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1 **Temporal evolution of magma and crystal mush storage conditions in the**
2 **Bárðarbunga-Veiðivötn volcanic system, Iceland**

3

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26 Geochemistry

27

28 **1. Introduction**

29 For more than 100 years, the concept of a melt-dominated and long-lived magma chamber
30 has been a commonly accepted paradigm in volcanology. However, our understanding of volcanic
31 plumbing systems and processes has improved in the last few decades due to new geophysical,
32 petrological and geological evidence (e.g., Ryan, 2018; Sinton and Detrick, 1992; West et al., 2001).
33 Magma plumbing systems beneath active volcanoes are now envisaged to be characterized by sets
34 of crustal reservoirs that are dominated by relatively liquid-poor crystal mushes (Edmonds et al.,
35 2019; Marsh, 2006). Crystal mushes are dynamic horizons made of a semi-rigid framework of
36 crystals within which the melt is distributed (Cashman et al., 2017; Maclennan, 2019). These
37 bodies cannot be erupted in their entirety due to their rheological properties, but mush fragments
38 can be disaggregated from the system by an ascending melt, and carried to the surface as
39 glomerocrysts, nodules or macrocrysts (Cashman et al., 2017). Evidence from mush fragments is
40 essential to constrain pressure (P), temperature (T), composition (X) and processes operating
41 within a plumbing system.

42 Magmas in Iceland commonly carry disaggregated fragments of crystal mushes (Cooper et
43 al., 2016; Halldórsson et al., 2018; Hansen and Grönvold, 2000; Neave et al., 2013, 2014, 2017;
44 Óskarsson et al., 2017; Passmore et al., 2012; Svavarsdóttir et al., 2017). Petrological investigations
45 of the geochemical relationship between macrocrysts and their carrier liquids have revealed that
46 some mush fragments are related to their carrier liquid (Neave et al., 2013), while others cannot
47 be cogenetic (Halldórsson et al., 2008). Identifying the location of a magma storage reservoir(s)

48 within a plumbing system and tracking its evolution with time is important for clarifying the
49 functioning of a volcanic system and for future eruption mitigation.

50 The Bárðarbunga-Veiðivötn volcanic system, located in the Eastern Volcanic Zone (EVZ), is
51 ideal for investigating the temporal and compositional evolution of a basaltic magma storage
52 reservoir(s) on the scale of a single volcanic system. This is because (1) it is one of the most active
53 volcanic systems in Iceland during the Holocene; (2) extrusives consist of phyrlic and ultra-phyric
54 rocks containing disaggregated crystal mush fragments (e.g., Halldórsson et al., 2018, 2008;
55 Hansen and Grönvold, 2000; Holness et al., 2019); and (3) Holocene tephtras from the Bárðarbunga
56 system (Óladóttir et al., 2011) exhibit distinctive compositional variations as a function of time
57 (Fig. 1). Furthermore, a fundamental observation from the Icelandic rift system is the apparent
58 effect of deglaciation on magma plumbing dynamics. This is well documented in the Reykjanes
59 Peninsula (Gee et al., 1998; Jakobsson et al., 1978), in the Western Volcanic Zone (Eason et al.,
60 2015; Sinton et al., 2005) and in the Northern Volcanic Zone (MacLennan et al., 2002; Slater et al.,
61 1998), although it remains unclear if and how magmatic plumbing systems were affected in the
62 EVZ.

63 In this work, we present mineral, groundmass glass and melt inclusion major and minor
64 elemental compositions from a temporally diverse (fully subglacial to historical) sample set from
65 the Bárðarbunga-Veiðivötn volcanic system in central Iceland. Each sample provides a snapshot of
66 the physical and chemical state of the volcanic system at the time of eruption. By linking
67 geochemical and petrological data with geothermobarometry calculations, we explore the
68 temporal evolution of magma storage conditions in the Bárðarbunga-Veiðivötn system. We first
69 chemically characterize a sample suite comprising fresh nodules, macrocrysts and glass. Secondly,
70 we employ a range of mineral-melt and melt-based thermobarometers (Hartley et al., 2018;
71 Neave et al., 2017; Putirka, 2008; Yang et al., 1996) to evaluate magma storage conditions for each

72 eruptive unit. On the basis of these observations, we make an attempt to reconstruct the
73 architecture of the Bárðarbunga-Veiðivötn magmatic system and the changes occurring within it
74 from the last glacial period to recent times.

75

76 **2. GEOLOGICAL SETTING AND SAMPLING**

77

78 **2.1 The Bárðarbunga-Veiðivötn volcanic system**

79 The Bárðarbunga volcanic system (Fig. 2) is the most extensive volcanic system in Iceland
80 with a total length of 190 km and an area of about 2500 km² (Thordarson and Höskuldsson, 2008).
81 With one eruption every 50 years in the last 1100 years, the Bárðarbunga volcanic system is one of
82 the most active system in Iceland (Larsen and Guðmundsson, 2014). The central edifice is split into
83 two subglacial volcanoes: Bárðarbunga, a ~2009 m-high caldera-bearing volcano situated under
84 the Vatnajökull ice cap, and Hamarinn, a smaller second central volcano located 20 km SW of
85 Bárðarbunga. The associated fissure swarm is commonly subdivided into two segments: the
86 Dyngjuháls fissure swarm extends 55 km north-northeast from Bárðarbunga into the Northern
87 Volcanic Zone, while the Veiðivötn fissure swarm extends 115 km southwest from Bárðarbunga
88 into the Eastern Volcanic Zone.

89 The southwest part of the Bárðarbunga volcanic system is commonly referred to as the
90 Bárðarbunga-Veiðivötn volcanic system. The Veiðivötn part consists of numerous well-developed
91 volcanic fissures orientated N45° (Larsen, 1984; Larsen and Guðmundsson, 2014; Thordarson and
92 Larsen, 2007). In the extreme southwest, the Veiðivötn fissure swarm propagates into the
93 Torfajökull volcanic system with production of both silicic and mixed products (Larsen, 1984;
94 Mørk, 1984; Zellmer et al., 2008). The latest eruption took place on the northern fissure swarm,
95 producing the 2014-15 Holuhraun lava (Pedersen et al., 2017).

96

97 **2.2 Sample description**

98 We have selected a suite of geologically well-characterised eruptive units from volcanic formations
99 situated in the Bárðarbunga-Veiðivötn volcanic system (Fig. 2 and Table 1).

100

101 *Ljósufjöll (Lj)*

102 Ljósufjöll, a subglacial volcanic ridge located within the Veiðivötn fissure system (Lj, Fig. 2). Studied
103 samples are from a glassy pillow lava, corresponding to Ljósufjöll formations b and c (lja and ljb)
104 described by Vilmundardóttir et al. (2000). Ljósufjöll is thought to have erupted early during the
105 last glacial period (Weichselian) and is therefore likely to be younger than 100 ka (Jóhannesson et
106 al., 1982). No prior petrochemical studies have been carried out on Ljósufjöll.

107

108 *Brandur (B), Fontur (F) and Saxi (S)*

109 Brandur, Fontur and Saxi are three early-Holocene tephra cones located to the east of
110 Þórisvatn lake (B, F, S, Fig. 2). Saxi and Fontur are aligned along a ~2.5 km-long linear fissure, while
111 Brandur is located 3 km west of the fissure on the edge of Þórisvatn lake. The craters consist of
112 unconsolidated, crystal-rich, fine-grained glassy material with plagioclase macrocrysts up to 4 cm
113 long, and abundant nodules of plagioclase, olivine and clinopyroxene (Halldórsson et al., 2008;
114 Hansen and Grönvold, 2000; Holness et al., 2007, 2019; Vilmundardóttir, 1977). These craters have
115 been suggested to be the source of the Þjórsárhraun lava (8.6 ka, e.g. Halldórsson et al., 2008;
116 Hansen and Grönvold, 2000; Hjartarson, 1988; Jakobsson, 1979). In addition to samples collected
117 for this study, we also collected data from samples previously studied by Hansen and Grönvold
118 (2000).

119

120 *Þjósárdalshraun (Th) and Drekahraun (Dr) (Tungnaá lava)*

121 Lava flows produced in the Veiðivötn fissure over the last 9 ka, but before the settlement of
122 Iceland in 874 AD, are collectively referred to as the Tungnaá lava sequence. At 45 km³, this is one
123 of the most voluminous lava sequences in Iceland (Vilmundardóttir, 1977). The source vents of the
124 Tungnaá lavas are now mostly buried by younger formations but were probably located in the
125 southern part of the Veiðivötn fissure swarm (Pinton et al., 2018). Þjósárdalshraun and
126 Drekahraun (Th and Dr, Fig. 2) are mid-Holocene lavas belonging to the Tungnaá sequence. They
127 are dated to between 3-4 ka BP by tephrochronology (Pinton et al. 2018). Drekahraun samples
128 consist of fresh and vesicular scoria collected west of Dreka vatn, near the lava source vents (Dr,
129 Fig. 2). The Þjósárdalshraun lava was largely emplaced to the north of Hekla volcano, flowing
130 westward following the Þjósá river and ultimately forming a field of rootless cones in Þjósárdalur
131 valley (Th, Fig. 2).

132

133 *Veiðivötn 1477 (V)*

134 The 1477 AD Veiðivötn eruption is the most recent eruption covered by our sample suite.
135 The eruption is considered to be the largest basaltic explosive eruption that has occurred in
136 Iceland in the last 1200 years (Thordarson and Larsen, 2007). It took place on a 65 km long fissure
137 and produced 5-10 km³ of highly fragmented basaltic tephra and small lava flows. This volume
138 includes both tephra and lavas ranging from basalt to rhyolite which erupted simultaneously as
139 the Veiðivötn magmas entered the Torfajökull silicic center to southwest (McGarvie, 1984; Mørk,
140 1984; Zellmer et al., 2008). Tephra from this eruption covered an area of 53 km² on land and has
141 been found as far afield as Ireland and Sweden (Larsen, 1984; Larsen and Guðmundsson, 2014).
142 Here we study fresh and glassy basaltic scoria from the central and southern part of the main
143 fissure specifically avoiding the mixed magmas near Torfajökull (V1477, Fig. 2).

144

145 **3. METHODS**

146

147 **3.1 Sample preparation, analytical and thermobarometry methods**

148 Thin sections of well-preserved and representative whole rock samples were made from each unit.
149 Plagioclase, olivine and clinopyroxene crystals (0.5-2.4 mm) were hand-picked from crushed
150 samples, mounted in epoxy resin and polished to expose glassy melt inclusions (MIs). Crystals
151 containing devitrified MIs were heated in a high-temperature furnace at 1210 ± 5 °C, which was
152 expected to exceed the crystallization temperature. The re-homogenized MIs were later exposed
153 at the surface. Major element compositions of macrocrysts ($n = 1530$), their host glass ($n = 328$)
154 and olivine- and plagioclase-hosted MIs ($n = 436$) were determined by electron microprobe (EPMA)
155 using a JEOL JXA-8230 SuperProbe at the University of Iceland. 1σ errors reported in this work are
156 based on multiple standard analyses collected during different analytical sessions. All melt
157 inclusion compositions have been corrected for the effect of post-entrapment crystallization (PEC)
158 on the inclusion walls.

159 We calculated magma storage temperatures based on glass compositions (Yang et al., 1996)
160 and mineral-glass pairs (Putirka, 2008). Crystallization pressures were calculated based on
161 clinopyroxene-melt pairs, following the method described by Neave and Putirka (2017), which has
162 a standard error of estimate (SEE) of ± 1.4 kbar, whereas the olivine-plagioclase-augite-melt
163 (OPAM) barometer (Hartley et al., 2018; Yang et al., 1996) was applied to estimate groundmass
164 glass and MI equilibration pressures ($SEE = \pm 1.3$ kbar). Full details of analytical methods,
165 homogenization experiments, PEC corrections and thermobarometry calculations are provided as
166 supplementary material (S1).

167

168 **4. Results**

169

170 **4.1 Petrography**

171 All our samples contain three main macrocryst (>500 μm) phases: olivine, clinopyroxene
172 and plagioclase. Minerals are present either as single grains scattered in the groundmass or in
173 polymineralic glomerocrysts (Fig. 3). Plagioclase is the most common mineral phase in all samples
174 (Fig. 3a). Plagioclase macrocrysts are generally euhedral and range from 500-6000 μm in size,
175 although crystals up to 3-4 cm are found in Brandur, Fontur and Saxi samples. Clinopyroxene
176 macrocrysts range between 500-1600 μm and often occur in glomerocrysts (Fig. 3b), although
177 large euhedral clinopyroxene is occasionally found (Fig. 3c). Olivine macrocrysts are typically 500-
178 2000 μm in size and are either euhedral or show rounded and resorbed habits (Fig. 3d-e). Cr-rich
179 spinel is sporadically found in the groundmass glass and is also widespread as inclusions in olivine
180 and plagioclase macrocrysts.

181 Naturally quenched melt inclusions are abundant in olivines and plagioclases from
182 Ljósufjöll, Brandur, Fontur, Saxi, and Veiðivötn 1477. The melt inclusions range in size from 10-150
183 μm (Fig. 3d-e). Plagioclase and olivine crystals from Þjórsárdalshraun and Drekahraun contain MIs
184 that are partially crystallized. Crystals in the tephra cones Brandur, Fontur and Saxi are surrounded
185 by a glassy to fine-grained matrix. Ljósufjöll samples display a coarse-grained groundmass (Fig. 3g)
186 composed of plagioclase, clinopyroxene, olivine and oxides, changing to a cryptocrystalline and
187 glassy matrix towards the pillow margins (Fig. 3h). Drekahraun and Veiðivötn 1477 samples
188 contain a fine-grained matrix with glassy portions at the tephra clast rims. Þjórsárdalshraun
189 samples have a holocrystalline groundmass.

190 Samples from all localities have plagioclase macrocrysts with cores exhibiting complex
191 internal textures. The inner part is either oscillatory or patchy zoned and always wrapped by
192 euhedral to subhedral rims. The thickness of Ljósufjöll plagioclase rims appears to be correlated
193 with the matrix texture (Fig. 3g-h). Plagioclase rims in contact with coarse-grained groundmass are
194 thicker (~30-70 μm) (Fig. 3g), while plagioclase rims scattered in cryptocrystalline to glassy matrix
195 are thinner (~10-40 μm) (Fig. 3h). Þjósárdalshraun plagioclase macrocrysts are normally zoned.

196 In backscattered electron (BSE) images, clinopyroxenes display bright and dark sectors (Fig.
197 3b). The clinopyroxenes are found either as glomerophytic clots or as fine to coarse intergrowths
198 of clinopyroxene and plagioclase forming next to plagioclase macrocrysts. In Veiðivötn 1477,
199 Ljósufjöll and the tephra cone samples, clinopyroxene also occurs as scattered single grains (Fig.
200 3c). Olivine crystals are often resorbed, especially in Brandur, Fontur and Saxi samples, although
201 euhedral crystals are found in all localities.

202 Abundant cm-size (up to 10 cm) olivine gabbro xenoliths are found in the Brandur, Fontur
203 and Saxi cones (Fig. 3e). The five studied nodules all contain 70-80 vol.% plagioclase in a subophitic
204 texture with some resorbed interstitial olivine and clinopyroxene. The framework is sustained by
205 transparent, light brown interstitial glass, which is locally crystallized to a fine intergrowth of
206 plagioclase, clinopyroxene \pm olivine (see also Hansen and Grönvold, 2000 and Holness et al., 2007).
207 Two xenoliths, 0.5-0.9 cm in size, were also found in the Þjósárdalshraun lava samples (Fig. 3f).
208 They consist of olivine-free gabbro, with plagioclase and clinopyroxene forming an ophitic texture,
209 and contain localized pockets of coarse- to fine-grained interstitial material (Fig. 3f).

210

211 **4.2 Mineral and glass chemistry**

212

213 Macrocryst compositions are summarized in Fig. 4a-c. For each mineral phase, variation diagrams
214 are also shown in Fig. 4d-f and in the supplementary material (Fig. S1.2-S1.4). Groundmass glass
215 and melt inclusion compositions are reported in Fig. 1 and Fig. 5. The full EPMA dataset is provided
216 as supplementary material (S2).

217

218 **4.1 Plagioclase**

219 Plagioclase macrocrysts commonly have bytownitic to anorthitic compositions. Ljósufjöll
220 plagioclase macrocryst cores display a narrow compositional range of An_{86-90.5}, whereas the
221 composition of the rims depends on the groundmass glass texture (Fig. 3g-h). Plagioclase rims in
222 contact with coarse-grained groundmass have more evolved compositions within the range An₆₅₋
223 ₇₁, while plagioclase rims adjacent to cryptocrystalline to glassy groundmass record compositions
224 within the range An₇₉₋₈₆ (Fig. 4a and 3d). Plagioclase macrocryst cores from Brandur, Fontur and
225 Saxi samples are in the range An_{83-91.5}, with Fontur plagioclase cores having slightly less
226 compositional variation of An_{86-90.5} (Fig. 4a). All plagioclase macrocrysts are surrounded by An₇₁₋₈₁
227 rims. Þjórsárdalshraun and Drekahraun plagioclase macrocryst core and rim compositions are
228 between An₈₄₋₉₁ and An₇₁₋₈₆, respectively (Fig. 4a and 4d), while plagioclases found in
229 Þjórsárdalshraun nodules are more homogeneous (An_{85-89.5}). Macrocryst cores and rims from
230 Veiðivötn 1477 display the largest compositional variation of all studied localities. Plagioclase core
231 compositions are An₇₈₋₉₁, while rims are An₆₆₋₇₇.

232

233 **4.2 Clinopyroxene**

234 Clinopyroxenes have augitic compositions. Ljósufjöll clinopyroxene cores and dark sectors
235 in BSE images have compositions of Mg# 84-87 ($Mg\# = [(MgO_{mol}) / (MgO_{mol} + FeO_{mol}^{tot})] * 100$), while
236 rims and bright sectors have Mg# 76-86 (Fig. 4b and 3e). Brandur, Fontur and Saxi cones contain

237 clinopyroxene with cores and dark sectors in the range Mg# 78-87 and rims and bright sectors in
238 the range Mg# 75-84 (Fig. 4b). Þjórsárdalshraun clinopyroxenes have Mg# 76.6-84.7, while
239 clinopyroxenes found in the nodules are more homogeneous, being in the range Mg# 83-85.
240 Drekahraun clinopyroxene cores and dark sectors range between Mg# 81.7-85, while rims and
241 bright sectors are in the range Mg# 79-84. Here, many reversely zoned clinopyroxenes occur with
242 Fe-rich cores (Mg# 70-72) surrounded by Fe-poor sectors (Mg# 82-84) (Fig. 4b and 3e).
243 Clinopyroxene dark sectors found in Veiðivötn 1477 samples vary in the range Mg# 79-85 and
244 bright sectors in the range Mg# 75.7-82. A few clinopyroxene crystals, occurring as large single
245 grains, are normally zoned with large Mg-rich cores of Mg# 84-85 overgrown by rims with Mg# 81-
246 82. Reversely zoned clinopyroxenes contain Fe-rich cores (Mg# 67-75.4) followed by oscillatory
247 zoning.

248 Clinopyroxene crystals can be strongly sector zoned, which is mostly reflected in their Ca,
249 Al and Ti contents (e.g.: Nakamura, 1973; Ubide et al., 2019). In order to minimise this
250 compositional effect, we plotted the $\text{Al}_2\text{O}_3/\text{TiO}_2$ vs Mg# for all clinopyroxene analyses (Fig. 4e). In
251 general, the $\text{Al}_2\text{O}_3/\text{TiO}_2$ increases with increasing Mg# and the $\text{Al}_2\text{O}_3/\text{TiO}_2$ variation is greater for
252 clinopyroxene with Mg# >83. Crystals with Mg# >83 show an $\text{Al}_2\text{O}_3/\text{TiO}_2$ span of 1.73 (1σ), in
253 contrast to clinopyroxenes with Mg# <83 where this span is only 0.97 (1σ). Ljósufjöll and tephra
254 cone clinopyroxenes record the largest dispersion, while recent samples have a narrower range. In
255 fact, middle-Holocene units and historical units do sample primitive clinopyroxene in terms of Mg#
256 and they register relatively low $\text{Al}_2\text{O}_3/\text{TiO}_2$ ratios and Cr_2O_3 contents. Indeed, Cr-rich ($\text{Cr}_2\text{O}_3 >0.8$
257 wt%) clinopyroxenes are exclusively sampled by the old units (Fig. S1.3c).

258

259

260 **4.3 Olivine**

261 In Ljósufjöll samples, unzoned olivine macrocrysts vary in composition from Fo_{84} to Fo_{87} .
262 Zoned olivines, on the other hand, have core and rim compositions of $\text{Fo}_{82.5-87}$ and $\text{Fo}_{71-77.5}$,
263 respectively (Fig. 4c and 3f). Olivine macrocrysts from Brandur, Fontur and Saxi have
264 homogeneous core compositions of $\text{Fo}_{82-87.5}$ and rims between Fo_{76-82} (Fig. 4c). Olivine cores from
265 Þjórsárdalshraun and Drekahraun vary in the range $\text{Fo}_{79.5-86}$ and $\text{Fo}_{82.4-86}$, respectively.
266 Þjórsárdalshraun olivine rims have compositions within the range $\text{Fo}_{73.6-79}$, while Drekahraun
267 olivine rims have compositions of Fo_{78-83} . Reversely zoned olivines, mostly occurring in Drekahraun
268 samples, have cores of $\text{Fo}_{66.5-82}$ encased by more primitive Fo_{80-82} rims (Fig. 4c and 3f). Sparse
269 olivines in Veiðivötn 1477 samples show a wide compositional range of cores and rims, between
270 Fo_{77-87} and $\text{Fo}_{75.4-85.5}$, respectively.

271 Olivine variation diagrams for all locations are shown in Fig. 4f and Fig. S1.4. Olivine cores
272 have NiO contents between 0.1 and 0.25 wt.% that decrease to 0.05 wt.% in the rims (Fig. 4f).
273 Fe/Mn ratio, diagnostic of parental magma compositional differences (e.g., Sobolev et al., 2007),
274 varies between 51 and 86 (Fig. S1.4a), with more variation observed in the most primitive crystals
275 ($\text{Fo} > 85$, $1\sigma=5.2$) compared to olivines with $\text{Fo} < 85$ ($1\sigma=4.3$).

276

277 **4.4 Groundmass glass**

278 Groundmass glass composition varies as a function of time (Fig. 1). From early-Holocene till
279 present, carrier melts become more evolved. Indeed, Ljósufjöll groundmass glass is the most
280 primitive (Fig. 1 and 5) with Mg# 57-60, MgO 7.5-9.3 wt% and TiO_2 0.95-1.1 wt%, being one of the
281 most primitive tholeiite glass compositions known from the EVZ (see Hansen and Grönvold, 2000;
282 Neave et al., 2014, 2017; Óladóttir et al., 2011; Passmore et al., 2012). Drekahraun and Veiðivötn
283 1477 groundmass glasses show a tight compositional range (Fig. 1 and 5). Drekahraun glass (Mg#
284 50.5-55) has MgO 7-7.5 wt% and TiO_2 1.6-1.8 wt%, whereas Veiðivötn 1477 groundmass glass

285 (Mg# 45.7-49.8) contains MgO 6-6.9 wt% and TiO₂ 1.7-2 wt%. The groundmass of samples from
286 Þjórsárdalshraun, assumed to represent melt compositions, refer to fine-grained pockets found in
287 the nodules. Þjórsárdalshraun nodule glass (Fig. 2f) displays a large chemical variability (Fig. 5)
288 (MgO 5.7-7.5 wt% and TiO₂ 1.9-2.3 wt%), perhaps due to microcrystals. The compositional
289 variation of groundmass glass from the tephra cones (B-F-S) is relatively large in comparison to
290 other localities (Fig. 1 and 5) and our data are in good agreement with previously published data
291 (Hansen and Grönvold, 2000). We note that Brandur generally has more primitive glass (Mg# 46-
292 53, MgO 6.1-7.5 wt.%) than Fontur and Saxi (Mg# 40-49 and 41.5-50.5, MgO 5.5-7.5 wt% and 5.5-7
293 wt%, respectively).

294

295 **4.5 Melt inclusions**

296 The majority of MIs from all localities, corrected for post-entrapment processes (S1), form
297 a group with Mg# 58-68 (Fig. 5a-d). The only locality where MI and interstitial glass compositions
298 overlap is Ljósufjöll. Ljósufjöll MIs have MgO between 8.0-9.7 wt%, while other localities record a
299 wider range (MgO 6.2-10.0 wt%). Among the most primitive melt inclusions (Mg# >65, MgO 8.5-
300 10.0 wt%, n=24), two are hosted in plagioclases (An₈₆₋₈₉) from Drekahraun and Þjórsárdalshraun
301 (MgO of MIs 8.5-8.7 wt%), one is a plagioclase-hosted (An₉₀) MI from Brandur (MgO 10.0 wt%),
302 and the other 21 MIs are hosted in olivines (Fo_{86.5-88}) from the tephra cones (MgO 9.3-10.0 wt%).

303 Evolved MIs (Mg# <55, n=56,) are widespread in all eruptive units except Ljósufjöll. They
304 are hosted in both plagioclase (An₈₃₋₈₈) and olivine (Fo₇₆₋₈₀). The groundmass glass and MI
305 variations of our dataset are found to be in excellent agreement with published whole rock and
306 glass compositions (Halldórsson et al., 2008, 2018; Hartley et al., 2018; Jakobsson, 1979; Óladóttir
307 et al., 2011; Svavarsdóttir et al., 2017) from Bárðarbunga volcanic system (pale blue fields in Fig.

308 5), although our samples do not include primitive MIs with Mg# as high as 71 as found in
309 Holuhraun samples (Bali et al., 2018; Hartley et al., 2018).

310

311 **4.6 Macrocryst compositions and mineral-melt equilibrium**

312 Figure 4a-c shows macrocryst compositional ranges, along with mineral compositions
313 calculated to be in equilibrium with the observed groundmass glass compositions, shown with
314 coloured bands. Equilibrium olivine compositions were calculated using a fixed Kd_{Fe-Mg}^{ol-liq} of 0.3,
315 following Roeder and Emslie (1970), and equilibrium clinopyroxene compositions following the
316 model of Wood and Blundy (1997). Equilibrium plagioclase compositions were calculated using
317 equation 33 from the model of Namur et al. (2011).

318 Plagioclase, olivine and clinopyroxene macrocryst rims are generally found to be close to
319 the compositions predicted to be in equilibrium with the groundmass glass, with the exception of
320 Ljósufjöll macrocryst rims. Macrocryst cores are always too primitive to be in equilibrium with
321 groundmass glass compositions. This feature is fairly common in mush-bearing magmas and has
322 been observed in other eruptive units from the EVZ (Halldórsson et al., 2008, 2018; Neave et al.,
323 2013, 2014, 2015).

324

325 **5. Geothermobarometry results**

326 **5.1 Clinopyroxene storage pressures and groundmass equilibration pressures**

327

328 The relative probability of clinopyroxene crystallization pressures are shown as kernel
329 density estimates (KDE) in Fig. 6 a-d. Clinopyroxenes from all eruptive units give comparable
330 crystallization pressure ranges of 0.5-4.5 kbar (Table 2 and Fig. 6). Each locality returns a well-
331 defined peak in the KDE (Fig. 6a-d), located at ~2 kbar. The mean calculated pressure across all

332 units is 2.2 ± 0.7 (1σ) kbar. Only a few clinopyroxenes ($n=14$, Mg#78-85) – the majority of which
333 are from Drekahraun and the tephra cone localities – return pressures higher than 3.5 kbar.
334 Assuming an average Icelandic crustal density of 2.86 g/cm^3 (Carlson and Herrick, 1990), our data
335 indicate a mid-crustal magma storage zone located at 7.8 ± 2.5 (1σ) km (Fig. 6 a-d).

336 In Fig. 6a-d, we plot KDEs of equilibration pressures calculated for both groundmass glasses
337 and melt inclusions. Groundmass glasses return a mean of 1.9 ± 0.8 (1σ) kbar, which is statistically
338 indistinguishable from our calculated clinopyroxene-liquid pressures (Table 2). The majority of
339 groundmass glasses are within the range 0.4–3.0 kbar. Similar equilibrium pressures are obtained
340 for the Bárðarbunga Holocene tephras (Óladóttir et al., 2011a), with a mean pressure of 2.6 ± 0.4
341 (1σ) kbar (Fig. 6). Therefore, although the studied carrier melts have distinct and variable chemical
342 compositions, they all last equilibrated with olivine, plagioclase and augite at essentially the same
343 depth (Fig. 6a-d), in a mid-crustal reservoir located at 6.8 ± 2.8 (1σ) km. Barometry calculations
344 carried out on 2014-15 Holuhraun samples reveal a magma storage zone located at about 7-8 km
345 depths, consistent with geophysical observations (Halldórsson et al., 2018; Hartley et al., 2018).

346

347 **5.2 OPAM melt inclusion equilibration pressures**

348 Out of 436 olivine and plagioclase-hosted MIs, 299 inclusions return probability fits >0.8
349 (table 2 and Fig. 6a-d). The oldest, subglacial unit Ljósufjöll, records a fairly large range of MI
350 trapping pressures, with broad peak at 3.3 kbar and a tail up to 6 kbar. Inclusions from the tephra
351 cones display a bimodal distribution with one peak at 3.0 kbar and another at 4.9 kbar. The high-
352 pressure peak is well defined for Brandur, Fontur and Saxi samples, with 108 MIs recording
353 pressures higher than 4.0 kbar, but not statistically significant for Ljósufjöll inclusions due to the
354 low number of samples (11 MIs show pressures between 4.0-6.0 kbar). Þjórsárdalshraun and
355 Drekahraun return MI equilibration pressures in the range 0.6-4.5 kbar, with a main peak at 2.7

356 kbar and a minor peak at 1.9 kbar. MI in Veiðivötn 1477 samples show a bimodal distribution but
357 the probability distribution is not well defined due to the small number of inclusions (n=28). The
358 calculated pressures range between 1.2-4.2 kbar with a most common equilibration pressure at
359 3.6 kbar and a second peak at 1.8 kbar.

360 We have explored the relationship between MI pressures and the composition of the host
361 crystals in all units (Fig. 7). In the subglacial unit (Fig. 7a), MIs hosted in Fo_{~86} olivine crystals record
362 a most probable peak at around 2.6 kbar (9.3 km), while plagioclase-hosted MIs show a main peak
363 at 3.9 kbar (13.9 km) and multiple secondary peaks, with the equilibration pressures up to a
364 maximum of 6.0 kbar (21.4 km). Brandur, Fontur and Saxi cones record an even larger range of MI
365 equilibration pressures (0.4-7.6 kbar) (Fig. 7b). Plagioclase-hosted MIs were entrapped within the
366 pressure range 3.5-7.6 kbar (12.5-27 km) at a most probable pressure of 4.9 kbar (17.5 km).
367 Olivine-hosted MIs were trapped at pressures within the range 0.4-7.0 kbar (1.5-25 km). MI
368 equilibration pressures increase with the forsterite content of the host olivine, which is mainly
369 noticeable in Fig. 7b, where we distinguish two different MI populations: (1) MIs trapped in Fo₇₆₋₈₆
370 crystals, with a most probable pressure of 2.8 kbar (10 km) and (2) MIs hosted in Fo₈₆₋₈₈ crystals,
371 which produce the high-pressure tail of the distribution, with secondary peaks at 4.3 kbar (15.3
372 km) and 6 kbar (21.4 km). Macrocrysts from Brandur, Fontur and Saxi were previously studied by
373 Hansen and Grönvold (2000). They concluded that macrophenocrysts crystallized between 7 and
374 40 km depth (2-11 kbar). Finally, middle-Holocene and historical units have a narrow range of MI
375 trapping pressures (Fig. 7c), of 0.7-4.3 kbar (2.5-15 km). Olivine-hosted MIs crystallized at a most
376 probable pressure range of 1.9-2.6 kbar (6.8-9.2 km), while plagioclase-hosted MI equilibration
377 pressures are more variable, with a main peak at 3.3 kbar (11.7 km).

378 One might argue that the highest pressures calculated for the primitive melt inclusions are
379 potentially not valid as, despite the numerical filtering, these melts might not be saturated in

380 clinopyroxene. Using Eq. 35 of Putirka (2008), we have calculated equilibrium melt compositions
381 for the most primitive clinopyroxene macrocrysts observed in our crystal cargo (i.e. Mg#₈₅₋₈₇). We
382 find that silicate melts with Mg#_{melt} = 61-65 are in equilibrium with Mg# 85-87 clinopyroxenes,
383 suggesting that only inclusions with Mg#>65 should be treated with caution (n=17, circles with
384 black thick outline in fig. 7b and 7c). Furthermore, model calculations with Petrolog (discussed in
385 detail later) suggest that clinopyroxene with Mg# ~88 will be on the liquidus of Mg# ~65 melts at
386 pressures of 2-6 kbar.

387

388 **5.3 Geothermometry**

389 KDEs for calculated temperatures are illustrated in Fig. 6e-h and mean temperature values
390 are reported in Table 3. There is little variation in melt temperature between the samples,
391 although there is some indication that the Ljósufjöll carrier melt was hotter than the carrier liquids
392 in the other eruptions. Calculated temperatures for Ljósufjöll groundmass glass range between
393 1185 and 1210 °C (mean at 1193 ± 4 (1 σ) °C), while samples from tephra cones, Tungnaá lava and
394 Veiðivötn 1477 give similarly lower temperatures, with a mean at 1165 ± 7 (1 σ) °C, 1170 ± 8 (1 σ) °C
395 and 1167 ± 3 (1 σ) °C, respectively. Furthermore, Ljósufjöll groundmass glass temperatures are
396 statistically indistinguishable from those derived from its melt inclusions. Across all samples, the
397 recovered carrier melt temperatures are within the $\pm 26^\circ\text{C}$ SEE of the thermometer (Eq. 16,
398 Putirka, 2008), but are sufficiently different to suggest that during the last glacial period, melts
399 erupted in the Bárðarbunga system were hotter than recent carrier magmas.

400 The clinopyroxene-melt thermometer returns wider temperature variations within
401 individual samples (Fig. 6e-h), but the sample average crystallization temperatures are remarkably
402 consistent with an overall mean crystallization temperature of 1188 ± 17 °C. We find no significant
403 temporal variation in crystallization temperature. Finally, the plagioclase-melt and olivine-melt

404 thermometers applied to MIs return temperatures with a narrowly focused peak at 1214 ± 10 (1σ)
405 °C, regardless of age.

406

407 **6. Discussion**

408 **6.1 Modelling fractional crystallization**

409 The glass compositional trends in Fig. 5a-b could, to a first order, be largely controlled by
410 fractional crystallization, although the complete variation in groundmass glass and MI major
411 element compositions is difficult to explain solely with fractional crystallization along a single
412 liquid line of descent. Results from our thermobarometric calculations suggest polybaric
413 crystallisation (Fig. 6). Therefore, we calculated liquid lines of descent (LLDs) from the average
414 composition of the most primitive melt inclusions for the whole dataset ($n=24$, Mg# >65) at
415 different pressures of 0.001, 2, 4 and 6 kbar. The starting composition has Mg# of ~ 66 , TiO₂ 0.97
416 wt%, MgO 9.5 wt% and CaO 13.4 wt% (white stars, Fig. 5a, b). Models were run using the
417 Petrolog3 software (Danyushevsky and Plechov, 2011), applying the pressure-sensitive mineral-
418 melt model of Ariskin et al. (1993). Oxygen fugacity was set at the QFM buffer, assuming similar
419 oxidation conditions to those measured in the most recent eruption of the Bárðarbunga volcano
420 (Bali et al., 2018; Halldórsson et al., 2018).

421 A similar approach was used for LLDs in Fig. 5 c-d, which show Al₂O₃/TiO₂ and TiO₂/K₂O as a
422 function of Mg#. Both diagrams show that the variability in these oxide ratios decreases with
423 decreasing Mg#. Al₂O₃/TiO₂ in melt inclusions varies between 6 and 37 ($1\sigma= 4.3$) (Fig. 5c), while in
424 groundmass glasses, it is between 5 and 15 ($1\sigma= 2.1$), with the Ljósufjöll carrier liquid having the
425 highest Al₂O₃/TiO₂ (~ 15). A similar diversity is also observed in TiO₂/K₂O (Fig. 5d), with primitive
426 melts recording the largest spread. Therefore, we modelled fractional crystallization at 2 kbar
427 considering different starting compositions to encompass the observed diversity in primitive melt

428 compositions (red stars in Fig. 5c, d). Out of the most primitive MIs (Mg# >60), we averaged MIs
429 with high and low Al₂O₃/TiO₂, ending up with (1) a melt composition with high Al₂O₃/TiO₂ (28.1 ±
430 3.3 (1σ)) and (2) a melt with low Al₂O₃/TiO₂ (9.5 ± 0.4 (1σ)).

431 Although the trend of CaO vs. Mg# (Fig. 5a) in the glasses is roughly covered by polybaric
432 fractional crystallisation, variation in TiO₂ (Fig. 5b) and oxide ratios such as TiO₂/K₂O (Fig. 5 c, d)
433 cannot be explained by simple fractional crystallization of a single parental melt composition.
434 Thus, regardless of the model and the starting composition adopted, a single LLD cannot
435 reproduce the observed chemical variability. Therefore, we suggest that neither isobaric nor
436 polybaric fractional crystallization alone is sufficient to describe the observed glass composition
437 variability.

438

439 **6.2 Evidence for concurrent mixing and crystallization**

440 Element ratio variability in olivine and clinopyroxene macrocrysts and MIs provides
441 evidence for the occurrence of diverse primary melts in the Bárðarbunga-Veiðivötn plumbing
442 system. Clinopyroxene records a notable decrease in the variability of Al₂O₃/TiO₂ (Fig. 4e) and
443 Cr₂O₃ (Fig. S1.3c) as clinopyroxene Mg# decreases. A similar behaviour is observed for olivine,
444 where there is a much greater spread in Fe/Mn among Fo-rich crystals than Fo-poor crystals (Fig.
445 S1.4a). Furthermore, we observe comparable trends in the oxide ratios of groundmass glasses and
446 melt inclusions (e.g., Al₂O₃/TiO₂ and TiO₂/K₂O in Fig. 5c and Fig. 5d, respectively), with melt
447 compositions at an early stage of magmatic evolution showing the greatest variability. This
448 behaviour has been observed in samples from several Icelandic eruptions with regards to trace
449 elements (e.g., Hartley et al., 2018; Maclennan et al., 2003; Maclennan, 2008; Neave et al., 2013).

450 In order to explain such variability at an early stage of magma history, we must invoke
451 heterogeneities in the mantle source or in the melting process. As primary melts form, they start

452 their history of magmatic evolution and progress through storage, crystallization and mixing
453 (Maclennan, 2008). The compositional variation of primitive crystals (Fig. 4e and Fig. S1.3c) and
454 their melt inclusions (Fig. 5c) has not been preserved with melt evolution, testifying that mixing
455 and compositional homogenization has occurred between MI entrapment and eruption. In our
456 samples, the most primitive MIs, hosted in olivine crystals of Fo~88, have Mg# ~67 and we do not
457 observe olivine crystals of Fo >88. We propose that our samples do not preserve near-primary
458 mantle-derived melts and most likely we are missing the earliest stages of the crystallization story.
459 The most primitive melts are therefore likely to have had even more variable compositions than
460 those preserved in our samples.

461 We compared our data with melt inclusion compositions from elsewhere in Iceland's EVZ.
462 Figure 8 shows melt inclusion compositions from the 1783 AD Laki eruption (Neave et al., 2013),
463 Skuggafjöll subglacial eruption (Neave et al., 2014), the 10 ka Grímsvötn tephra series (Neave et
464 al., 2015) and the recent 2014-15 Holuhraun eruption (Bali et al., 2018). Naturally quenched MIs
465 from Laki are fairly evolved with Mg# extending to much lower values (Mg# 32-60) and a very
466 narrow range of SiO₂/TiO₂ (Fig. 8), and seem to follow a slightly different trend from our data.
467 Conversely, melt inclusions from the 10 ka Grímsvötn (MgO up to 10.5 wt%) and Skuggafjöll (MgO
468 up to 10.3 wt%) samples show large variations in SiO₂/TiO₂, closely matching the most primitive
469 compositions reported in this work. Finally, melt inclusions from Holuhraun span the
470 compositional variation recorded by our data (Fig. 8), although more primitive melts (Mg#~71),
471 relative to our dataset, are captured by the Holuhraun melt inclusion record.

472 In summary, major and minor elements of macrocrysts, as well as melt inclusions, preserve
473 evidence for compositionally diverse parental melts that might reflect heterogeneities in the
474 mantle source or different degrees of partial melting (Maclennan, 2008). This needs to be further
475 investigated by trace elements and stable and radiogenic isotopes. However, the compositional

476 variability decreases as mixing and fractional crystallization progresses, producing the magma
477 composition documented by the carrier melt.

478

479 **6.3 Characteristics of the crystal cargo**

480 In the case of Ljósufjöll macrocrysts, equilibrium with the carrier liquid is exclusively
481 registered by plagioclase with intermediate compositions (An_{79-86}), although olivine cores are close
482 to equilibrium (Fig. 4). We suggest that macrocrysts incorporated within the glassy groundmass
483 are representative of the original cargo and that the macrocrysts scattered within the coarse-
484 grained groundmass would have experienced post-emplacement crystallization of the outermost
485 rims (An_{65-71}). The fact that Ljósufjöll samples come from a pillow lava, whose interior had longer
486 time to cool and evolve, supports the chemical evidence for the two macrocrysts types.

487 In all units, macrocryst cores could not have crystallized from the respective carrier melts
488 (Fig. 4a-c). In Fig. 9a-b, we show KDEs (coloured areas) for plagioclase and olivine macrocryst core
489 compositions against time. We also report the most primitive mineral compositions (vertical bars),
490 which are calculated to crystallize from the most primitive melt inclusions from each magmatic
491 unit. Finally, we show a comparison of olivine and plagioclase macrocryst core compositions
492 (dotted curves) in samples of subglacial (Skuggafjöll; Neave et al. 2014), early Holocene (10 ka
493 Grímsvötn; Neave et al. 2015) and historical (1783 Laki and 2014-2015 Holuhraun eruptions;
494 Neave et al. 2013, Halldórsson et al., 2018) eruptions that all took place in central Iceland.

495 Macrocryst cores are close to the predicted equilibrium compositions with melt inclusions
496 (Fig. 9a-b), and MIs appear to be in equilibrium with plagioclase, clinopyroxene and olivine with
497 maximum values of $An \sim 86-87$, $Mg\# \sim 85-86$ (not shown) and $Fo \sim 86-87$, respectively. As a result,
498 high-MgO melts represented by the most primitive melt inclusions could have crystallized the
499 majority of clinopyroxene and olivine core compositions acquired in this work, although this does

500 not apply to plagioclases with the highest An. Plagioclase macrocrysts with $An > 87$ are widespread
501 at all localities, and must have crystallized from melts that are not preserved in our petrological
502 record.

503 Ljósufjöll has a very restricted range of plagioclase macrocryst compositions (An_{87-91}). The
504 range of plagioclase compositions expands with time: for younger eruptive units the plagioclases
505 are skewed towards more evolved compositions (Fig. 9a), and high-An cores are rare in the
506 middle-Holocene and historical units. Similar relationships are observed for olivine, with the range
507 of core compositions extending to lower Fo with time (Fig. 9b). Comparing our results with crystal
508 compositions from the 2014-2015 Holuhraun eruption (Halldórsson et al., 2018), we find that the
509 Holuhraun cargo records a wider distribution of plagioclase macrocryst compositions (An_{69-92})
510 compared to Veiðivötn 1477, but a similar distribution of olivine macrocryst compositions (Fo_{76-88}).

511 A similar variation of macrocryst compositions is observed when we compare available
512 samples within the EVZ of similar age (Fig. 9a-b). We find that the subglacial Skuggafjöll (Neave et
513 al., 2014) and 10 ka Grímsvötn (Neave et al., 2015) samples record a narrow range in plagioclase
514 core compositions (An_{83-92}), while historical eruptions such as Laki have much more variable
515 plagioclase compositions (An_{65-90}) (Neave et al., 2013). Interestingly, this partially also applies to
516 olivine macrocrysts. The subglacial Skuggafjöll eruption products have a restricted olivine
517 compositional range (Fo_{84-87}), but olivines from the 10 ka Grímsvötn show a broader and bimodal
518 (Fo_{69-77} and Fo_{83-87}) compositional distribution. The 1783 Laki eruption has several olivine
519 populations in the range Fo_{67-87} . However, the compositional range in Laki crystal cargo might be
520 affected by availability of a much larger dataset compared to the other eruptions. Taken together,
521 these data suggest that more evolved crystals have been erupted with time, both within the scale
522 of a single volcanic system and, potentially, on the scale of the EVZ and central Iceland.

523

524 **6.4 Assessing the temporal variability of the Bárðarbunga-Veiðivötn volcanic** 525 **system**

526

527 In the case of Iceland in the postglacial period (<12 ka), the crust has been affected by isostatic
528 adjustments (Sigmundsson, 1991) due to ice removal and glacial rebound effects (Le Breton et al.,
529 2016), with magma eruption rates 20-30 times higher than at present day (Maclennan et al., 2002;
530 Sigvaldason et al., 1992). Modelling studies (Eksinhol et al., 2019; Jull and McKenzie, 1996) and
531 chemical constraints (Eason et al., 2015; Gee et al., 1998; Hardarson and Fitton, 1991; Maclennan
532 et al., 2002; Sinton et al., 2005; Slater et al., 1998) have provided evidence to link this eruption
533 pulse

534 either to the release of pooled magma enhanced by a change of the stress field in the crust
535 (Gudmundsson, 1986; Sigvaldason et al., 1992) or to an increase in the decompression melting
536 rate of the mantle (Jull and McKenzie, 1996; Maclennan et al., 2002) caused by the unloading of
537 the ice cap above Iceland.

538 The evolution of the Bárðarbunga-Veiðivötn volcanic system in the late Pleistocene and Holocene
539 can likely be explained within this framework and we propose three different stages (Fig. 10):

540 (1) A steady-state glacial stage when a 2000 m-thick ice cap (Sigmundsson, 1991) pressed
541 down the crust, sampled by the subglacial unit (Fig. 10a). In this period the magmatic system was
542 characterized by magma storage regions distributed over a large crustal interval of ~5-21 km (Fig.
543 10a). The accuracy of the OPAM barometer does not allow us to resolve the vertical arrangement
544 of the deeper storage zones (i.e., multiple stacked sills) in the mid- to lower crust (Kelemen et al.,
545 1997). In this period, the crystal-mush system was characterized by highly primitive macrocrysts
546 that were picked up by a primitive and homogeneous carrier liquid.

547 (2) A changing state of the magmatic system, when decompression and glacial rebound
548 effects associated with ice unloading occurred (Jull and McKenzie, 1996; Le Breton et al., 2016;
549 Maclennan et al., 2002; Slater et al., 1998), represented by the early-Holocene tephra cones (Fig.
550 10b). Our results (Fig. 7) on Brandur, Fontur and Saxi samples can be interpreted as a result of an
551 increase in magma production rates (Jull and McKenzie, 1996; Maclennan et al., 2002) that
552 promoted the input of primitive melts from the mantle and crystallization of primitive macrocrysts
553 in a deep-seated storage zone(s). We find that crystallization of primitive olivine ($Fo_{>85-86}$) and
554 plagioclase ($An_{>85}$) macrocryst cores took place at a mid- to lower-crustal depths of 15-22 km (Fig.
555 10b). Some of these macrocrysts were transported upwards into mid-crustal reservoir(s) (7-11
556 km), where more evolved phases also crystallized. Furthermore, although it is still unresolved how
557 deglaciation might have affected the chemistry of erupted melts, it is clear that early postglacial
558 products record a greater variability in MgO contents, and are depleted in incompatible elements
559 compared with lavas erupted when the ice load was thought to be relatively stable or absent
560 (Eason et al., 2015; Gee et al., 1998; Jull and McKenzie, 1996; Maclennan et al., 2002; Sinton et al.,
561 2005; Slater et al., 1998). This is consistent with our observation of a larger spread in the MgO
562 contents of carrier liquids erupted in the early Holocene (Fig. 1).

563 (3) A steady-state stage, which we observe at the present, with the magmatic system being
564 unaffected by short-term ice unloading effects, sampled by middle-Holocene and historical units
565 (Fig. 10c). MI pressures likely indicate comparable crystallization depths of olivine and plagioclase
566 macrocrysts in a storage zone(s) in the mid-crust. MI equilibration pressures estimated for the
567 2014-2015 Holuhraun eruption (Hartley et al., 2018) are all below 5 kbar (dotted KDE in fig. 7c),
568 with the most probable pressure at 3.2 kbar (11.4 km), which is consistent with our data. We infer
569 that since the middle-Holocene, magmas erupted in the Bárðarbunga-Veiðivötn system have

570 mainly carried evolved olivine and plagioclase macrocrysts (Fig. 9) that were stored at mid-crustal
571 depths around 7-13 km (Fig. 10c).

572 All these clues rule out the involvement of a deep reservoir during historical eruptions, at
573 least as a direct source for extrusives. The absence of a deep signature can be interpreted as a
574 result of the re-establishment of a new pressure equilibrium. Shallow storage region(s) may then
575 dominate the plumbing architecture. As seen at Etna volcano, passageways of melts in a magmatic
576 system can change with time (Kahl et al., 2013). From this perspective, the deep reservoir(s) might
577 be bypassed by melts coming from the mantle in historical time, most likely because new and
578 stable magma pathway has been established. Deep earthquake swarms (~22 km) beneath the
579 Bárðarbunga system, which have been associated with deep melt injections and melt movement
580 (Hudson et al., 2017) are also of importance here. However, the 2014-15 Holuhraun eruption
581 products preserve no petrologic record of crystals or melt inclusions from this depth. Our
582 combined data suggest that since the middle-Holocene, crystals and melts from the deep storage
583 zone have not been directly transferred to the surface. This is perhaps due to a more effective
584 homogenisation within the mid-crustal storage zone(s). In contrast, the deep reservoir(s) was
585 clearly sampled in the early Holocene and during subglacial eruptions, indicating that during this
586 period the magmatic system had a different architecture that permitted direct ascent of magma
587 from deeper storage regions to the surface.

588

589 **7. Conclusions**

590

- 591 1. Studied samples contain evidence of interaction with a crystal mush reservoir(s)
592 and entrainment of crystal mush fragments.

593

594 2. Olivine and plagioclase macrocryst compositions vary with time. The older
595 formations are dominated by primitive crystals, whereas in the mid-Holocene to historic
596 formations their chemistry is skewed towards more evolved compositions.

597

598 3. Macrocryst cores and primitive melt inclusions exhibit a great chemical variability,
599 which is likely linked to compositionally heterogeneous primary melts. The decrease of this
600 variability, highlighted by the carrier liquid and macrocryst compositions, provides
601 evidence for concurrent mixing and crystallization of compositionally diverse melts within
602 the system.

603

604 4. Clinopyroxene-melt and OPAM barometries return a temporally consistent
605 crystallization pressure of 2.2 ± 0.7 (1σ) kbar and 1.9 ± 0.8 (1σ) kbar, corresponding to a
606 depth of 7.3 ± 2.7 (1σ) km, which are consistent with a relatively shallow reservoir located
607 in the middle crust. These estimates are in perfect agreement with petrological and
608 geophysical results obtained for 2014-15 Holuhraun eruption.

609

610 5. The subglacial and early-Holocene formations preserve a crystal cargo that
611 originated from a deep reservoir with An- and Fo-rich macrocrysts crystallized at about
612 17.5 km (4.9 kbar) depth. In contrast, mid-Holocene to historical samples record only
613 shallower crystallization pressures of 2-4 kbar (7-13 km). It appears that since the middle-
614 Holocene no crystals from the deep storage zone are transferred directly to the surface,
615 which could indicate the establishment of new magma pathways and more complete
616 homogenization of melts in the shallow storage zone during this period.

617

618 6. At the end of the last glacial maximum, the isostatic rebound caused a significant
619 change in the stress field of the crust and an increase in the melting rate in the upper
620 mantle, triggering a large pulse in magmatic activity (Le Breton et al., 2016; Maclennan et
621 al., 2002; Slater et al., 1998). Following this, during the early-Holocene, the lower crust was
622 continuously supplied with fresh batches of melt that allowed magma from deep regions to
623 erupt at the surface. Once crustal pressure equilibrium has been re-established following
624 the early-Holocene, shallower storage levels dominated the system architecture with more
625 efficient mixing and homogenisation of melts prior to eruption.

626

627

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852 **Figure captions**

853

854 **Fig. 1.** Temporal variation of MgO content in glasses from the Bárðarbunga-Veiðivötn volcanic
855 system. Subglacial (Ljósufjöll), early-Holocene (Brandur, Fontur and Saxi tephra cones), middle-
856 Holocene (Drekahraun) and Veiðivötn 1477 data are presented in this study. ‘Holocene
857 Bárðarbunga’ glass compilation from Óladóttir et al. (2011). 2014-2015 Holuhraun eruption glass
858 composition from Halldórsson et al. (2018). Temporal evolution trends are also observed in CaO,
859 FeO and TiO₂ contents (not shown).

860

861 **Fig. 2.** Geological map of the southernmost part of the Bárðarbunga-Veiðivötn system. The general
862 geology of the area is indicated with greyscale colours. Eruptive units studied in this work are
863 marked in colours. Historic lavas are younger than 1100 BP; prehistoric lavas are older than 1100
864 BP. Triangles show the exact sampling location within each unit. The insert map in the upper left
865 corner shows the outline of the active neovolcanic zones (in grey) and the Bárðarbunga-Veiðivötn
866 system (BV) in red. RR: Reykjanes Ridge; RVB: Reykjanes Volcanic Belt; SVB: Snæfellsnes Volcanic
867 Belt; WVZ: Western Volcanic Zone; MIB: Mid-Iceland Belt; BV: Bárðarbunga-Veiðivötn system; EVZ:
868 Eastern Volcanic Zone; OVB: Öraefajökull Volcanic Belt; NVZ: Northern Volcanic Zone. Geological
869 map compiled by Haukur Jóhannesson and Kristján Sæmundsson published by the Icelandic
870 Museum of Natural History and Iceland Geodetic Survey.

871

872 **Fig. 3.** Backscattered electron (BSE) images showing the main petrographic and chemical features
873 of the studied samples. a) Plagioclase macrocryst from the Veiðivötn 1477 eruption with an
874 oscillatory zoned interior surrounded by a darker rim. b) Glomerophytic clot from Drekahraun.
875 Sector zoned clinopyroxene crystals that have grown with plagioclases. c) Clinopyroxene
876 macrocryst from Veiðivötn 1477 showing resorbed dark cores enclosed by less primitive rims. d)
877 Normally zoned olivine with a large melt inclusion from Brandur. e) Saxi tephra cone nodule with
878 olivine and plagioclase macrocrysts and clinopyroxene in glomerophytic clots. f) Nodule from
879 Þjórsárdalshraun: plagioclase crystals surrounded by homogeneous clinopyroxene; interstitial glass
880 pockets are also observed. g) Plagioclase crystal from Ljósufjöll with crystalline melt inclusions,
881 surrounded by a coarse-grained groundmass. h) Plagioclase crystal from Ljósufjöll within a glassy
882 groundmass. plg= plagioclase; ol= olivine; cpx= clinopyroxene; mi= melt inclusion; gl= interstitial
883 glass; Fo=olivine fosterite content; An=plagioclase anorthite content; Mg#= clinopyroxene Mg#.

884

885 **Fig. 4.** The range in (a) An content of plagioclase, (b) Mg# of clinopyroxene, and (c) Fo content of
886 olivine for each locality. Core and rim compositions are depicted as circles and diamonds,
887 respectively. Clinopyroxene dark and bright sectors refer to sector zones in BSE images. Coloured
888 bands represent the mineral compositions calculated to be in equilibrium with the carrier liquid
889 for each sample, where the glass composition is taken as representative of the carrier melt.
890 Numbers in each corner state the number of point analyses in minerals for that specific locality
891 and mineral phase. In general, macrocryst cores are more primitive than the equilibrium
892 compositions whilst rims are more evolved and tend to overlie the equilibrium bands. Lj: Ljósufjöll;
893 B: Brandur; F: Fontur; S: Saxi; Th: Þjórsárdalshraun; Dr: Drekahraun; V: Veiðivötn 1477. (d)-(f)
894 Variation diagrams showing the TiO₂ content vs An content of plagioclase (d), Al₂O₃/TiO₂ vs Mg#
895 clinopyroxene (e), and NiO content vs Fo content of olivine (f) macrocrysts. 1σ error is smaller than
896 the symbol sizes unless otherwise shown.

897

898 **Fig. 5.** Variation diagrams showing (a) CaO content, (b) TiO₂ content, (c) Al₂O₃/TiO₂, and (d)
899 TiO₂/K₂O vs Mg# of groundmass glasses (circles) and melt inclusions (triangles) from the studied
900 samples. Liquid lines of descent (LLD) in (a) and (b) are calculated starting from the same melt
901 composition (white stars) at different pressures. LLDs in (c) and (d) are calculated at 2 kbar, using
902 two different starting compositions (red stars). See main text for details. All LLDs were calculated
903 using Petrolog3 (Danyushevsky and Plechov, 2011). Blue fields denote published data from
904 Bárðarbunga-Veiðivötn system (Halldórsson et al., 2008, 2018; Hartley et al., 2018; Jakobsson,
905 1979; Óladóttir et al., 2011; Svavarsdóttir et al., 2017). Low K₂O data within the ellipse were
906 acquired from glassy to cryptocrystalline areas which could be affected by quench modifications.
907 1σ error is smaller than the symbols.

908

909 **Fig. 6.** Kernel density estimate plots, with bandwidth 0.3, of calculated pressures (a)-(d) and
910 temperatures (e)-(h) for all studied units. Clinopyroxene-melt pressures are calculated using the
911 Neave and Putirka (2017) barometer; OPAM barometry (Hartley et al., 2018; Yang et al., 1996) was
912 applied both to glasses and PEC-corrected melt inclusions. Reported MI pressures represent
913 compositions with a returned probability of fit higher than 80%. Temperatures are calculated using
914 a glass-only thermometer and cpx, ol and plag-melt thermometers from Putirka (2008). See the
915 main text for more details. The dotted curve in each plot shows the pressure and temperature
916 distribution for Holocene Bárðarbunga glasses from Óladóttir et al. (2011). Errors (± next to the

917 legend) indicate the standard error of estimate of the thermometers and barometers used. Table
918 2 lists the numbers of processed cpx, glass and MIs for each locality.

919

920 **Fig. 7.** Relationship between host mineral composition and melt inclusion equilibration pressure,
921 for (a) subglacial unit (Ljósufjöll) (b) early-Holocene units (Brandur, Fontur and Saxi cones) and (c)
922 middle-Holocene/historical units (Tungnaá lavas and Veiðivötn 1477). Kernel density estimates to
923 the right of the plots show the relative probability of equilibration pressures for olivine- and
924 plagioclase-hosted melt inclusions. MI pressure distributions for Holohraun eruption is reported as
925 dotted curve. Error bars refer to the OPAM standard error of estimate (SEE=1.3 kbar). MIs
926 outlined with dotted lines have Mg#>65 and are the least likely to be three-phase saturated.

927

928 **Fig. 8.** SiO₂/TiO₂ vs Mg# of melt inclusions from this work along with published data from the
929 subglacial Skuggafjöll eruption (Neave et al., 2014), the 10 ka early-Holocene Grímsvötn eruption
930 (Neave et al., 2015), the historical 1783 Laki eruption (Neave et al., 2013) and the 2014-15
931 Holuhraun eruption (Bali et al., 2018). Our melt inclusions span the compositional variability
932 recorded by other eruptions from the EVZ. Also indicated are depleted and enriched end-member
933 melt compositions for the Northern Volcanic Zone (NVZ, red stars) and the Western Volcanic Zone
934 (WVZ, white stars) (Shorttle and Maclennan, 2011). 1σ error is within the symbol sizes.

935

936 **Fig. 9.** Kernel density estimate (KDE) curves showing the relative probability of plagioclase and
937 olivine compositions and how the probability has changed with time (a) An content of plagioclase
938 macrocryst cores (b) Fo content of olivine macrocryst cores. Coloured bars indicate the mineral
939 composition in equilibrium with the most primitive MIs within each magmatic unit. Dotted and
940 dashed lines show KDE for published subglacial and Holocene eruption data for the EVZ.
941 Skuggafjöll (Neave et al., 2014); 10ka Grímsvötn (Neave et al., 2015); 1783 Laki (Neave et al.,
942 2013); 2014-2015 Holuhraun (Halldórsson et al., 2018).

943

944 **Fig. 10.** Schematic cartoon summarizing the proposed evolution of the Bárðarbunga-
945 Veiðivötn magmatic system over time, based on barometry, thermometry and chemical data. a)
946 Subglacial time. The magmatic system is distributed over a wide range of depths, with a main
947 storage zone in the middle crust. At that time, crystal mush bodies were mostly composed of
948 primitive macrocrysts. b) Early-Holocene time when the magmatic system underwent glacial

949 rebound effects. The magmatic system is characterized by (1) a storage zone in the mid-crust at
950 around 10 km depth, mostly made up of evolved macrocrysts and (2) a deep-crustal zone(s) at 17
951 km depth, where more primitive macrocrysts crystallize. The increase in magma productivity
952 facilitated movement of magma from deep regions to the surface. c) The magmatic system
953 configuration since the middle-Holocene. The magmatic system is most likely dominated by a mid-
954 crustal reservoir, with crystal mush horizon(s) made up of evolved macrocrysts. Once crustal
955 pressure equilibrium has been established, mid-crustal storage zone(s) dominate the magmatic
956 system configuration, with new magma passageways being established and melts being
957 homogenized in the middle crust. Primitive plagioclase and olivine crystals are coloured in grey
958 and dark green respectively; evolved plagioclase and olivine crystals are coloured in white and
959 light green respectively. Bár: Bárðarbunga edifice; Veið: Veiðivötn area; plg: plagioclase; ol: olivine;
960 cpx: clinopyroxene; dsl: deactivated/disconnected storage levels; xx=crystal.

961

Figure 1

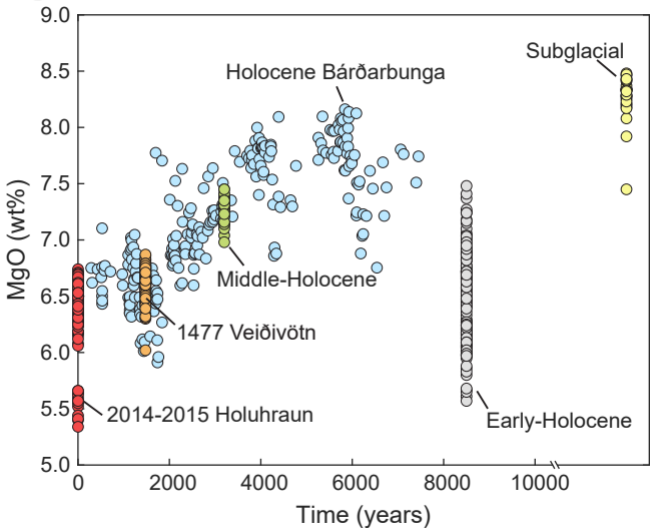


Figure 2

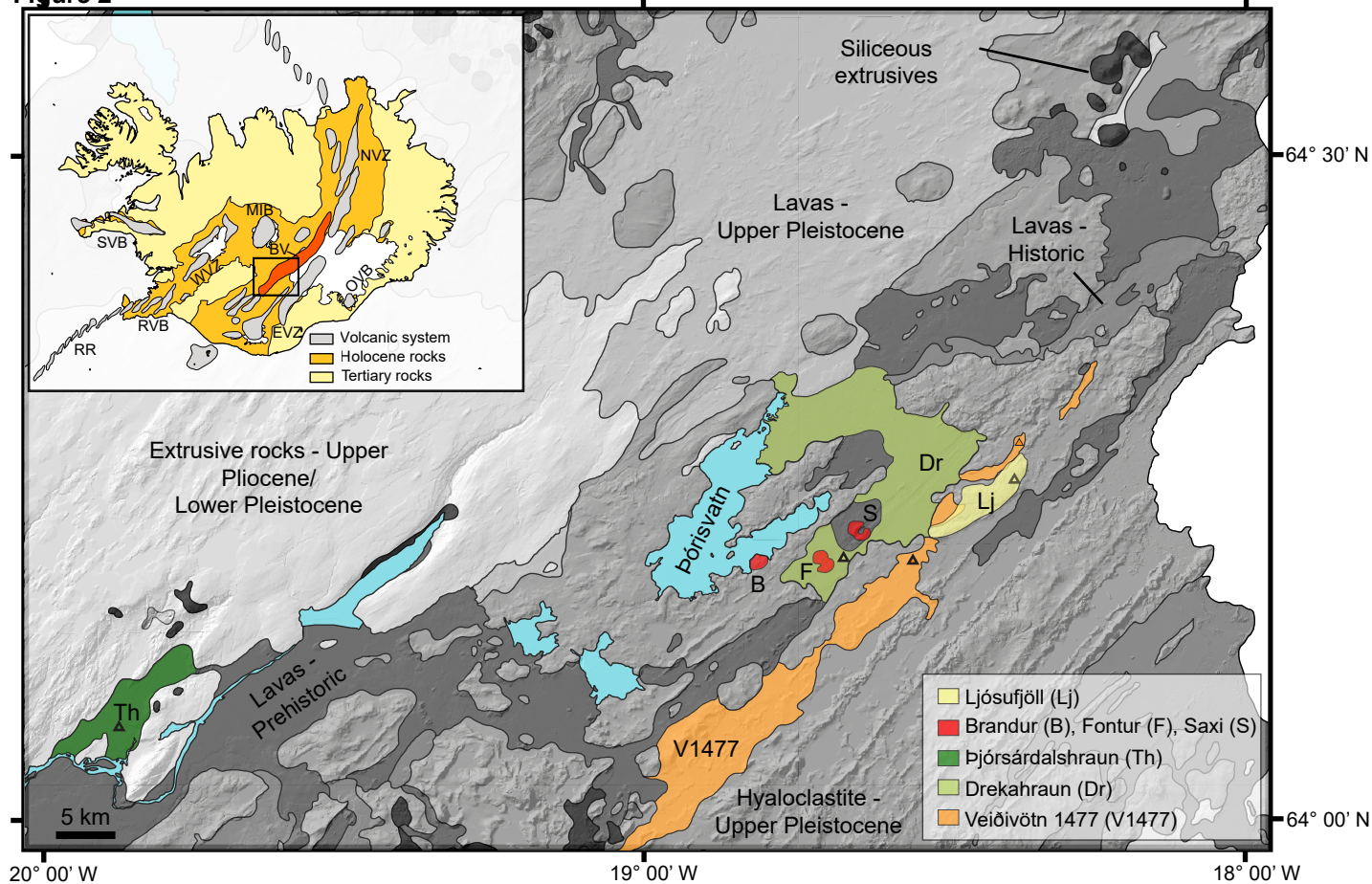


Figure 3

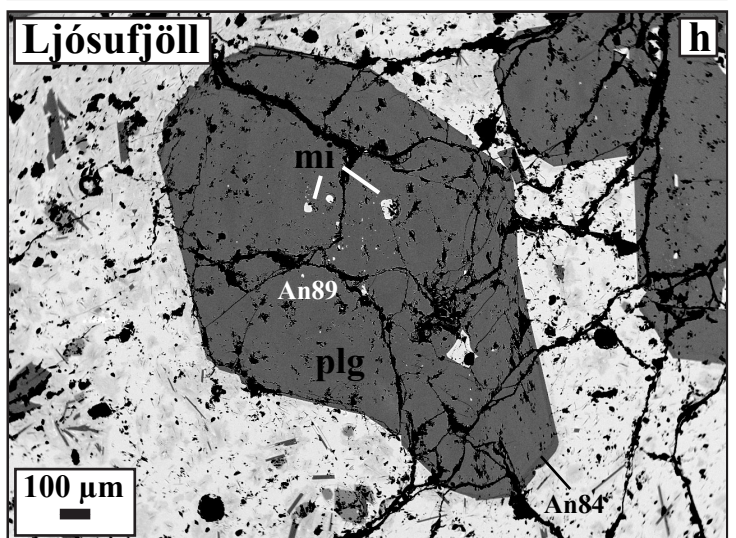
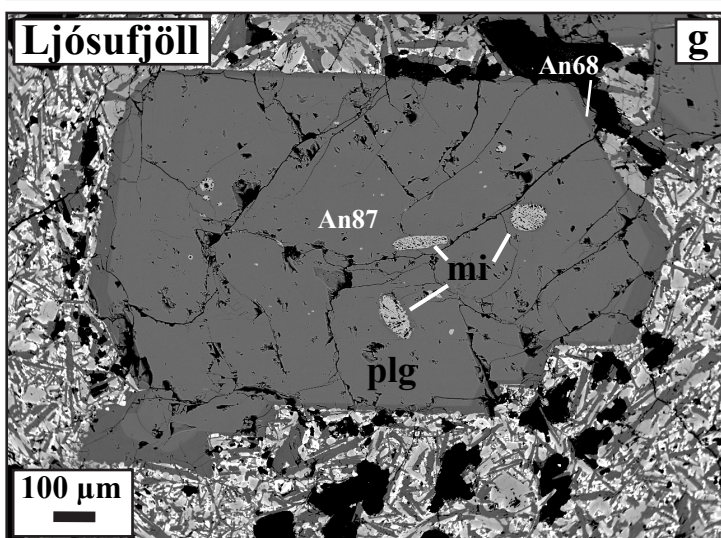
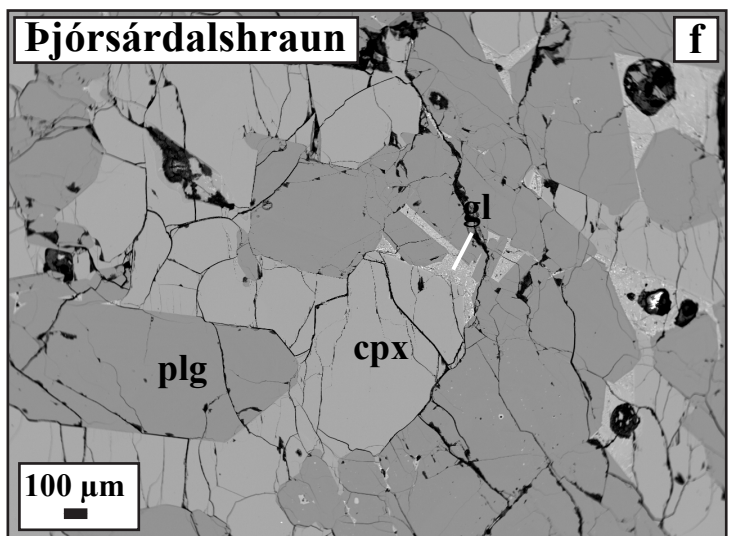
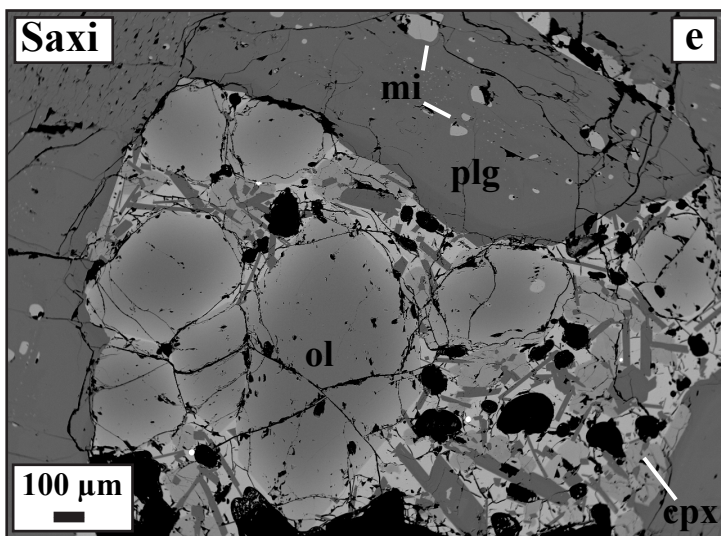
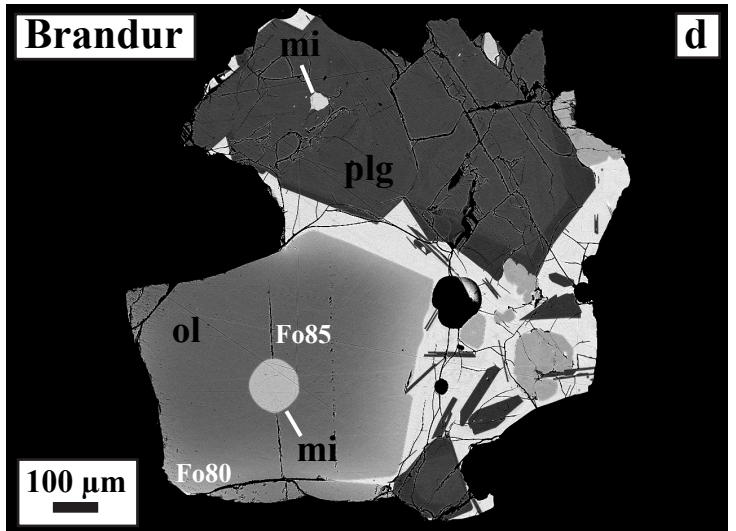
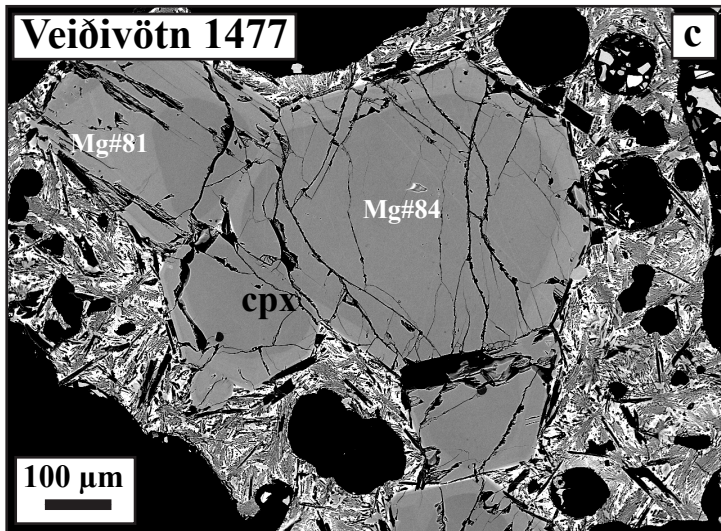
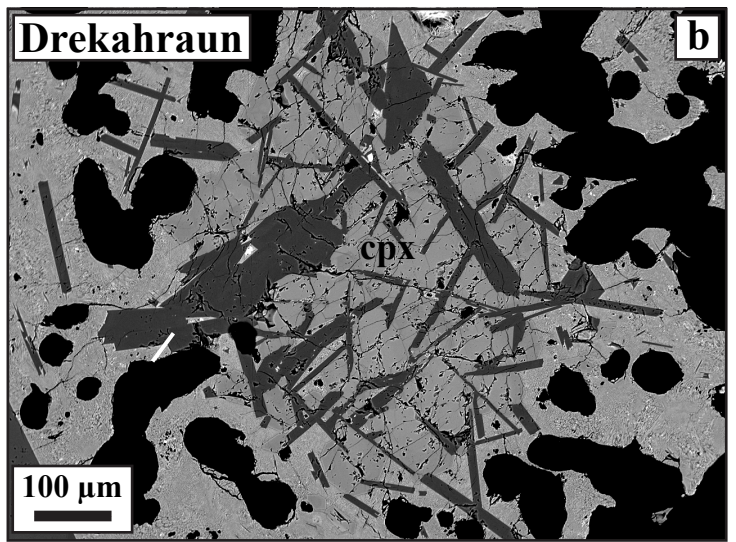
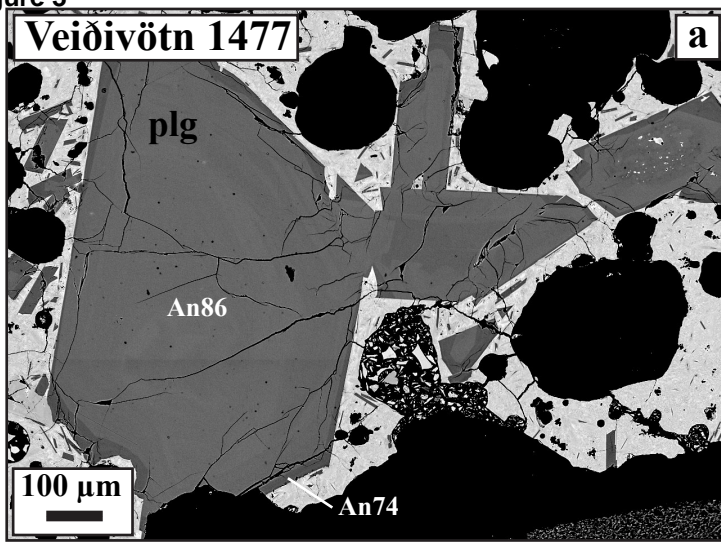


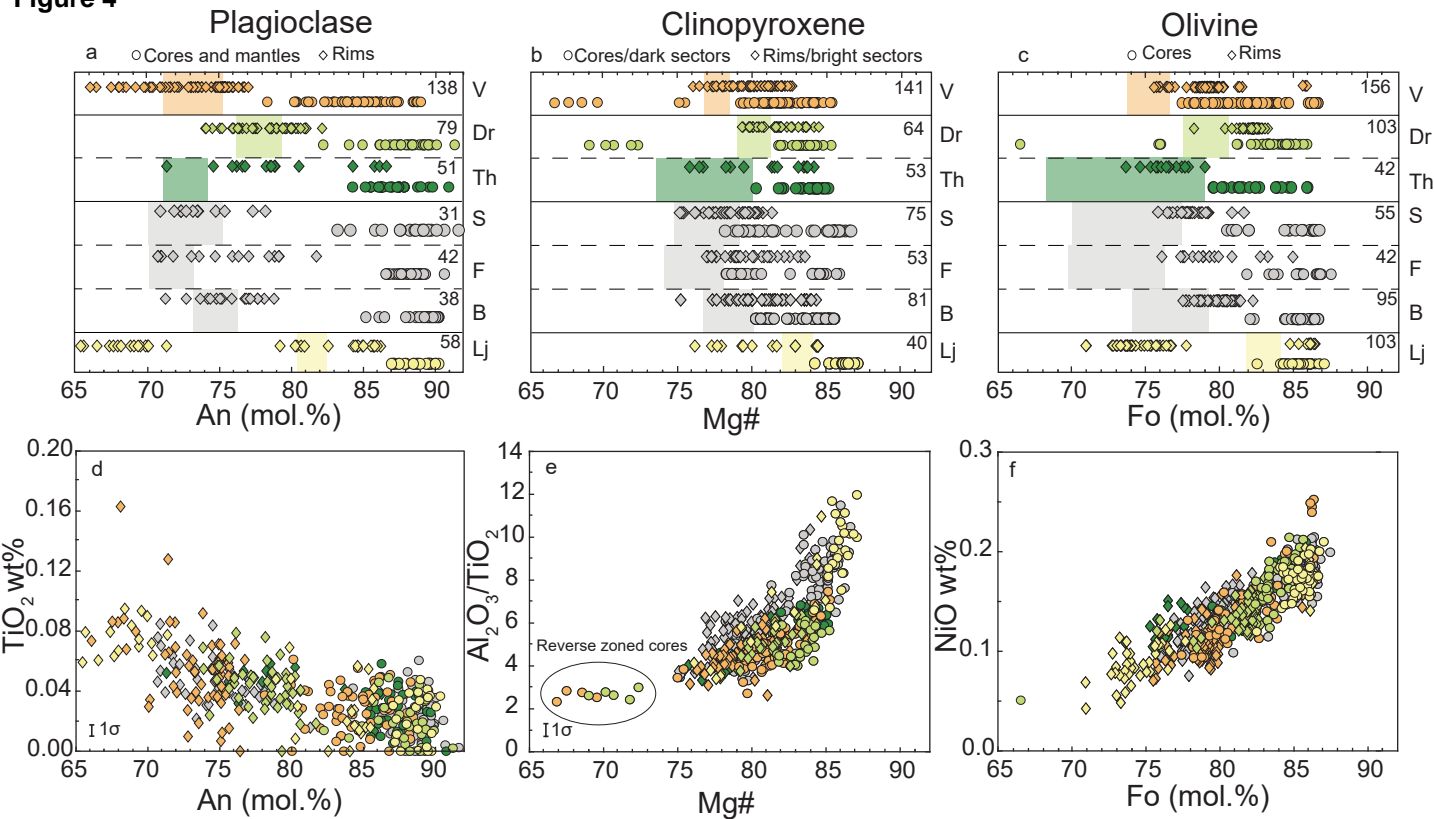
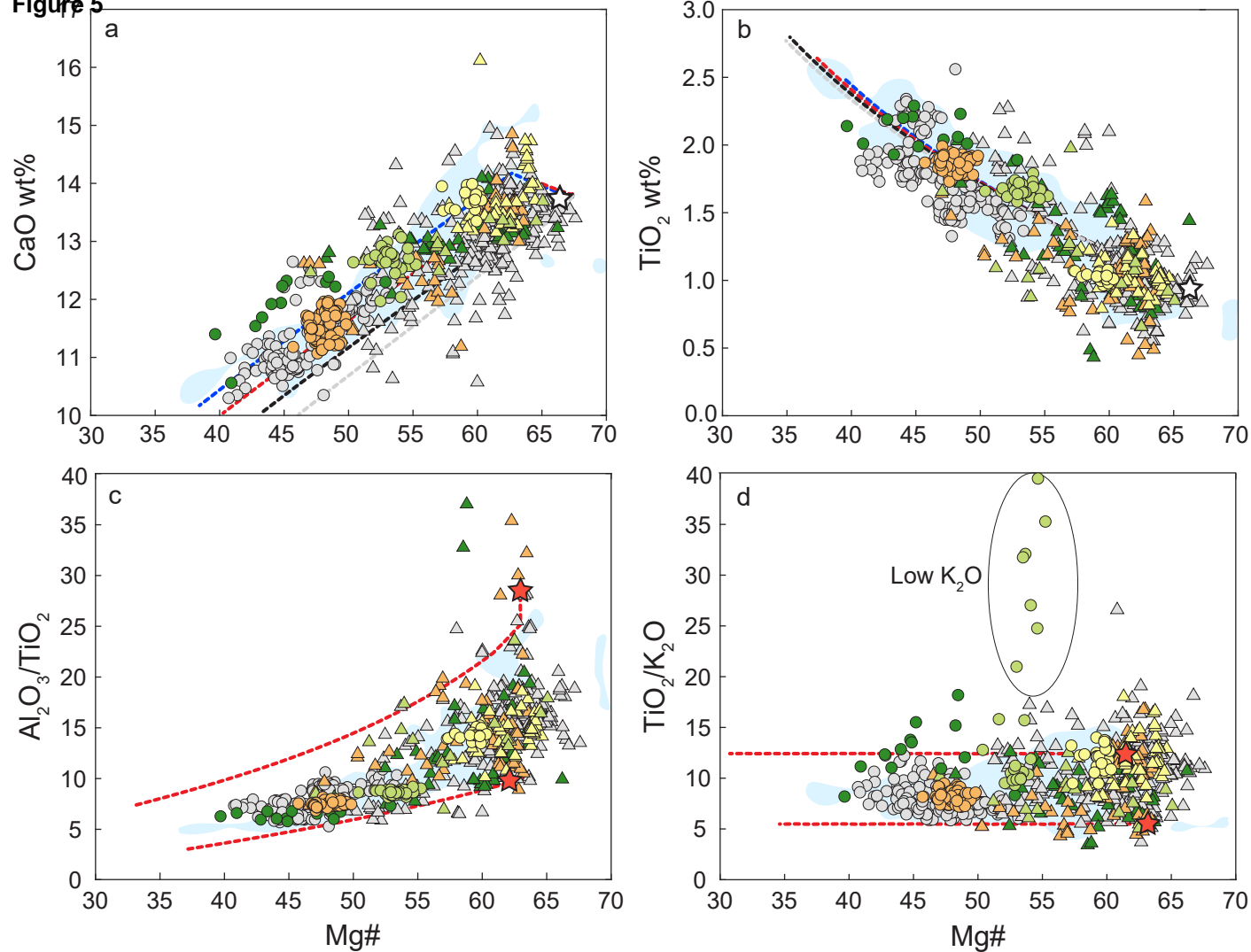
Figure 4

Figure 5

Groundmass glass

● Ljósufjöll ● Brandur, Fontur, Saxi ● Þjórsárdalshraun ● Drekahraun ● Veiðivötn 1477

Melt inclusions

▲ Ljósufjöll ▲ Brandur, Fontur, Saxi ▲ Þjórsárdalshraun ▲ Drekahraun ▲ Veiðivötn 1477

----- 0.001 kbar ----- 2 kbar ----- 4 kbar ----- 6 kbar

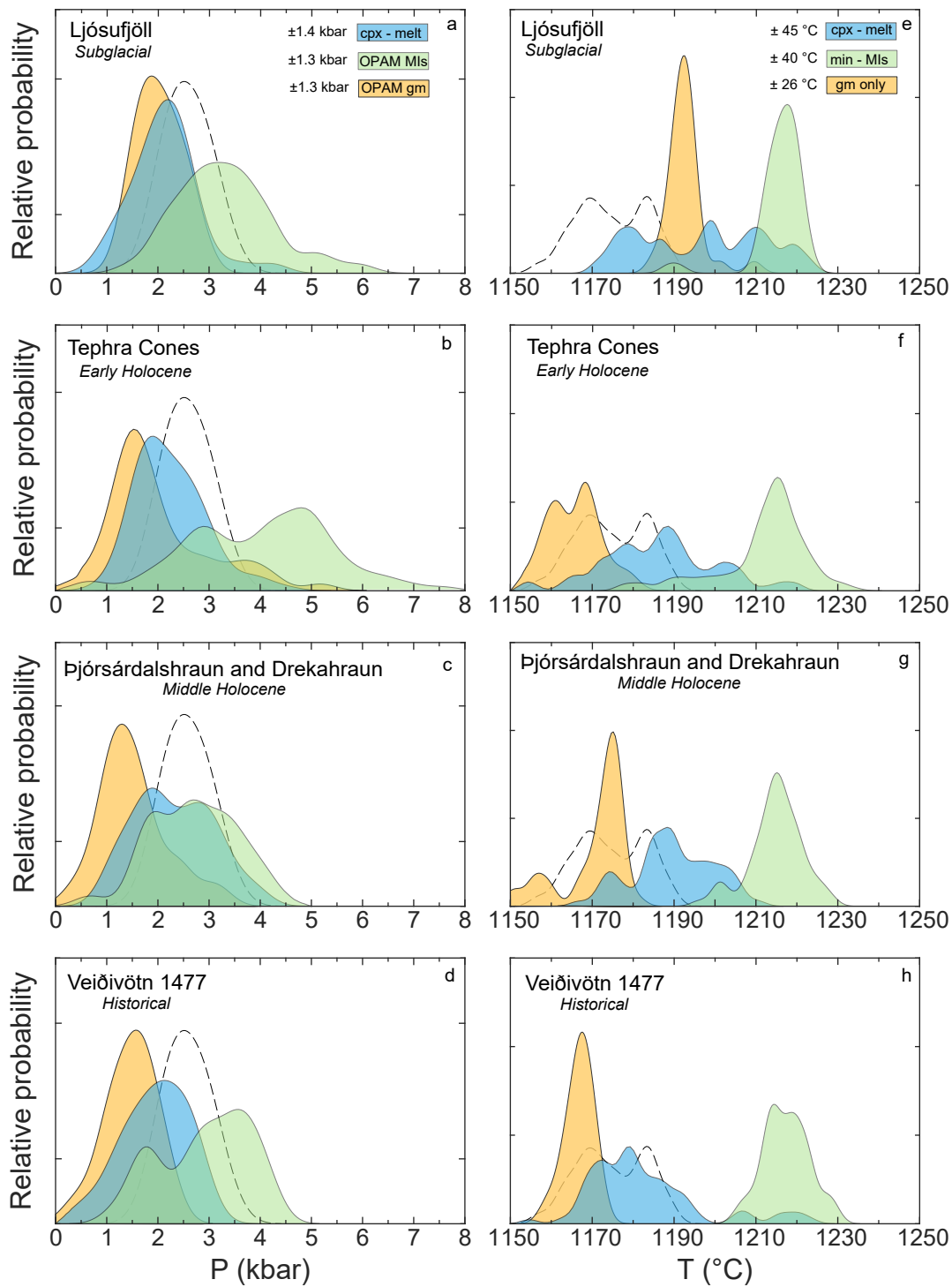
Figure 6

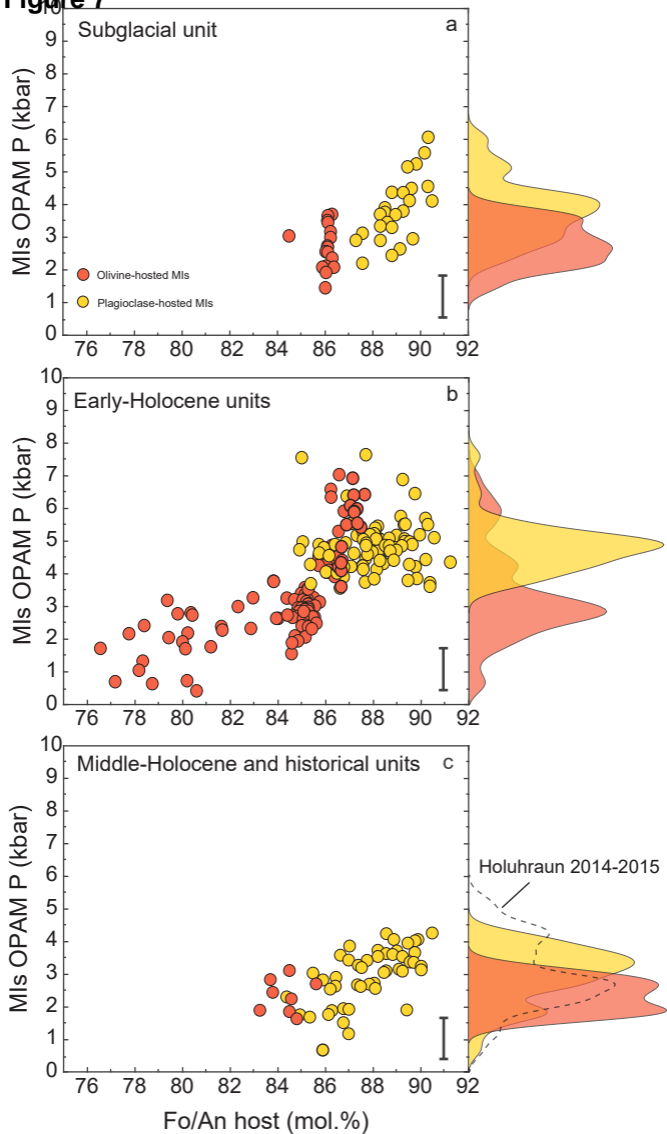
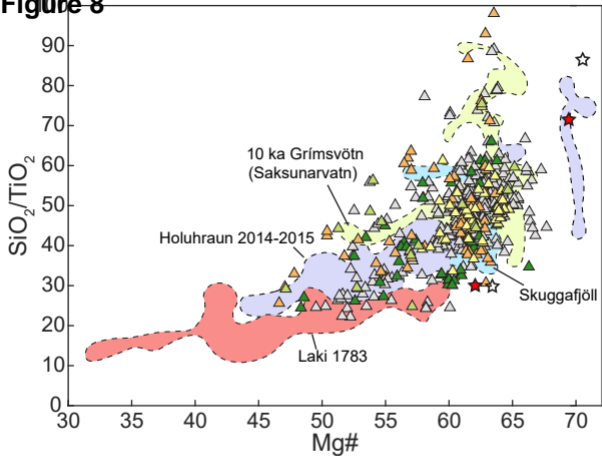
Figure 7

Figure 8

- △ Ljósufjöll △ Brandur, Fontur, Saxi ▲ Þjórsárdalshraun ▲ Drekhraun ▲ Veiðivötn 1477
★ NVZ (Shorttle and MacLennan 2011) ☆ WVZ (Shorttle and MacLennan 2011)

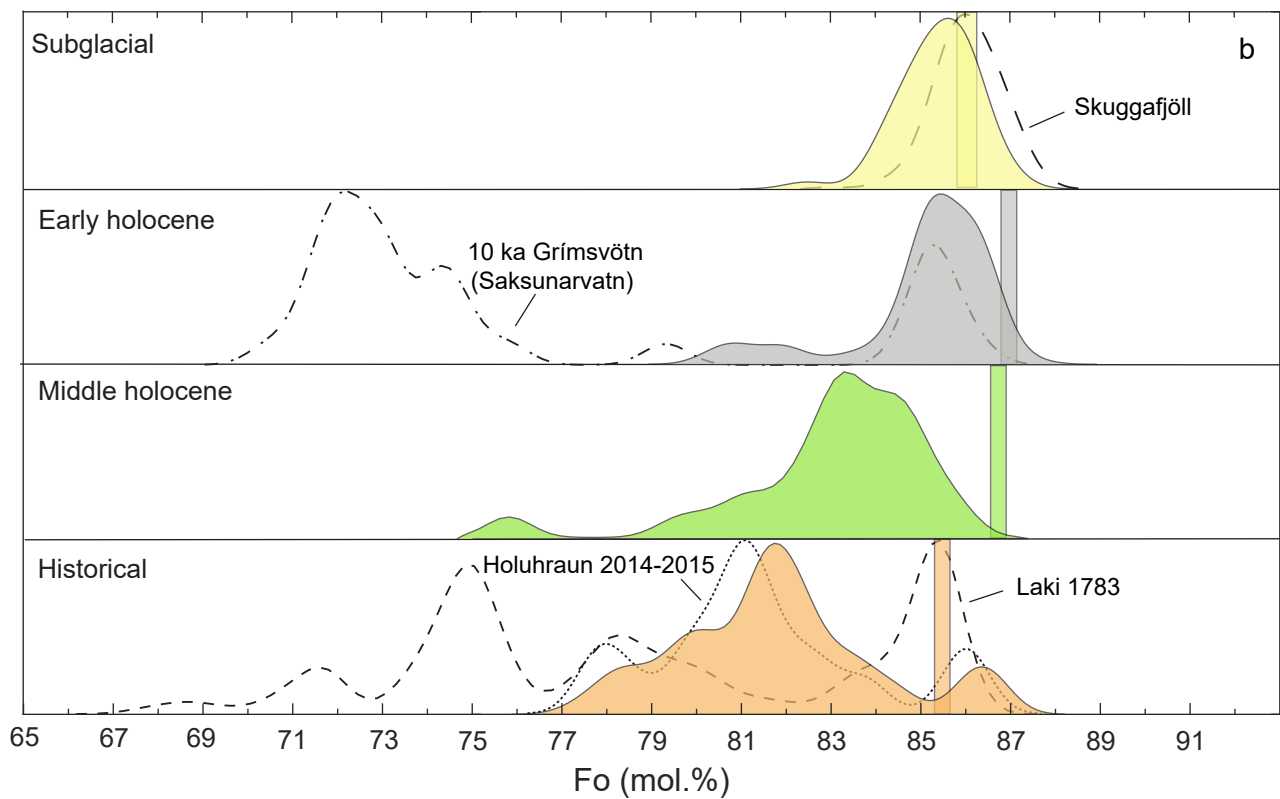
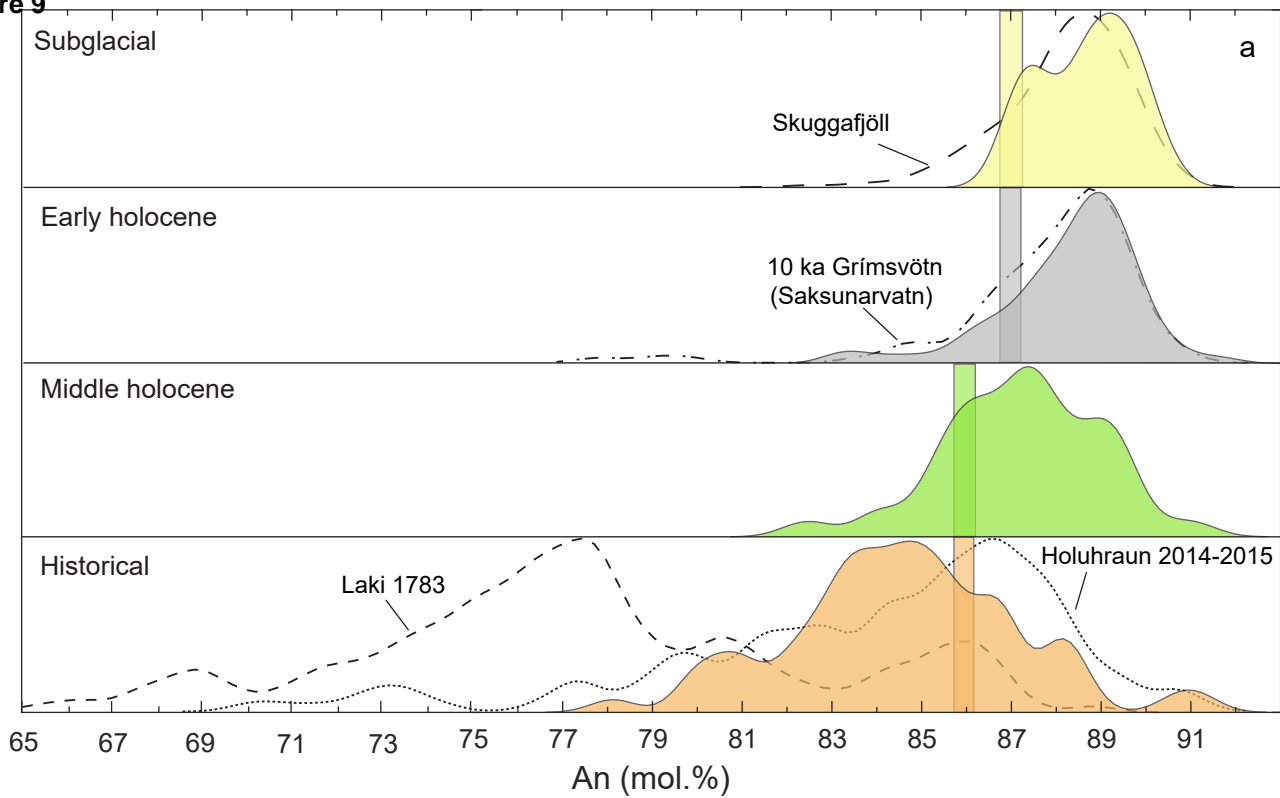
Figure 9

Figure 10

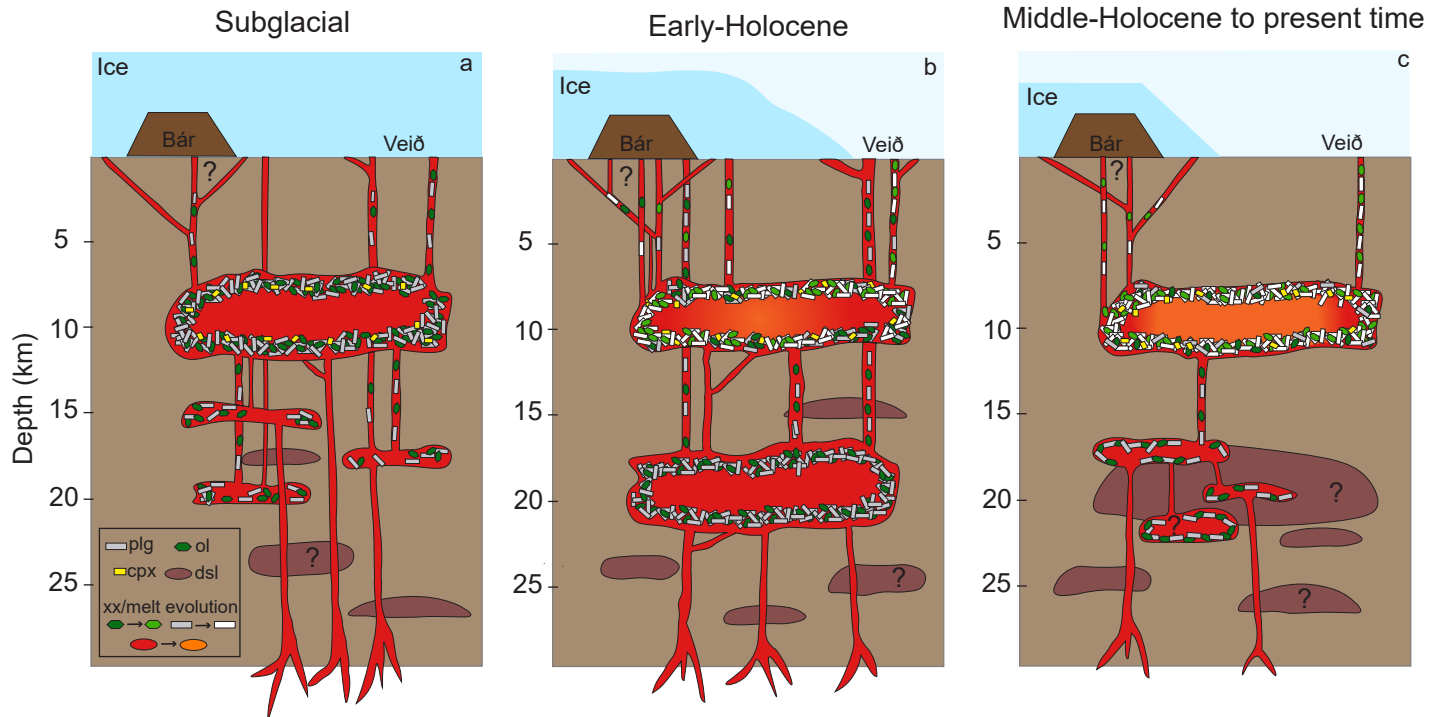


Table 1[Click here to download Table: Table 1.docx](#)**Table 1.** Sample list, description and location.

Sample name	Age	Coordinates		Description	Previous work
		Latitude (N)	Longitude (W)		
Ljósufjöll	Subglacial	64° 16' 51.2"	18° 23' 40.8"	pillow lava	-
Brandur	Early Holocene	64° 13' 55.6"	18° 48' 8.6"	lava, tephra, nodules	Hansen and Grönvold (2000), Holness et al. (2007)
Fontur	Early Holocene	64° 15' 17.9"	18° 38' 0.2"	lava, tephra, nodules	Hansen and Grönvold (2000), Halldorsson et al. (2008), Holness et al. (2007)
Saxi	Early Holocene	64° 13' 49.2"	18° 42' 15.7"	lava, tephra, nodules	Hansen and Grönvold (2000), Halldorsson et al. (2008), Holness et al. (2007)
Þjórsárdalshraun	Middle-Holocene	64° 08' 43.2"	19° 49' 26.8"	lava with nodules	-
Drekahraun	Middle-Holocene	64° 14' 0.4"	18° 39' 57.2"	fresh scoria	-
Veiðivötn 1477	Historical (1477 AD)	64° 14' 29.2"	18° 31' 50.7"	fresh scoria	-

Table 2[Click here to download Table: Table 2.docx](#)**Table 2.** Number of clinopyroxenes, groundmass glasses and MIs processed for geobarometry estimates along with pressure results. Pressures are in kbar.

	cpx-melt barometry				OPAM barometry									
	Clinopyroxenes				Groundmass glasses				Melt inclusions					
	n. analysis	n. cpx-melt	P		n. analysis	P _f >80%	P		n. analysis		P _f >80%		P	
			Mean	σ			Mean	σ	ol-hosted	plg-hosted	ol-hosted	plg-hosted	Mean	σ
Ljósufjöll	43	40	2.1	0.6	29	26	2.0	0.4	18	30	17	24	3.3	0.9
Brandur	59	45	2.2	0.7	69	42	1.6	0.6	86	37	73	22	3.6	1.5
Fontur	34	24	2.2	0.6	33	6	3.5	0.9	24	54	20	31	4.4	1.1
Saxi	47	35	2.3	0.7	63	5	3.2	1.5	39	37	17	33	4.4	1.3
Þjórsárdalshraun	41	35	1.9	0.7	13	11	1.5	0.8	5	33	5	23	2.7	0.9
Drekahraun	63	45	2.6	0.8	30	6	1.5	0.2	0	17	0	6	2.9	1.0
Veiðivötn 1477	144	90	1.9	0.6	91	15	1.5	0.5	4	52	3	25	2.9	0.9
Total	431	314			328	111			176	260	135	164		

P_f = probability of fit ; σ= standard deviation

Table 3[Click here to download Table: Table 3.docx](#)**Table 3.** Mean temperature (°C) values along with the standard deviation (σ) for all studied localities.

	Groundmass glasses			Melt inclusions			Cpx - melt		
	n	Mean	σ	n	Mean	σ	n	Mean	σ
Ljósufjöll	29	1193	4	48	1216	6	40	1197	15
Brandur	69	1170	4	123	1213	9	45	1194	22
Fontur	33	1161	7	78	1213	8	24	1186	12
Saxi	63	1163	5	76	1209	16	35	1186	20
Þjórsárdalshraun	13	1160	9	42	1214	7	35	1187	10
Drekahraun	30	1174	2	17	1216	5	45	1188	15
Veiðivötn 1477	91	1167	3	56	1217	5	90	1182	14