that they were erupted from a compositionally uniform magma body in a deep-seated (>10 km) reservoir at the base of the crust below Ice-

Eruption column and distal haze

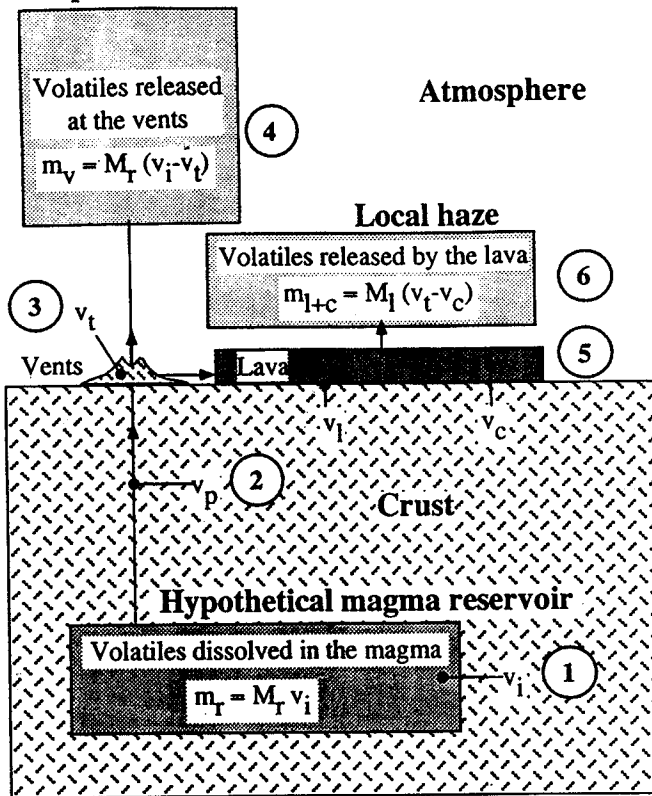


Fig. 13 Illustration outlining a volatile budget model for basaltic fissure eruptions. 1 Hypothetical magma reservoir containing magma of the mass M_r (taken here to be equal to erupted volume of magma) and total amount of dissolved volatiles m_r ; the measured concentration of volatiles in glass inclusions, v_i , represents the concentration of volatiles dissolved in the magma. The total mass of volatiles dissolved in the magma prior to eruption is given by the lowermost equation. 2 Quenched magma fraction due to interaction with external water, represented by phreatomagmatic tephra at the surface containing volatile concentrations v_p . 3 Vent accumulations of pyroclastic material containing concentrations of volatiles v_t , which represents volatile fraction of m_r which did not escape the magma as it emerged from the vents. 4 Eruption column (lava fountain plus convecting plume) containing total amount of volatiles released at the vents, m_v , as given by the uppermost equation. 5 Lava flow of mass M_l and volatile concentration v_l in lava selvages and v_c in crystalline lava. 6 Amount of volatiles, m_{l+c} , released by the degassing of lava during and after emplacement is given by the middle equation

land (Sigmarsson et al. 1991). Based on the chemical similarities between group-1 inclusions and the erupted products (Table 3), we conclude that the inclusions represent the pre-eruption Laki magma. The high volatile concentrations in the inclusions compared with concentrations in groundmass glass of quenched eruption products supports this conclusion. Therefore, we use the petrologic method, modified from Óskarsson et al. (1984) and Devine et al. (1984), to estimate the volatile yield of the Laki eruption.

In fissure eruptions, such as Laki, the release of volatiles by the magma can be viewed as occurring in two main stages (Fig. 13): (a) vent degassing of magma during strombolian explosive activity or Hawaiian-type

lava-fountaining; and (b) degassing of magma during and after flow of lava. Thus, it should be possible to obtain information on the degree of degassing during an eruption by comparing the pre-eruption volatile content to the content of volatiles retained in the products of each phase, although the crystallization of magma as it passes through each stage must be accounted for. The volatile concentrations in rapidly cooled pyroclasts and glassy selvages of proximal lavas are indicators of the amount retained by the magma as it flowed away from the vents. Glass selvages from the base of lava flow units and tephra clasts from the rootless cone deposits indicate the residual amount of volatiles in molten lava upon emplacement at some distance from source. Volatile contents of crystalline lavas represent the final residue of gases which did not escape from the lava into the atmosphere.

Assuming that the total mass erupted equals the mass of magma that contributes to the volatile emission during the eruption, then the volatile budget of the eruption can be schematically modeled as illustrated in Fig. 13. The total mass (m_r) of single volatile species (x) dissolved in magma residing in a reservoir (r) at depth and effectively involved in the eruption is:

$$m_{x,r} = M_r v_{x,i}$$

where M_r is mass of magma which is degassed effectively during the eruption and $v_{x,i}$ is the mass fraction of a gaseous element x in inclusions (i). The total mass ($m_{x,v}$) of each volatile species which escapes from the magma at the vents (v) and is dispersed into the atmosphere by the eruption column is estimated by:

$$m_{x,v} = (v_{x,t} - v_{x,i}) M_r$$

where $v_{x,t}$ is the average mass fraction of the gaseous species x in strombolian tephra and proximal lava (t). Similarly, the mass released into the atmosphere from the lava during emplacement ($m_{x,l}$) and after emplacement ($m_{x,c}$) is estimated according to:

$$m_{x,l} = (v_{x,t} - v_{x,l}) M_l$$

$$m_{x,c} = (v_{x,l} - v_{x,c}) M_l$$

where $v_{x,l}$ and $v_{x,c}$ are the mass fraction of the gaseous species x in glassy lava selvages and crystalline lava, respectively, and M_l is the total mass of lava.

Application

The data used to calculate the volatiles released in various stages of the Laki eruption are presented in Table 6. The mass of each volatile species carried to the surface by the erupting magma is calculated by:

$$m_{x,r} = M_r v_{x,i} = V_r \rho \epsilon v_{x,i}$$

where V_r is total erupted volume taken as 15.1 km³ (Thordarson and Self 1993), ρ is the magma density, 2750 kg m⁻³ (Metrich et al. 1991), $v_{x,i}$ is the mass fraction of individual elements in glass inclusions, and ϵ is

Eruption column and distal haze

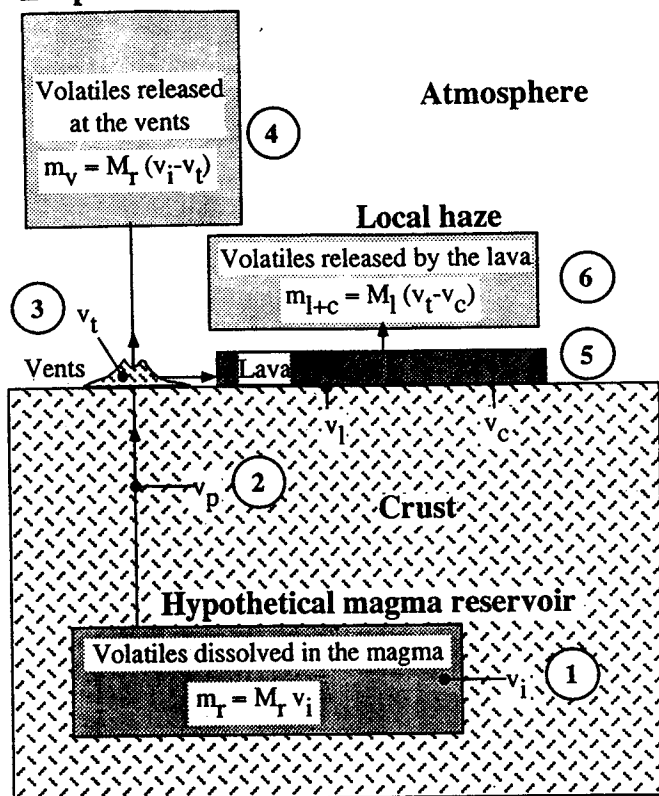


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land (Sigmarsson et al. 1991). Based on the chemical similarities between group-1 inclusions and the erupted products (Table 3), we conclude that the inclusions represent the pre-eruption Laki magma. The high volatile concentrations in the inclusions compared with concentrations in groundmass glass of quenched eruption products supports this conclusion. Therefore, we use the petrologic method, modified from Óskarsson et al. (1984) and Devine et al. (1984), to estimate the volatile yield of the Laki eruption.

In fissure eruptions, such as Laki, the release of volatiles by the magma can be viewed as occurring in two main stages (Fig. 13): (a) vent degassing of magma during strombolian explosive activity or Hawaiian-type

lava-fountaining; and (b) degassing of magma during and after flow of lava. Thus, it should be possible to obtain information on the degree of degassing during an eruption by comparing the pre-eruption volatile content to the content of volatiles retained in the products of each phase, although the crystallization of magma as it passes through each stage must be accounted for. The volatile concentrations in rapidly cooled pyroclasts and glassy selvages of proximal lavas are indicators of the amount retained by the magma as it flowed away from the vents. Glass selvages from the base of lava flow units and tephra clasts from the rootless cone deposits indicate the residual amount of volatiles in molten lava upon emplacement at some distance from source. Volatile contents of crystalline lavas represent the final residue of gases which did not escape from the lava into the atmosphere.

Assuming that the total mass erupted equals the mass of magma that contributes to the volatile emission during the eruption, then the volatile budget of the eruption can be schematically modeled as illustrated in Fig. 13. The total mass (m_x) of single volatile species (x) dissolved in magma residing in a reservoir (r) at depth and effectively involved in the eruption is:

$$m_{x,r} = M_r v_{x,i}$$

where M_r is mass of magma which is degassed effectively during the eruption and $v_{x,i}$ is the mass fraction of a gaseous element x in inclusions (i). The total mass ($m_{x,v}$) of each volatile species which escapes from the magma at the vents (v) and is dispersed into the atmosphere by the eruption column is estimated by:

$$m_{x,v} = (v_{x,t} - v_{x,i}) M_r$$

where $v_{x,t}$ is the average mass fraction of the gaseous species x in strombolian tephra and proximal lava (t). Similarly, the mass released into the atmosphere from the lava during emplacement ($m_{x,l}$) and after emplacement ($m_{x,c}$) is estimated according to:

$$m_{x,l} = (v_{x,l} - v_{x,i}) M_l$$

$$m_{x,c} = (v_{x,l} - v_{x,c}) M_l$$

where $v_{x,l}$ and $v_{x,c}$ are the mass fraction of the gaseous species x in glassy lava selvages and crystalline lava, respectively, and M_l is the total mass of lava.

Application

The data used to calculate the volatiles released in various stages of the Laki eruption are presented in Table 6. The mass of each volatile species carried to the surface by the erupting magma is calculated by:

$$m_{x,r} = M_r v_{x,i} = V_r \rho \epsilon v_{x,i}$$

where V_r is total erupted volume taken as 15.1 km³ (Thordarson and Self 1993), ρ is the magma density, 2750 kg m⁻³ (Metrich et al. 1991), $v_{x,i}$ is the mass fraction of individual elements in glass inclusions, and ϵ is

Table 6

a) Averaged concentration (in ppm) of volatile species in the Laki eruption products						
	S	Cl	F	H ₂ O	CO ₂	Σ
<i>Inclusions</i>						
Laki type (<i>v_i</i>)	1675 ± 225	310 ± 59	665 ± 100	6325 ^a	8530 ^a	17505
<i>Tephra</i>						
Strombolian (<i>v_t</i>)	490 ± 82	225 ± 74	475 ± 89	1915 ^a	1480 ^a	4615
<i>Lava</i>						
Lava selvage (<i>v_l</i>)	350 ± 30	185 ± 55	435 ± 80			
Crystalline lava (<i>v_c</i>)	195 ^a	150 ^a	315 ^a	620 ^a	100 ^a	1380
<i>Total in solidified eruption products</i>						
Lava + tephra (<i>v_s</i>)	205 ± 10	150 ± 5	320 ± 10	655	135	1465
b) Volatile abundances (in ppm) used in calculating the amount released by various eruption stages						
	ΔS	ΔCl	ΔF	ΔH ₂ O	ΔCO ₂	ΔΣ
Total amount released (<i>v_i</i> - <i>v_s</i>)	1470	160	345	5670	8395	16040
Amount released at vents (<i>v_i</i> - <i>v_t</i>)	1185	85	190	4410	7050	12890
Amount released during lava transport (<i>v_i</i> - <i>v_l</i>)	140	40	40	-	-	-
Amount released after lava emplacement (<i>v_i</i> - <i>v_c</i>)	155	35	115	1260 ^b	1345 ^b	3165 ^b
c) Estimates on degree (%) of magma degassing at various eruption stages						
	S	Cl	F	H ₂ O	CO ₂	Σ
Total degassing (<i>v_i</i> - <i>v_s</i>)	88	51	52	90	98	92
Amount retained in solidified products (<i>v_s</i>)	12	49	48	10	2	8
Degassing at vents (<i>v_i</i> - <i>v_t</i>)	71	27	29	70	82	74
Degassing during lava transport (<i>v_i</i> - <i>v_l</i>)	8	13	6	-	-	-
Degassing after lava emplacement (<i>v_i</i> - <i>v_c</i>)	9	11	17	20 ^b	16 ^b	18 ^b

^a From Óskarsson et al. (1984) and Sigvaldason and Óskarsson (1986). Numbers in *italics* probably overestimate the CO₂ concentration in the Laki products and should be regarded as maximum values

^b Calculated as the total amount released by the lava. See text for definitions and further discussion

Table 7 Estimates on the mass (in megatons) of volatiles released by the Laki magma at various stages in the eruption

	SO ₂	HCl	HF	H ₂ O	CO ₂	Σ
Original mass (<i>m_r</i>)	139.1	13.2	29.1	262.6	354.3	798.3
Total mass released (<i>m_r</i>)	122.1	6.8	15.1	235.4	348.6	728.0
Mass retained in solidified products (<i>m_s</i>)	17.0	6.4	14.0	27.2	5.7	70.3
Mass released at vents (<i>m_r</i> - <i>m_v</i>)	98.4	3.6	8.3	183.1	292.8	585.0
Mass released during lava transport (<i>m_r</i> - <i>m_l</i>)	11.6	1.7	1.8	-	-	-
Mass released after lava emplacement (<i>m_r</i> - <i>m_c</i>)	12.5	1.5	5.0	52.3 ^a	55.8 ^a	143.4 ^a

Total volume of magma used in the calculations of 15.1 km³ (lava 14.7 km³; tephra 0.4 km³) is from Thordarson and Self (1993) and the assumed magma density of 2750 kg m⁻³ is from Metrich et al. (1991). See text for definitions and further discussion. Numbers in

italics probably overestimate the CO₂ mass released by the Laki eruption and should be regarded as an absolute maximum

^a Calculated as the total amount released by the lava

the constant required to convert the pure element to the species assumed to be present in the magma (e.g., SO₂, HCl, and HF). As an example, the average inclusion concentration of sulfur, 1677 ppm (0.1677 wt.%), transforms into ~139 Mt of SO₂ carried from depth to the surface by the erupting magma (Fig. 14; Table 7).

The total mass of each volatile species (*m_{x,b}*) released by the eruption is taken as the difference between the inclusion value and the amount retained in the solidified eruption products (*m_{x,s}*), or:

$$m_{x,b} = m_{x,r} - m_{x,s} = (M_r v_{x,i}) - (v_{x,t} M_t + v_{x,c} M_l)$$

where *M_t* is the mass of tephra and other symbols are as defined previously. Application of this equation indicates that more than 85% of the SO₂ and H₂O escaped from the magma during the eruption (Table 6) and that the total amounts released were ~139 Mt SO₂ and ~263 Mt H₂O (Fig. 14; Table 7). Although the CO₂ values reported in Table 6 may be overestimates (see above), we include them in our calculations. The results

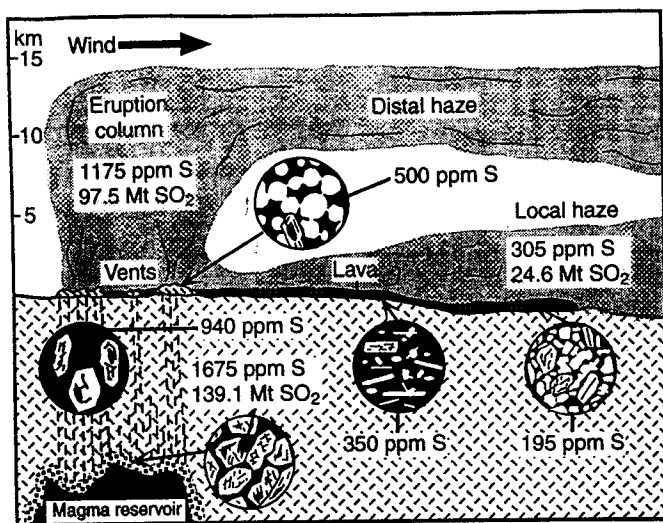


Fig. 14 Illustration of the Laki eruption viewed from the southwest (not to scale) showing the amount of S retained in samples from various eruption stages and their relationship to the degassing of the magma. Also shown is the total amount of SO_2 dissolved in the Laki magma prior to eruption and the estimated SO_2 yield by various eruption stages. The gas columns and plumes generated by the magma degassing are shown as *fine stippled areas*. *Distal haze* denotes the gas liberated at the vents and carried to high altitudes by the eruption columns, and *local haze* denotes that produced by gas rising from the lava flow. Other features are as indicated in Fig. 4

imply that the Laki magma released a maximum of ~ 354 Mt CO_2 (Table 7). The fractions of HCl and HF released are considerably smaller, approximately 50% of the total mass originally dissolved in the magma. This implies that the mobility and solubility of Cl and F in the erupting liquid are governed by somewhat different chemical processes (Carroll and Webster 1994). The Laki eruption liberated a total of 6.8 and 15.1 Mt of HCl and HF, respectively. Our estimates suggest that the total volatile release by the eruption was ~ 730 Mt (Table 7).

The groundmass-glass sulfur concentrations in the phreatomagmatic tephra (mean 944 ± 217 ppm) fall between that of the inclusions and the strombolian tephra. Furthermore, the estimated melt temperature of the phreatomagmatic tephra is near the reservoir temperature of the Laki magma (Fig. 12). This evidence suggests that the phreatomagmatic tephra was produced by early quenching and fragmentation of partly degassed magma within the conduit by explosive interaction of water and magma. The consistently lower, but uniform, S concentration of the strombolian tephra, however, implies that conduit degassing was near completion when fragmentation and venting occurred. The identical S, Cl, and F abundances in the strombolian tephra and proximal lava samples (Tables 4 and 5) support this conclusion. Approximately 70% of the total magmatic SO_2 and H_2O was released as the magma emerged from the fissures (Table 6), and approximately 98 Mt SO_2 and 183 Mt H_2O were discharged into the atmosphere above the Laki fissures at times of vigorous explosive

activity (Fig. 14; Table 7). Moreover, taking the CO_2 concentrations in Table 6 at face value, the Laki magma released $\sim 80\%$ of its carbon dioxide (~ 293 Mt CO_2) through vent degassing. The vent degassing of HCl and HF was considerably less, 27 and 29%, respectively, of the total magmatic mass of these volatiles, signifying an upper tropospheric/lower stratospheric loading of the order of 3.6 Mt for Cl and 8.3 Mt for F.

Volatile concentrations of medial and distal lava, which are very uniform after being corrected for degree of crystallinity (Table A-2, Appendix II), can be taken as the fraction of volatiles retained in the magma upon emplacement in the medial and distal areas of the lava field. The difference in the concentration of the strombolian tephra and the lava selvages (Table 6) can be used to estimate the volatile mass released during lava emplacement (Fig. 13). Similarly, the difference between the concentration in lava selvages and crystalline lava denotes the mass released during cooling and crystallization of the lava after the flow had stopped its advance. According to these arguments ~ 11.6 Mt SO_2 , ~ 1.7 Mt HCl, and ~ 1.8 Mt HF were liberated from the moving lava during transport from vents, followed by release of ~ 12.5 Mt SO_2 , ~ 1.5 Mt HCl, and ~ 5.0 Mt HF during cooling and crystallization (Fig. 14; Table 7).

Discussion

Magma degassing and implications for eruption dynamics

The S concentrations in Laki glass inclusions are consistent with observed S-FeO solubility relations in submarine tholeiites (Wallace and Carmichael 1992; Mathiez 1976), suggesting that the Laki magma was near sulfur saturation prior to the eruption (Fig. 15). Even though the analysis of Óskarsson et al. (1984) overestimates the CO_2 concentrations in Laki inclusions, the results show clearly that the melt was above its CO_2 saturation limit before it erupted (Metrich et al. 1991). Furthermore, the concentrations of other volatiles measured in Laki inclusions imply that the melt was undersaturated with H_2O , Cl, and F (Johnson et al. 1994; Carroll and Webster 1994). These data imply that magma over-pressurization induced by CO_2 saturation was responsible for the onset of the Laki eruption and suggest an early exsolution of volatiles (primarily CO_2) either in the reservoir or in the lowest part of the feeder dike(s).

The S content of the groundmass glass from Laki products reflects progressive magma degassing during its rise through the crust and after its emergence from the vents as tephra and lava (Fig. 15). The differences in the melt temperatures and composition of the tephra types can be reconciled if the phreatomagmatic tephra was quenched at greater conduit depths than the strombolian tephra. The abundance of highly vesicular clasts

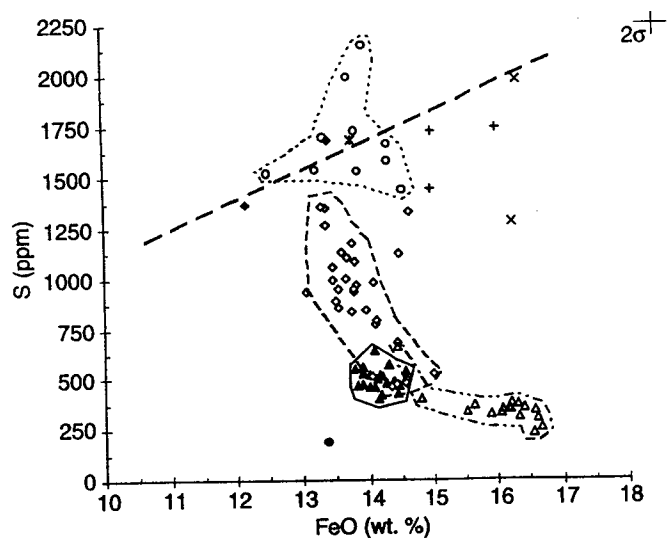


Fig. 15 S (ppm) variation as a function of FeO (wt.%) in Laki samples. Note apparent negative correlation of S and FeO shown by phreatomagmatic tephra and strong clustering of strombolian tephra samples (see text for discussion). *Open circles* Laki-type inclusions; *asterisk* mean of Laki-type inclusions; *filled diamonds* inclusions with high MgO content; *pluses* inclusions with high FeO content; *crosses* inclusions with high FeO and TiO₂ content; *open diamonds* phreatomagmatic tephra; *filled triangles* strombolian tephra; *open triangles* lava selvages; *filled circle* mean of crystalline lava (whole rock). *Heavy broken line* indicates the sulfur solubility trend in submarine basalts as shown by Wallace and Carmichael (1992) and Mathez (1976)

in the phreatomagmatic tephra demonstrate that vesiculation and foam formation was well under way prior to the explosive interaction of water and magma which produced the deposits. Moreover, the data (Figs. 9b and 15) show that the magma had exsolved ~20–70 wt.% of its original S content when melt fragmentation and quenching occurred preventing further degassing. The more evolved composition (Fig. 7) and mature vesicle structure of the strombolian clasts (Fig. 8a) demonstrate that the strombolian tephra was produced by fragmentation of a foam layer, presumably at very shallow conduit levels. Cashman and Mangan (1994) have suggested that such magmatic foams are produced by “runaway” bubble nucleation and growth on timescales of hundreds of seconds, where vesiculation and magma fragmentation occur almost simultaneously. Episodic recurrence of this process may explain the relatively low, but uniform, S concentrations in all of the strombolian fall units (Fig. 15). The high abundances of Pele’s hair in the strombolian tephra suggest that it was erupted by hot gas-rich fountains at high velocities (Thorarinnsson 1984; Ólafsson et al. 1984; Shimozuru 1994), which may explain the fused outer surfaces of strombolian clasts (Fig. 8b). Furthermore, the analytical data imply that the lava contained ~500 ppm S when it flowed from the vents, or alternatively that the magma liberated ≥70 wt.% of its original S content through conduit and vent degassing. The results of Haughton et al. (1974) indicate that the sulfur capacity

of the Laki magma at 1 atm pressure is 400 ppm, which is similar to the measured sulfur concentrations in strombolian tephra and the proximal lava, implying that the vent degassing of S was limited by its solubility.

The large fraction of volatiles released at the vents implies effective separation of volatiles and melt during ascent. In this context it is important to recognize that the Laki eruption was episodic, on the scale of several days to weeks, with intense fountaining and voluminous lava outpourings repeated each time a new pulse of magma reached the surface. Also note that the low ratio of tephra to lava volume (~1/35) shows that the bulk of the magma was erupted as lava. Estimated rise velocities of the magma pulses are in the range of 0.04–0.32 m/s (Thordarson 1990) and are similar to the advance velocities (0.03–0.08 m/s) of the lava surges produced by the eruption episodes (Thordarson and Self 1990). This suggests that the dominant lava emissions occurred at eruption velocities <1 m/s. Such low velocities imply an eruption in the homogeneous flow regime (gas and melt rise with the same velocity; Vergnolle and Jaupart 1986). However, the 800- to 1450-m-high fountains at Laki indicate magma exit velocities in the range of 130–170 m/s (Fig. 7b; Wilson and Head 1981), suggesting that the initial magmatic explosive phases of the episodes involved eruption velocities which were two orders of magnitude higher. Such intense fountains are driven by gas jets containing entrained melt fragments (Vergnolle and Jaupart 1986). Heights of fountains are controlled primarily by the contents of exsolved magmatic gas (Head and Wilson 1987). To produce the recorded fountain heights, the Laki magma had to have exsolved 50–75% of its original volatile content (Wilson and Head 1981). These considerations suggest that the explosive phases resulted from separated two-phase flow in the magma conduit, implying that transitions from separated to homogeneous flow regime occurred during each eruption episode (Vergnolle and Jaupart 1986).

As suggested above, the closed-cell spherical foam structure of the strombolian tephra is best explained by vesiculation and subsequent fragmentation at shallow conduit levels. However, this process alone cannot account for the high degree of vent degassing suggested by our data, because the volatile mass exsolved by the magma upon venting indicates a bulk gas-to-melt volume ratio of >100. As shown previously, the volume of a single fall unit is <3% of the total volume of magma erupted in separate episodes. Consequently, the cumulative bubble volume of the tephra can only be a fraction (<<0.1) of the total gas volume upon eruption, indicating that large quantities of gas exsolved from the melt without leaving clear physical evidence of its presence. These considerations suggest that most of the gas erupted as a free volatile phase, which supports the inferred evolution of separated conduit flow during the explosive phases. Thus, we conclude that the explosive phases at Laki involved an early and accelerated exso-

lution of volatiles (primarily CO_2), followed by a second vesiculation burst at shallow conduit levels which was caused by H_2O and SO_2 oversaturation of the melt (Bottinga and Javoy 1991).

Based on the considerations of volatile degassing at Laki, the dynamics of the eruption may be viewed as follows: As the magma began its rise to the surface, volatile exsolution was accelerated, liberating initially mostly CO_2 . Rapid coalescence of bubbles led to formation of a separated two-phase flow in the feeder dike which enhanced the driving-pressure gradient in the magma column (Vergnolle and Jaupart 1986), resulting in intense fountaining activity at the surface and efficient liberation of CO_2 , SO_2 , and H_2O . This initial bulk degassing resulted in a gradual, but rapid, transition from separated to homogeneous flow regime and deceleration of magma rise velocities. We speculate that during this transition period part of the magma became saturated with H_2O and SO_2 , triggering "runaway" vesiculation. These vesicles accumulated rapidly at the top of the magma column to form a gas-rich foam layer which disintegrated upon extrusion to produce the strombolian fall units. Trailing the foam head was more voluminous, less vesicular, partly degassed magma, corresponding to the lava-producing phase of the eruption episodes. Cyclic recurrence of these phenomena can explain the episodic nature of the eruption and the uniformity of the texture and clast morphology of the strombolian tephra deposits. The dominance of separated flow regimes at the beginning of each episode implies that the bulk of the volatiles were discharged during the vigorous explosive activity. The explosive phases also produced the highest eruption columns of the Laki eruption (Thordarson and Self 1993) and therefore are likely to have discharged large quantities of volatiles at or above the tropopause. Consequently, the effective altitude of atmospheric volatile loading from eruptions such as Laki is best evaluated by the maximum height reached by the eruption columns.

Fluorine and chlorine degassing

Many previous studies based on whole-rock analyses of crystalline lava and glassy tephra have argued that Cl and F are not effectively lost from the magma during an eruption (e.g., Sigvaldason and Óskarsson 1976, 1986; Aoki and Kanisawa 1979; Ishikawa et al. 1980; Aoki et al. 1981). These arguments are supported primarily by experimental Cl and F solubility data, theoretical and model considerations, and the relatively small variation in measured Cl and F concentrations ($\leq 20\%$) in samples representing various phases in a single eruption. However, Metrich (1990) and Metrich and Clocchiatti (1989) showed that the 1763 AD Etna magma lost 45% of its original Cl upon eruption. Results of Devine et al. (1984) indicate up to 50% Cl loss by magmas of intermediate compositions; similar results have been ob-

tained from products of basaltic fissure eruptions in shallow marine and subaerial environments (Unni and Schilling 1978).

Our results show that the Laki magma liberated ~50% of its original Cl and F abundances during the eruption. Approximately half of the Cl and F released into the atmosphere was discharged at the Laki vents. The remainder was liberated during lava transport and during cooling of the lava after it was emplaced (Table 6). The nearly equal degrees of degassing for these two elements appear consistent with their similar chemical characteristics and expected compatibility at magmatic temperatures, especially in basaltic melts free of apatite and hydrous mineral phases (Carmichael et al. 1974). Laki inclusions have Cl and F contents which are well below HCl and HF saturation levels in basaltic melts (Carroll and Webster 1994); thus, fluxing by other effervescing volatiles is a likely explanation for the effective removal of these elements from the Laki magma (Iwasaki and Katsura 1967; Rowe and Schilling 1979). Similar analyses of products from other basaltic fissure eruptions in Iceland also indicate that approximately half of the magmatic Cl and F was released into the atmosphere upon extrusion (Thordarson 1995).

Atmospheric loading of SO_2 and rate of degassing with time

In the case of short-lived explosive eruptions with well-documented atmospheric perturbations, such as El Chichón 1982 and Pinatubo 1991, the potential amount of H_2SO_4 aerosols generated from the measured SO_2 emission by the eruption is only slightly less than the maximum measured stratospheric aerosol loading (e.g., Hofmann 1987; McCormick et al. 1995). Obviously, this does not hold for predominantly effusive fissure eruptions, such as Laki, where the ratio of tephra to lava is less than 1:20 and the eruption columns just manage to penetrate the tropopause. In assessing the SO_2 loading of long-duration (months to years) lava-producing fissure eruptions, it is important to evaluate the dynamics of degassing, the timing, and the degree of degassing by separate eruptive phases and episodes. One must also consider the volume erupted by each phase and episode, changes in discharge, and eruptive behavior.

Contemporary descriptions of the Laki eruption indicate that the highest rates of atmospheric volatile mass loading (i.e., magma degassing) coincided with periods of vigorous eruption and peak magma discharge at the Laki fissures (Thordarson and Self 1993; Thordarson 1995), implying a direct correlation between magma and gas discharge. The Laki eruption produced ~9.0 km³ of lava and tephra in the first 40 days, approximately 60% of the total volume produced. Consequently, Laki released ~73.0 Mt of SO_2 into the atmosphere during this period, equivalent to atmospheric mass loading of ~1.7 Mt per day. With 70% of this amount liberated at the vents and carried by the eruption columns to altitudes between 6 and 13 km, the

Laki eruption is estimated to have injected ~ 1.3 Mt per day of SO_2 into the upper troposphere and the lower stratosphere during this period. For comparison, the annual SO_2 discharge by the present Pu'u'O'o-Kupaianaha eruption at Kilauea, Hawaii, is ~ 0.7 Mt (Elias et al. 1993) or approximately half of the *daily* output by the Laki fissures. The estimated SO_2 yield of the May to August 1783 Asama eruption in Japan is 1.0–3.5 Mt (Thordarson 1995), equivalent to the amount of SO_2 released by Laki in a few days.

The Laki eruption produced approximately 4.0 km^3 of lava and tephra between August and October, half the amount produced during the first 40 days of the eruption (Thordarson and Self 1993). Magma production at the Laki fissures in August and September of 1783 is estimated as 3.5 km^3 in 60 days (Thordarson 1990, 1991) yielding 0.5–0.6 Mt SO_2 emitted daily, dropping to ~ 0.2 Mt per day in October and less than 0.1 Mt per day during the waning stages of the eruption.

The moving Laki lava flow released approximately 25 Mt of SO_2 into the troposphere (Table 8), with an almost equal amount released by degassing of liquid lava during and after emplacement, consistent with contemporary descriptions of gases rising from the advancing lava and a 2-year postemplacement emission of steam and other gases (Steingrímsson 1783, 1788). The lava emissions represent approximately one fifth of the total SO_2 released by the Laki eruption.

A total atmospheric SO_2 loading of ~ 122 Mt by the Laki event gives a potential H_2SO_4 -aerosol yield of ~ 250 Mt. Assuming that the aerosol particles consisted of 75% H_2SO_4 and 25% H_2O (Thomason and Osborn 1992), this is equivalent to ~ 185 Mt of pure H_2SO_4 . According to the degassing rates given above, the eruption released sufficient mass of SO_2 in the first 2 months to generate ~ 175 Mt of H_2SO_4 . This value is similar to an independent estimate of ~ 200 Mt (Stothers 1996) for the accumulative total column H_2SO_4 -aerosol yield from Laki in the same period using information on atmospheric turbidity over Europe. The agreement of two totally different methods of estimating the atmospheric loading indicates that the petrologic method gives reliable results for Laki. These values cannot be directly compared with earlier aerosol mass estimates based on the amount of Laki sulfate fallout in Greenland (e.g., Clausen and Hammer 1988), because the findings of Fiacco et al. (1994) indicate that the fallout represents only the portion of the Laki aerosol cloud which remained aloft in the atmosphere for more than 1 year. The volatile contribution due to degassing of the Laki lava flow is only 18% of the total amount released by the magma. These gaseous emissions were confined mostly to the lowest regions of the troposphere and therefore only important on a regional scale in Iceland. A more comprehensive discussion on the environmental and climatic effects of the Laki eruption is presented by Fiacco et al. (1994) and Thordarson, 1995.

Appendix I

Sample locations and representation

Systematic sampling of vent and rootless-cone tephra, and lava, was conducted to obtain an even spatial and temporal coverage of the eruption products (Figs. 1 and 4; see also Table 1). The phreatomagmatic tephra and the strombolian tephra samples (Table 1) provide information about the volatile and magma composition during magma rise in the conduit and upon vent extrusion. The groundmass glass of lava selvages and rootless-cone tephra represent the liquid lava which was quenched rapidly upon emplacement at the respective sample location. These samples provide information about the chemical and the physical properties of the liquid lava at various distances from source. Samples VRC-9, VRC-11, L-23, and LEIRC-3-1 to LEIRC-20A-8 represent the lava stream which emerged from the fissure segments SW of Laki and advanced down the Síða highlands through the Hellisá river valley before entering the Skaftár River gorge (Thordarson and Self 1993). VRC-9 and VRC-11 are tephra samples from rootless cones located ~ 2 km south of the fissures in Varmárdalur (location 9) and sample L-23 is from the basal selva of lava flow unit near Kálfá (location 7) and field relations suggest that both units were formed before 30 June 1783 (Thordarson 1990, 1991). The lava and tephra samples from the Leidólfssfell rootless cone group (location 12) were produced by explosive interaction of lava and water around 17 June 1783 (Steingrímsson 1783, 1788; Thordarson 1990, 1991). Sample locations 7 and 12 are 16 and 21 km from the source, respectively, and thus come from the medial part of the early formed lavas. The distal lava of the same flow field is represented by samples L-24 and L-25, which compose the basal and surface selva, respectively, of a single 3.5- to 6.0-m-thick flow unit (Sal-9) at Eldvatnsbrú (location 8). This flow unit was emplaced sometime between 14 June and 12 July 1783. The samples L-20, L-21 (location 6), BRC-9 (location 10), and NIRC-7 (location 11) are from flow units which belong to the lavas produced by the fissures NE of Laki between 1 August and 1 October. At location 6, a distance of 14 km from source, the Laki lava is composed of two emplacement units (E1 and E2), each containing several flow units. Sample L-20 is from the surface glass selva of the uppermost flow unit of the lower emplacement unit (E1FU4), whereas L-21 is the basal selva of a flow lobe within the lower half of the upper emplacement unit (E2FU2). Sample BRC-9 is tephra from the Blængur rootless cone group, located 0.5 km south of fissures northeast of Laki and sample NIRC-7 is tephra from the Innrieyrar rootless cone group located 1.5 km north of the fissures. Both samples are representative of the proximal lava.

Appendix II

Table AII-1 Major element and volatile composition of the samples used in this study

Laki inclusions:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	N	Host	
Group 1. Laki composition																		
L-i-1	49.88	2.87	12.73	14.55	0.24	6.05	10.14	2.69	0.47	0.19	99.82	1430	270	660	42.55	1	P	
L-i-2	49.61	2.93	12.06	13.74	0.22	6.31	10.00	2.91	0.53	0.34	98.65	1985	250	820	44.98	1/2	P	
L-i-3	49.61	2.85	13.07	13.88	0.24	5.77	10.39	3.01	0.42	0.25	99.50	1520	300	705	42.56	6/5	P	
L-i-4	49.72	2.88	12.50	14.33	0.23	5.91	10.51	2.85	0.41	0.30	99.65	1570	290	530	42.35	4/3	P	
L-i-5	49.25	2.93	13.40	13.98	0.24	5.78	10.25	2.98	0.42	0.27	99.50	2140	290	665	42.40	2/3	P	
L-i-6	49.83	2.73	13.56	13.24	0.17	5.91	10.49	2.85	0.38	0.33	99.50	1530	360	720	44.28	1	O	
L-i-7	49.24	3.07	13.17	14.33	0.25	5.57	10.38	2.83	0.47	0.31	99.62	1660	270	530	40.92	2/3	O	
L-i-8	49.48	3.21	13.41	13.84	0.18	5.32	10.68	2.74	0.36	0.24	99.46	1720	440	690	40.63	2	O	
L-i-9	49.69	3.25	13.29	13.35	0.19	5.83	10.66	2.85	0.36	0.24	99.72	1690	350	550	43.77	1	O	
L-i-10	50.51	2.84	13.36	12.51	0.22	5.35	10.98	2.64	0.38	0.31	99.10	1515	270	780	43.24	1/2	C	
Group 2. High FeO																		
L-i-11	48.70	3.00	12.48	16.01	0.24	5.51	10.25	2.46	0.39	0.34	99.39	1740	185	450	38.02	2	O	
L-i-12	49.05	2.98	12.01	15.00	0.23	6.38	10.60	2.43	0.57	0.26	99.51	1720	260	530	43.12	2	O	
L-i-13	50.05	3.10	11.74	14.99	0.23	5.99	10.38	2.41	0.48	0.32	99.69	1435	295	630	41.61	2	P	
Group 3. High FeO and TiO ₂																		
L-i-14	48.86	3.40	13.15	16.36	0.24	4.78	9.61	2.58	0.42	0.33	99.73	1970	310	710	34.23	2/3	P	
L-i-15	48.87	3.45	13.00	16.26	0.19	4.62	9.50	2.55	0.48	0.37	99.29	1275	285	610	33.61	4/3	P	
Group 4. High MgO																		
L-i-16	49.65	2.17	13.45	12.17	0.21	6.96	11.51	3.01	0.38	0.20	99.72	1244	262	618	50.44	4/5	P	
L-i-17	49.13	2.00	12.03	13.41	0.23	8.47	11.37	2.47	0.39	0.17	99.67	1675	198	562	52.95	4/5	P	
Strombolian tephra:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Ves.%	Cr.%	N
LS1-01-11	49.97	3.05	13.24	14.19	0.23	5.41	10.08	2.81	0.45	0.35	99.78	418	150	468	40.43	65	1	3
LS1-01-12	50.13	3.01	13.28	14.15	0.22	5.43	10.14	2.69	0.49	0.36	99.90	402	159	431	40.62	45	1	3
LS1-04-12	50.09	3.08	13.21	14.27	0.21	5.47	9.83	2.83	0.45	0.32	99.75	480	225	515	40.57	50	3	12/9
LS1-04-14	50.11	3.10	13.24	14.49	0.19	5.37	9.87	2.75	0.46	0.35	99.92				39.78	70	2	3
LS1-04-17	49.98	3.04	13.19	14.45	0.21	5.47	9.77	2.79	0.45	0.28	99.63	425	235	445	40.26	75	1	4
LS1-04-18	50.01	3.06	13.18	14.36	0.22	5.37	9.75	2.85	0.44	0.32	99.55	480	225	500	39.98	88	1	4
LS2-06-11	49.96	3.01	13.38	14.12	0.24	5.44	10.15	2.83	0.46	0.35	99.92	498	148	455	40.70	55	5	3
LS2-06-12	49.93	3.08	13.20	14.56	0.26	5.31	9.87	2.80	0.49	0.36	99.86	534	147	518	39.41	55	3	3/2
LS3-02-31	49.96	3.01	13.24	14.00	0.21	5.42	9.90	2.84	0.45	0.31	99.34	455	245	445	40.81	71	1	3/4
LS3-02-32	49.92	3.02	13.20	14.17	0.21	5.44	9.81	2.82	0.44	0.26	99.29	515	245	525	40.64	84	1	3/4
LS3-02-33	49.93	3.01	13.21	13.92	0.21	5.37	9.89	2.71	0.46	0.34	99.05	470	245	525	40.75	78	1	3
LS3-02-34	49.82	3.02	13.22	13.84	0.23	5.45	9.86	2.59	0.44	0.33	98.79	465	245	450	41.24	82	1	3
LS3-02-21	49.98	3.04	13.18	13.92	0.22	5.39	9.89	2.56	0.44	0.27	98.90	520	320	530	40.83	84	1	3
LS3-02-22	49.61	3.00	13.12	14.08	0.20	5.44	9.92	2.75	0.47	0.28	98.87	640	250	310	40.78	90	0	3
LS3-02-23	50.18	3.04	13.31	13.79	0.22	5.43	9.85	2.53	0.46	0.32	99.12	545	280	485	41.22	77	0	3
LS3-02-24	49.20	2.94	13.19	13.92	0.23	5.37	9.77	2.67	0.44	0.34	98.07	555	175	415	40.76	88	0	3
LS3-03-11	49.92	2.96	13.15	14.00	0.19	5.43	9.92	2.86	0.43	0.30	99.15	515	315	450	40.88	78	1	3
LS3-03-12	49.85	3.04	13.08	14.20	0.24	5.33	9.88	2.80	0.44	0.27	99.14	510	320	525	40.09	57	1	3/4
LS3-03-13	49.84	3.04	13.28	14.30	0.19	5.44	9.91	2.63	0.47	0.32	99.41	570	200	485	40.40	61	1	3/4
LS3-03-15	49.23	3.03	13.14	14.08	0.21	5.43	9.77	2.71	0.42	0.26	98.28	455	250	510	40.73	72	1	3
LS4-01-11	50.10	3.12	13.11	14.38	0.23	5.20	10.12	2.80	0.49	0.36	99.92	484	164	496	39.19	45	5	3
LS4-01-12	50.03	3.15	13.06	14.46	0.25	5.22	10.08	2.85	0.48	0.36	99.94	461	134	482	39.14	48	2	3
Phreatomagmatic tephra:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Ves.%	Cr.%	N
LP1-U2-1	49.67	2.80	13.38	13.69	0.22	5.85	11.02	2.23	0.62	0.25	99.73	1090	256	345	43.19	2	5	7/8
LP1-U2-2	49.50	2.74	13.50	13.47	0.21	5.98	11.07	2.34	0.49	0.24	99.54	1050	190	415	44.18	0	1	5/6
LP1-U2-3	49.88	2.77	13.45	13.37	0.18	5.88	11.20	2.28	0.55	0.26	99.81	1115	210	460	43.93	0	1	2/3
LP1-U2-4	49.67	2.74	13.32	13.68	0.21	5.75	10.96	2.32	0.55	0.25	99.46	995	200	455	42.82	1	2	3/4
LP1-U2-21	49.52	2.72	13.25	13.79	0.17	6.01	11.02	2.21	0.71	0.23	99.63	930	180	460	43.71	54	5	2/4
LP1-U2-61	50.19	3.10	13.17	14.58	0.25	5.26	9.65	2.30	0.71	0.25	99.47	490	130	335	39.14	55	2	1/3
LP1-U2-62	49.53	2.98	13.25	14.56	0.23	5.64	10.12	2.45	0.72	0.33	99.81	1260	240	265	40.85	10	1	2
LP1-U3-2	49.61	2.84	13.46	13.61	0.23	5.71	11.12	2.41	0.46	0.28	99.71	1125	185	485	42.77	49	7	2/4
LP1-U5-22	49.34	2.69	13.40	13.82	0.22	5.93	10.95	2.41	0.69	0.24	99.68	1075	450	320	43.34	38	1	2
LP1-U5-42	49.71	3.08	13.22	13.97	0.23	5.33	9.74	2.74	0.44	0.29	98.76	835	240	485	40.48	50	10	2/3
LP1-U5-61	49.76	3.01	13.23	14.11	0.22	5.35	9.92	2.76	0.45	0.31	99.13	770	295	460	40.32	69	1	2
LP1-U5-62	49.98	3.05	13.06	14.12	0.21	5.33	9.80	2.82	0.44	0.30	99.09	785	350	440	40.21	59	1	2
LP2-05-11	49.90	3.00	13.46	13.91	0.23	5.25	10.14	2.86	0.47	0.35	99.58	958	143	426	40.21	28	1	3
LP2-05-12	49.52	2.38	13.70	12.60	0.21	6.46	11.70	2.58	0.32	0.25	99.71	980	122	381	47.75	0	1	2
LP2-05-13	50.10	2.96	13.27	14.27	0.23	5.40	10.21	2.89	0.46	0.33	100.12	507	137	505	40.25	80	2	3/1

(Continued)

Table AII-1 (Continued)

Lava selvages:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Cr.%	N	D (km)
L-20	49.75	3.78	11.79	16.31	0.27	4.52	9.37	2.78	0.56	0.43	99.55	460	270	613	33.07	35	16/11	14
L-21	50.03	3.86	11.84	16.05	0.25	4.66	9.48	2.78	0.57	0.36	99.89	434	309	608	34.10	23	75/39	14
L-23	50.31	3.64	11.49	16.57	0.26	4.77	8.81	2.56	0.90	0.37	99.69	442	228	521	33.88	30	3/3	16
L-24	49.78	3.80	11.85	16.17	0.27	4.69	9.43	2.66	0.55	0.36	99.56	467	217	604	34.05	38	4/16	45
L-25	49.78	3.82	11.77	16.20	0.29	4.69	9.29	2.75	0.54	0.39	99.51	470	226	594	34.01	35	5/3	45
Rootless cone tephra RCT-groundmass glass:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum	S	Cl	F	Mg#	Cr.%	N	D (km)
VRC-9-1	49.89	3.48	12.12	15.52	0.25	4.80	10.00	2.77	0.53	0.39	99.76	451	182	496	35.53	35	3/3	2
VRC-11-11	49.66	3.68	11.98	15.90	0.24	4.67	9.90	2.82	0.56	0.42	99.83	435	168	471	34.35	35	3/2	2
VRC-11-12	49.95	3.67	12.00	16.08	0.26	4.59	9.79	2.80	0.57	0.42	100.12	447	165	644	33.70	28	3/3	2
BRC-9-31	50.06	3.15	13.28	14.45	0.19	5.25	9.88	2.77	0.44	0.36	99.84	740	270	465	39.32	12	4/4	0.5
BRC-9-32	48.81	4.37	13.04	14.81	0.22	4.78	9.70	2.61	0.79	0.73	99.87	459	268	397	37.31	15	3/3	0.5
BRC-5-1	48.91	3.39	12.05	16.92	0.27	4.76	10.09	2.66	0.43	0.37	99.86	914	120	454	33.37	2	3/3	0.5
NIRC-7-11	49.62	3.60	11.97	15.64	0.25	4.65	10.12	2.84	0.62	0.42	99.74	435	201	482	34.62	18	2/3	1.5
NIRC-7-12	49.87	3.71	11.90	16.22	0.26	4.63	9.65	2.21	0.78	0.41	99.62	448	173	534	33.70	18	2/3	1.5
LEIRC-3-1	49.54	3.86	11.69	16.56	0.27	4.64	9.37	2.75	0.58	0.44	99.72	335	190	575	33.31	45	8/5	21
LEIRC-3-2	49.67	3.94	11.71	16.61	0.25	4.59	9.33	2.75	0.56	0.48	99.87	435	250	535	32.97	45	8/5	21
LEIRC-3A-1	49.48	4.34	11.45	17.69	0.24	4.10	9.03	2.59	0.65	0.62	100.18				29.22	50	4	21
LEIRC-41-3	49.86	3.91	11.81	16.26	0.20	4.79	9.50	2.69	0.53	0.46	100.02				34.40	30	3	21
LEIRC-42-3	49.71	3.92	11.54	16.67	0.25	4.51	9.19	2.81	0.57	0.40	99.57	340	215	540	32.54	30	5/5	21
LEIRC-20A-8	49.86	3.88	11.75	16.34	0.22	4.63	9.36	2.76	0.56	0.39	99.76	405	265	495	33.55	30	5/5	21
RCT whole-rock composition:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum				Mg#			
LEIRC-02	49.60	2.74	13.31	14.03	0.22	5.92	10.46	2.67	0.51	0.41	99.87				42.91	21		
LEIRC-03A	49.59	2.66	13.36	13.66	0.24	6.08	10.60	2.74	0.53	0.41	99.87				44.23	21		
LEIRC-12	49.07	2.80	13.52	13.96	0.23	5.85	10.87	2.65	0.50	0.42	99.85				42.75	21		
New whole-rock analyses of Laki lava:																		
Label	SiO ₂	TiO ₂	Al ₂ O ₃	FeO	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅	Sum				Mg#			
L-3	50.04	2.77	13.88	13.73	0.22	5.55	10.42	2.45	0.42	0.28	99.76				41.88	1		
L-9	50.10	2.74	13.86	13.97	0.22	5.49	10.43	2.39	0.43	0.29	99.92				41.21	1		
LEI-04	49.58	2.71	13.27	13.98	0.23	5.93	10.41	2.81	0.54	0.41	99.87				43.04	21		
LEI-12	50.05	2.66	13.49	13.33	0.22	5.88	10.68	2.61	0.51	0.44	99.85				44.01	21		
L-14	50.17	2.80	13.96	13.96	0.22	5.52	10.42	2.42	0.41	0.29	100.17				41.35	47		

Host, mineral hosting inclusion; P, plagioclase; O, olivine; C, clinopyroxene; D, distance from vents in km; Mg# = 100 Mg/(Mg + Fe*), Fe* = total Fe; Ves.% and Cr.%, vesicularity and crystallinity respectively; N, number of analyses

Table AII-2 Correction of measured S, Cl and F values in Laki lava selvages and rootless cone tephra for the effect of crystallization during lava emplacement

Label	Mg#	FeO	Cr.%	S	S ^a	Cl	Cl ^a	F	F ^a	N
Lava selvage										
L-20	33.07	16.31	32	460	375	270	220	615	500	11
L-21	34.10	16.05	23	435	330	310	235	610	460	39
L-23	32.73	16.42	30	440	340	230	175	520	400	6
L-24	34.05	16.17	38	490	355	195	145	545	395	16
L-25	34.01	16.17	35	470	350	225	165	595	440	3
Rootless cone tephra										
VRC-9-1	35.53	15.52	35	451	334	182	135	496	367	3
VRC-11-11	34.35	15.90	35	435	322	168	124	471	349	3
VRC-11-12	33.70	16.08	28	447	349	165	129	644	503	3
BRC-9-31	39.32	14.45	12	740	660	270	240	465	415	3
BRC-9-32	37.31	14.81	15	459	400	265	230	395	345	3
BRC-5-1	33.37	16.92	2	914	896	120	117	454	445	3
NIRC-7-11	34.62	15.64	18	435	368	201	171	482	408	3
NIRC-7-12	33.70	16.22	18	448	380	173	147	534	452	3
LEIRC-3-1	33.31	16.56	45	335	230	190	130	575	395	8/5
LEIRC-3-2	32.97	16.61	45	435	300	250	170	535	370	8/5
LEIRC-42-3	32.54	16.67	30	340	260	215	165	540	415	5/5
LEIRC-20A-8	33.55	16.34	30	405	310	265	205	495	380	5/5

^a Corrected values; N, number of analyses

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