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1 **A tephra lattice for Greenland and a reconstruction of volcanic events**
2 **spanning 25-45 ka b2k**

3
4 Bourne, A.J.^{1*}, Cook, E.¹, Abbott, P.M.¹, Seierstad, I.K.,² Steffensen, J.P.², Svensson, A.²,
5 Fischer, H.³, Schüpbach, S.³, Davies, S.M.¹

6
7 ¹ Department of Geography, College of Science, Swansea University, Swansea, UK

8 ² Centre for Ice and Climate, Niels Bohr Institute, University of Copenhagen, Denmark

9 ³ Climate and Environmental Physics, Physics Institute, University of Bern, Switzerland

10

11 *Corresponding Author. Email: a.j.bourne@swansea.ac.uk

12

13 **Abstract**

14 Tephra layers preserved within the Greenland ice-cores are crucial for the independent
15 synchronisation of these high-resolution records to other palaeoclimatic archives. Here we
16 present a new and detailed tephrochronological framework for the time period 25,000 –
17 45,000 yrs b2k that brings together results from 4 deep Greenland ice-cores. In total, 99
18 tephra deposits, the majority of which are preserved as cryptotephra, are described from the
19 NGRIP, NEEM, GRIP and DYE-3 records. The major element signatures of single glass
20 shards within these deposits indicate that 93 are basaltic in composition with 43 originating
21 from Grimsvötn, 20 are thought to be sourced from the Katla volcanic system and 17 show
22 affinity to the Kverkfjöll system. Robust geochemical characterisations, independent ages
23 derived from the GICC05 ice-core chronology, and the stratigraphic positions of these
24 deposits relative to the Dansgaard-Oeschger climate events represent a key framework that
25 provides new information on the frequency and nature of volcanic events in the North

26 Atlantic region between GS-3 and GI-12. Of particular importance are 19 tephra deposits
27 that lie on the rapid climatic transitions that punctuate the last glacial period. This framework
28 of well-constrained, time-synchronous tie-lines represents an important step towards the
29 independent synchronisation of marine, terrestrial and ice-core records from the North
30 Atlantic region, in order to assess the phasing of rapid climatic changes during the last glacial
31 period.

32

33 **Keywords**

34 Tephrochronological framework; tephrostratigraphy; cryptotephra; Greenland ice-cores;
35 Iceland; rapid climate changes

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39 **Introduction**

40 The Greenland ice-cores have provided an unprecedented insight into the nature of abrupt
41 climatic changes (Dansgaard–Oeschger (DO) events) during the last glacial period (e.g.
42 Dansgaard et al 1993; NGRIP members, 2004). With an independent annually-resolved
43 chronology (Andersen, et al., 2006, Rasmussen et al., 2006; Svensson et al., 2006; Vinther et
44 al., 2006), these records represent significant archives for establishing the history of volcanic
45 events during this time-interval. Both volcanic aerosol (ice acidity and sulphate records) and
46 tephra particulate matter (or glass shards) preserved in the ice permit the reconstruction of
47 volcanic history, but only the volcanic glass shards allow the geochemical identification of
48 the volcanic source and their employment as isochronous marker horizons between disparate
49 archives. A major disparity exists between the number of volcanic events recorded by these
50 two methods, with over 800 events being identified in the GISP2 sulphate record over the

51 past 110,000 years (Zielinski et al., 1996) but only 68 tephra deposits have been recognised to
52 date from four of the deep ice-cores (Abbott and Davies, 2012 and references therein; Coulter
53 et al., 2012; Bourne et al., 2013). As such, there is untapped potential to explore the full
54 record of tephra deposits in the Greenland ice-cores as early work focused predominantly on
55 the presence of easily identifiable visible layers (Grönvold et al., 1995, Zielinski et al., 1996).
56 More recently investigations have moved to search for cryptotephra deposits that contain a
57 low concentration of volcanic glass particles or shards and, as such, are invisible to the naked
58 eye (e.g. Abbott et al., 2012; Davies et al., 2010; Coulter et al., 2012). However these studies
59 focused on a limited number of samples, typically around peaks in ice acidity and sulphate
60 thought to relate to volcanic activity. It has become apparent, however, that in some
61 instances, glass shards from volcanic events can be present in the ice without an associated
62 acidity or sulphate peak suggesting the relationship between the two records of volcanism
63 may be more complex than previously thought (e.g. Davies et al., 2010, in press).

64

65 Here we investigate the cryptotephra content within four deep ice-cores from Greenland
66 spanning 25-45 ka b2k as part of the TRACE project (Tephra constraints on RApid Climate
67 Events). TRACE employs tephra deposits to facilitate the high-precision correlation of
68 palaeoclimatic archives that preserve a record of rapid climate changes that characterised the
69 last glacial period. A systematic search for cryptotephra deposits is undertaken to reduce an
70 over-reliance on chemical indicators in order to build a comprehensive framework of
71 volcanic events preserved within Greenland ice-core records. A lattice of this kind, which
72 combines robust geochemical signatures with well-constrained age estimates, is essential for
73 the wider application of tephrochronology and especially to circumvent any potential mis-
74 correlation that may arise due to an incomplete record of volcanic history. Common tephra
75 deposits that can be traced between the Greenland ice-cores and North Atlantic marine

76 records will provide a robust chronological foundation to test the lead/lag relationships
77 between the atmospheric and oceanic systems over rapid climatic events and permit an
78 assessment of potential causal mechanisms. We report the discovery of 73 new tephra
79 deposits – all of which are available for the precise correlation of marine, terrestrial and ice-
80 core records spanning 25 - 45 ka b2k. This framework represents a significant advancement
81 on the previously published results from this period with just 26 tephra deposits identified in
82 the Greenland ice-cores by Davies et al., (2010) and Bourne et al., (2013). We highlight
83 which of the deposits are potentially most valuable for the synchronisation of palaeoclimate
84 archives. Moreover, our focus on four different ice-cores, provides an insight into the tephra
85 dispersal and preservation patterns over the Greenland ice sheet and also presents an
86 independent method (and test) by which an ice-core chronology can be transferred between
87 cores.

88

89 Until recently only a handful of tephras could be traced between different ice-cores including
90 the widespread Saksunarvatn and North Atlantic Ash Zone II (NAAZII; Z2) deposits
91 identified as visible layers in three ice-cores (Grönvold et al., 1995; Ram et al., 1996;
92 Zielinski et al., 1997; Mortensen et al., 2005; Svensson et al., 2008). Recently, however, the
93 intensified focus on cryptotephra deposits in different ice-cores has allowed Rasmussen et al.,
94 (2013) to use 5 new coeval tephras in NEEM and NGRIP in tandem with acidity match points
95 to transfer the GICC05 timescale to the NEEM ice-core. A further 9 tephra deposits were
96 used as an independent test of this timescale transfer. A similar approach was applied
97 between NGRIP and GRIP by Seierstad et al., (in review) where 20 new tephra pairs support
98 the synchronisation of these two records. The tephra deposits utilised for the aforementioned
99 timescale transfer processes are components of the overall framework for Greenland
100 presented here.

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Methods

Sampling was undertaken on four deep Greenland ice-cores: NGRIP, NEEM, GRIP and DYE-3 (Figure 1). Observations made by Davies et al., (2008; 2010), Abbott et al., (2012) and Coulter et al., (2012) have shown that glass shard particles can be present in the ice without an associated sulphate peak. Therefore a more continuous sampling approach was employed to explore the volcanic record preserved only in cryptotephra form. Sampling was based on the following criteria:

1. Ice spanning rapid climatic transitions (particularly the warming transitions);
2. The likely position of widespread volcanic events yet to be located in the Greenland ice, such as the Campanian Ignimbrite eruption of the Campi Flegrei, dated to 39.28 ± 0.11 ka (de Vivo *et al.*, 2001) and the Dawson tephra deposit from the Aleutian Arc - Alaska Peninsula region of southwestern Alaska, dated to 30,433–30,014 cal yrs BP (Demuro et al., 2008);
3. For the NEEM ice core, the likely positions of selected tephra deposits previously identified in the NGRIP ice core by Davies et al., (2008, 2010) and the presence of glass shards in low-resolution (1.1 m) water samples collected from the NEEM continuous flow analysis (CFA) set-up.

This amounted to 113.3 m of NGRIP ice between 1823.80 m and 2178.00 m and 97.35 m between 1617.55 m and 1845.25 m in the NEEM ice core. The GRIP and DYE-3 cores were largely sampled to investigate the second criterion and therefore the ice sampled is limited to 97.9 m between 1998.15 m and 2231.35 m in the GRIP ice core and 34.10 m between 1865.60 and 1914.00 m in the DYE-3 ice core (Table 1).

125 Ice cross-sections of 2 cm² were removed from the edge of 55 cm long archive core sections
126 stored at the University of Copenhagen. These 55 cm long samples were then cut into 3 sub-
127 samples of either 15 or 20 cm length for NGRIP, NEEM and GRIP. As the DYE-3 record is
128 a lower temporal resolution at these depths, the DYE-3 samples were cut into 6 sub-samples
129 of 10 or 5 cm length. These individual ice samples were melted at room temperature and
130 centrifuged to concentrate any particulate matter. The particulate material was dried onto
131 frosted microscope slides and embedded in epoxy resin. Samples were then examined for
132 tephra shards using optical light microscopy. Any samples containing 5 or more glass shards
133 were subsequently prepared for geochemical analysis. Thin sections of the tephra shards
134 were produced by grinding and polishing the samples using silicon carbide paper and 9, 6 and
135 1 µm diamond suspension.

136

137 Electron-probe microanalysis (EPMA) of the identified glass shards took place during seven
138 analytical periods at the Tephra Analytical Unit at the University of Edinburgh. A Cameca
139 SX-100 electron microprobe with five vertical wavelength dispersive spectrometers was
140 employed to analyse oxide values for 10 major and minor elements within individual glass
141 shards. Both a 3 and 5 µm beam diameter were used, according to the grain-size of the
142 samples, and the operating conditions followed those outlined by Hayward (2012).
143 Secondary standard analyses of Lipari Obsidian and BCR2G basalt were run at the beginning
144 and end of each day, as well as at regular intervals between samples. The full geochemical
145 results, including the operating conditions, beam diameter employed for each sample and
146 standard data are provided in the Supplementary data.

147

148 In all cases, the tephra deposits identified have been given a unique label. This is derived
149 from the name of the ice core and the basal depth of the sample containing the glass shards.

150 For example, the label for the tephra layer in NGRIP sample 2065.45 – 2065.65 m will be
151 NGRIP 2065.65 m.

152

153 The tephra horizons can be assigned ages using the annual-layer counted chronology, the
154 GICC05 timescale for the NGRIP core (see; Andersen et al., 2006; Svensson et al., 2006,
155 Svensson et al., 2008 for details of the layer counting). This timescale has been transferred to
156 the NEEM and GRIP ice cores using a series of reference horizons (chemo-stratigraphy as
157 well as tephra horizons), which allows GICC05 ages to be assigned to any tephra horizons
158 identified within NEEM and GRIP (Rasmussen et al., 2013; Seierstad et al., in review). The
159 GICC05 timescale has errors on the ages based on the concept of maximum counting errors
160 (MCE), which can be viewed as 2σ errors (see Rasmussen et al., 2006; Andersen et al., 2006;
161 Svensson et al 2008). There is no GICC05 chronology for the DYE-3 sections studied here,
162 therefore ages for tephra deposits found in that record are approximations and are inferred
163 from their stratigraphic position and wiggle matching of the DYE-3 isotope record to the
164 NGRIP isotope record. Correlation of tephra deposits between ice-cores may improve the
165 precision of these ages.

166

167 The most likely volcanic source for each tephra deposit is suggested based on comparison to
168 the best available published data. Due to the limited preservation of pre-Holocene deposits on
169 Iceland, no proximal tephra records in the 25-45 ka time-interval are available for comparison
170 (Haflidason et al 2000). Furthermore, distally-preserved tephra data-sets from Icelandic
171 eruptions between 25 and 45 ka are sparse and dominated by the Grimsvötn-sourced Faroe
172 Marine Ash Zones described in Wastegård et al., (2006). The major producers of basaltic
173 tephra during the Holocene are the Grimsvötn, Katla, Veidvötn-Bárdarbunga, Kverkfjöll and
174 Vestmannaeyjar systems (Larsen and Eiríksson, 2007), whilst the central volcanoes that have

175 predominantly erupted silicic tephra during the Holocene are Hekla, Askja, Öräfajökull,
176 Torfajökull, Snæfellsjökull, Eyjafjallajökull and Katla. We employ Holocene glass data-sets
177 (Larsen et al., 2002, Meara, 2012; Óladottir et al., 2008, 2011a and b) and whole rock data
178 (Jakobsson, 1979; 2008) from Icelandic samples/records for these most productive source
179 volcanoes. These data-sets are also supplemented by distal tephra glass occurrences from
180 both Holocene and last glacial eruptions (Boygale, 1994; Hunt et al., 1995; Dugmore and
181 Newton, 1998; Haflidason et al., 2000 and references within, Davies et al., 2001 ; Wastegård
182 et al., 2001, 2006 ; Andrews et al., 2002; Mortensen et al., 2005).

183

184 Correlation of tephra layers between ice-cores is initially based upon major element
185 geochemistry. However, where major element geochemistry alone is not distinctive, the
186 Greenland event stratigraphy, which is based on high-resolution Greenland $\delta^{18}\text{O}$ and calcium
187 records (Rasmussen et al., in review), can be used to discriminate between tephra with similar
188 geochemical signatures positioned in different climate periods (i.e. different interstadials and
189 stadials). In cases where multiple eruptions with similar geochemical composition are located
190 in very similar stratigraphic positions, a potential tephra correlation can be tested according to
191 whether it is consistent with the depth-depth relationship from the chemo-stratigraphic tie-
192 points between the cores which assumes that the ratio of the layer thickness is slowly varying
193 (Seierstad et al., in review, Rasmussen et al., 2013).

194

195 Tephra correlations identified between NGRIP and NEEM and NGRIP and GRIP are
196 presented in Rasmussen et al., (2013) and Seierstad et al., (in review) respectively. These
197 individual tephra deposits are described for the first time and are present in the descriptive
198 biplots of Figure 3 – 7. However, details of the correlations are not dealt with here. The full
199 geochemical data for these deposits are outlined for the first time in the supplementary

200 information. Similarly, tephra geochemical data from Davies et al., (2010) and Bourne et al.,
201 (2013) are not plotted in biplots alongside the new data here and the full data-sets are
202 available in the original publications. However, all previously published tephra deposits from
203 Davies et al., (2010), Bourne et al., (2013), Rasmussen et al., (2013), Seierstad et al.
204 (submitted) are included in the overall tephrochronological framework tabulated in Tables 2
205 and 3.

206

207 **Results**

208 Within our time-window, 42 cryptotephra deposits were identified in NGRIP, of which 39 are
209 basaltic in composition. Twenty tephra deposits were identified in NEEM, of which 17
210 exhibit a basaltic affinity and one is a visible layer. Twenty-two cryptotephra deposits were
211 identified in GRIP all of which are basaltic in composition and 15 cryptotephra deposits were
212 identified in DYE-3 with 14 of basaltic composition (Figure 2). A number of these deposits
213 fall close to rapid transitions on the $\delta^{18}\text{O}$ records (Figure 2).

214

215 For clarity, the results from all four ice cores will be considered in four time periods (Figure
216 2). These periods are determined by the areas of GRIP and DYE-3 that were sampled and
217 consist of: Period 1 from 25 ka to 32 ka b2k encompassing GS-3 to GS-5.2, period 2 from 32
218 ka to 37 ka b2k encompassing GI-5.2 to GS-8, period 3 from 37 ka to 41 ka b2k
219 encompassing GI-8 to GS-10 and period 4 from 41 ka to 46 ka b2k encompassing, GI-10 to
220 GI-12 (Table 1). The number of tephra layers, amount of ice sampled and average grain size
221 in each time period is shown in Table 1. These results include six NGRIP tephra deposits in
222 Period 1 which were previously published by Davies et al., (2010) and twenty tephra deposits
223 in Period 3 (5 in NEEM and 15 in NGRIP) that were previously published by Bourne et al.,
224 (2013) (Table 1).

225

226

227 *Period 1 – 25-32 ka b2k, GS-3 to GS-5.2*

228 The ice sampled in this time period consisted of 55.55 m from NGRIP (46 % of total ice from
229 the period), 30.8 m from NEEM (44 % of total ice), 52.8 m from GRIP (48 % of total ice)
230 and 15.4 m from DYE-3 (Table 1). A total of 37 tephra layers are present within this time
231 period, 8 from NGRIP, 9 from NEEM, 12 from GRIP and 8 from DYE-3. Within this time
232 period six tephra deposits have been previously identified in NGRIP (Davies et al., 2008,
233 2010) and the stratigraphic positions, ages, chemical compositions and source volcanoes of
234 both new and published tephra layers are shown in Table 2.

235

236 Average grain-size for these tephra layers varies between cores with larger shards present in
237 DYE-3 with an average grain size of 57.3 μm (Table 1). In contrast, the average grain size of
238 the GRIP tephra deposits is 36.8 μm , 33.5 μm for NGRIP deposits and 27.1 μm for the
239 NEEM deposits (Table 1).

240

241 Within this time period 35 of the deposits are basaltic in composition with just two of
242 rhyolitic composition, NGRIP 1888.10 m and NEEM 1636.45 m (Figure 3A). With only one
243 glass shard analysed from NEEM 1636.45 m it is difficult to pinpoint a source volcano
244 (Figure 4). NGRIP 1888.10 m, however, shows closest affinity to products from Hekla
245 (Figure 4). Several of the basaltic deposits exhibit similar geochemical signatures (Figure
246 3A, Table 2). The majority of the glass shard analyses for NGRIP, NEEM and GRIP tephra
247 deposits reveal homogenous and tightly-clustered populations. However, one or two outlying
248 analyses are observed in some of the deposits e.g. GRIP 2002.20 m, GRIP 2070.20 m, and
249 NEEM 1669.25 m (Figure 3C and D). In contrast, the 8 deposits present in DYE-3 exhibit

250 marked heterogeneous geochemical signatures. For instance SiO_2 and FeO/TiO_2 values for
251 shards from DYE-3 1865.80 m vary between 46.70 and 59.62 %wt and 3.19 and 7.00 %wt
252 respectively (Figure 3A and E). As such glass shards from this one sample fall within
253 geochemical fields for Katla, Grimsvötn and Veidivötn (Figure 3E). This heterogeneity is
254 common to all DYE-3 tephra deposits and therefore it is not possible to assign these deposits
255 to one source volcano. The only tephra deposit that does exhibit some homogeneity is DYE-
256 3 1869.15 m, which can be tentatively assigned to the Grimsvötn volcanic source.

257

258 Tephra deposits of transitional alkali and tholeiitic basalt composition dominate this period
259 with 12 deposits clustering within the Katla field and 13 deposits falling within the
260 Kverkfjöll/Grimsvötn fields (Figures 3B, C and D). Two distinct tholeiitic tephra deposits
261 are separated from the Katla and Kverkfjöll geochemical clusters observed in Figure 3B and
262 C. NGRIP 1894.05 m and NEEM 1636.65 m are geochemically distinct from the other
263 basaltic horizons with an FeO/TiO_2 ratio of >6 (Figure 3B). Although Veidivötn is a likely
264 source for NGRIP 1894.05 m, the source for NEEM 1636.65 m is uncertain with SiO_2 values
265 ranging between 49.22 and 52.17 %wt and an FeO/TiO_2 ratio of between 7.66 and 8.69 %wt
266 (Figure 3B and C).

267

268 It is very difficult to separate the Katla tephra deposits based on geochemical results alone
269 (Figure 3F). Small variations, however, can be observed between some of the tephra deposits
270 e.g. NEEM 1669.25 m reveal CaO values of 11-11.7 wt% whereas the CaO values for GRIP
271 2049.50 m and NEEM 1656.50 m range between 9.5 and 11 wt%. There is however, a great
272 deal of overlap between these Katla deposits and difficulties may arise in using these for
273 correlating to tephra deposits in other sequences when only one deposit is present. The
274 stratigraphic position of these Katla deposits should, therefore, be used in tandem with the

275 geochemical results to avoid any potential mis-correlations. For instance, NGRIP 1882.10 m
276 and NEEM 1648.90 m have been correlated according to their geochemical signatures and
277 their stratigraphic position within GS-4 by Rasmussen et al., (2013) (Table 2). These are the
278 only transitional alkali basalts deposited during GS-4 and, thus, can be discriminated from the
279 other layers shown in Figure 3B and E. Likewise, GRIP 2070.20 m is the only tephra of
280 Katla origin deposited during GI-5.1 and NGRIP 1929.95 m, NEEM 1677.60 m and GRIP
281 2079.40 m all relate to the same volcanic event during GS-5.2 and are, thus, stratigraphically
282 distinct. However three layers are located in GS-3 (NEEM 1626.15 m, NGRIP 1855.80 m
283 and NGRIP 1861.55 m) with a further three tephra deposits in GS-5.1 which cannot be
284 discriminated from one another using geochemistry or stratigraphy.

285

286 Tephra deposits originating from Kverkfjöll also exhibit similar geochemical signatures but
287 can be discriminated based on very small differences in their TiO₂ values (Figure 3G). For
288 example GRIP 2002.20 m has TiO₂ values of 3.6-3.7% whereas GRIP 2064.35 m has TiO₂
289 values of 3.2-3.3%. Some layers for example NEEM 1664.95 m and NEEM 1671.85 m
290 cannot be discriminated using geochemical signatures (Figure 3G), however their
291 stratigraphic position (GS-5.1 and GI-5.1 respectively) does allow for discrimination between
292 these layers (Table 2).

293

294 ***Period 2 – 32-38 ka b2k, GI-5.2 to GS-8***

295 The ice sampled in this period consisted of 50.05 m from NGRIP (40 % of total ice in this
296 period) and 26.4 m from NEEM (37 % of total ice) (Table 1). Nine tephra layers are present
297 in this time period, 5 from NGRIP and 4 from NEEM (Table 2, Figure 4 and 5). GRIP and
298 DYE-3 were not sampled during this time period. The NGRIP grain size average is 30.6 µm,

299 with an average maximum grain size of 51.0 μm . The NEEM average grain size is 18.6 μm ,
300 with an average maximum shard size of 35 μm (Table 2).

301

302 Seven of the nine deposits in this time period are basaltic in composition, with two
303 transitional alkali deposits and four tholeiitic deposits (Figure 5A). NGRIP 1954.70 m,
304 NGRIP 1952.15 m and NEEM 1690.35 m originate from Katla, Iceland (Figure 5B-D).
305 NGRIP 1952.15 m and NEEM 1690.35 m exhibit overlapping geochemical signatures and
306 are believed to relate to the same volcanic event (Rasmussen et al., 2013) (Figure 5).
307 Despite the Katla origin, NGRIP 1954.70 m is distinctive from NGRIP 1952.15 m and
308 NEEM 1690.35 m (apart from 2 outliers) due to its lower SiO_2 and higher TiO_2 values
309 (Figure 5B-D). The remaining four layers originate from Kverkfjöll (Figure 5B-D) and can
310 be separated into two groups based on their Al_2O_3 and TiO_2 values (Figure 5B). Moreover,
311 these tephra deposits can also be distinguished based on their stratigraphic positions as
312 NGRIP 1950.50 m and NEEM 1689.25 m fell during GI-5.2 and NGRIP 1973.16 m and
313 NEEM 1702.40 m were deposited during GI-6. NGRIP 1950.50 m and NEEM 1689.25 m
314 have lower TiO_2 and higher Al_2O_3 values (Figure 5C) and were assigned to the same volcanic
315 event by Rasmussen et al., (2013). NEEM 1693.45 m and NGRIP 2009.15m are dacitic to
316 rhyolitic in composition and show geochemical affinity to Hekla products (Figures 4 and 5a).

317

318 ***Period 3 – 37-41 ka b2k, GI-8 to GI-9***

319 Within this time period 45.65 m of NGRIP ice (91 % of the available ice-core representing
320 this interval), 26.4 m of NEEM ice (86 % of total ice), 45.10 m of GRIP ice (100% of total)
321 and 18.70 m of DYE-3 ice (100% of total) was sampled and 19 previously unreported tephra
322 layers were identified (Table 1). Of these, two are from NEEM, 10 from GRIP and 7 from
323 DYE-3 (Table 2, Figure 4 and 6). In addition to these layers another 20 tephra deposits in this

324 time period were reported by Bourne et al., (2013) and are included in the overall framework
325 in Table 2 but are not plotted in Figure 6. Of these twenty tephras, 15 were identified in
326 NGRIP and 5 in NEEM and the geochemical results reveal that they all fall within the
327 compositional envelope of the marine Faroe Marine Ash Zone III deposit (Bourne et al.,
328 2013).

329

330 DYE-3 again has the largest average grain size in this time period (54.6 μm , Table 3)
331 compared to the GRIP average of 45.6 μm , the NGRIP deposits of Bourne et al., (2013) have
332 an average grain size of 29.3 μm and all the NEEM tephras deposits have an average grain
333 size of 25.0 μm .

334

335 Of the new deposits reported here, 17 are basaltic in composition (Figure 6A), with 16
336 originating from Grimsvötn and one, GRIP 2213.05 m, originating from Katla (Figure 6 B-
337 C). DYE-3 1895.55 m, DYE-3 1901.80 m, DYE-3 1904.10 m, and DYE-3 1904.15 m are
338 more homogenous than those identified in Period 1, and plot within the Grimsvötn field
339 (Figure 6A-C). However DYE-3 1900.80 m and DYE-3 1912.35 m still show geochemical
340 heterogeneity (Figure 6B and C) but all show geochemical affinity to Grimsvötn. NEEM
341 1784.46 m has a mafic composition, falling in the basaltic andesite composition of the TAS
342 plot (Figure 6A). The only rhyolitic tephra from this period is DYE-3 1898.65 m (Figure 6A)
343 and based on TiO_2 and FeO values is thought to originate from Hekla, as is NEEM 1784.46
344 m (Figure 4). The 17 eruptions from Grimsvötn are geochemically very similar, however
345 small differences in the TiO_2 values do allow these eruptions to be split into three main
346 groupings (Figure 6C) but with only limited stratigraphical separation. Within the highest
347 TiO_2 grouping (3.0-3.5 %wt) both GRIP 2227.15 m and GRIP 2227.90 m are located in GS-
348 10, and thus stratigraphic position cannot be used as an additional discriminatory tool for

349 these tephra deposits (Figure 6C, Table 2). Slightly lower TiO₂ values (2.75-3.00 % wt) for
350 GRIP 2195.45 m, GRIP 2197.25 m, DYE-3 1895.55 m, DYE-3 1901.80 m, DYE-3 1904.10
351 m and DYE-3 1904.15 m, give these a somewhat distinctive character. Both GRIP layers are
352 positioned in GI-8c, meaning stratigraphic discrimination is not possible (Table 2, Figure
353 6C). Finally in the lowest TiO₂ group (2.00-2.75 %wt) NEEM 1747.10 m and GRIP 2190.65
354 m are located in GI-8c (Table 2), and GRIP 2200.75 m, GRIP 2201.50 m, and GRIP 2207.00
355 m are located in GS-9 (Table 2), meaning some limited additional discrimination based on
356 stratigraphy is possible. Thus, small geochemical variations allow the discrimination of
357 tephra deposits within these three groupings, but using their stratigraphic positions as an
358 added discrimination tool in this particular context is limited.

359

360 This time period was sampled intensively to detect whether the Campanian Ignimbrite (CI)
361 tephra layer was present in a Greenland ice-core. This is one of the largest eruptions of the
362 Late Quaternary in the Mediterranean region (Pyle et al., 2006) and dated to 39.28 ± 0.11 ka
363 (de Vivo et al., 2001). Tephra from this eruption was not present in NGRIP or NEEM
364 (Bourne et al., 2013) and no tephra layers with trachy-phonolitic geochemistry, typical of the
365 CI were identified in GRIP or DYE-3 either.

366

367 ***Period 4 – 41-46 ka b2k, GS-10 to GS-12***

368 Within this time period 30.25 m of NGRIP ice (34 % of available ice) and 8.8 m of NEEM
369 ice (18% of total ice) was sampled (Table 1), no ice was sampled from GRIP or DYE-3.
370 Eight tephra deposits are present in this time period, all of them from NGRIP, as only a small
371 amount of NEEM ice was sampled. The stratigraphic positions, ages, chemical compositions
372 and source volcanoes of the tephra layers are shown in Figure 7 and are summarised in Table
373 2. The average grain size of glass shards within the NGRIP deposits in this time period is

374 24.1 μm , which is consistent with the results from this core location in other time periods.
375 Each of the deposits in this time period are tholeiitic basaltic in composition and all originate
376 from the Grimsvötn volcano, although NGRIP 2163.35 m also contains a sub population of
377 dacitic shards that appear to originate from Hekla (Figure 4), suggesting two closely spaced
378 eruptions that are not stratigraphically resolved in the 20 cm sample. All of the Grimsvötn
379 deposits, are geochemically very similar, however they can be split into two groups based on
380 the CaO composition with NGRIP 2150.90 m, NGRIP 2162.05 m and NGRIP 2185.70 m
381 forming one group with lower CaO and TiO_2 values (Figure 7B and C). These 3 deposits can
382 also be separated stratigraphically and fell during GI-11 (NGRIP 2150.90 m), GS-12 (NGRIP
383 2162.05 m) and GI-12c (NGRIP 2185.70 m) (Figure 2, Table 2). NGRIP 2162.60 m, NGRIP
384 2163.35 m, NGRIP 2164.10 m and NGRIP 2188.25 m form the second group with a lower
385 FeO/TiO_2 ratio and higher CaO values (Figure 7C and D). Stratigraphically NGRIP 2188.25
386 m can be distinguished from the other 3 layers, as it is positioned in GI-12c, as opposed to
387 GS-12. NGRIP 2186.80 m is the most geochemically distinct layer in this time period with
388 lower FeO and higher CaO values and can, thus, be easily discriminated from the younger
389 NGRIP tephra in this period (Figure 7D).

390

391 **Discussion**

392

393 An investigation of tephra deposits preserved within 4 different Greenland ice-cores provides
394 a detailed record of Icelandic volcanism over the glacial period between 25 and 45 ka b2k
395 (Table 2). Together with the previously published tephra deposits in Davies et al., (2008,
396 2010) and Bourne et al., (2013), 99 tephra layers are identified during this interval. This
397 framework represents a significant advancement in our understanding of the Icelandic
398 volcanic history and is an important first step towards widening the use of tephra horizons for

399 the synchronisation of the ice-cores with other palaeoclimatic archives. Some tephra deposits
400 within this framework will be more valuable than others as marker horizons, but a detailed
401 history of volcanic events is important to preclude any potential mis-correlations. When
402 assessing the potential of individual tephra deposits for correlation purposes, the most
403 valuable deposits will be: i) robustly characterised and geochemically distinct, ii) widespread
404 in extent and iii) well-dated and deposited close to an event of rapid change (Davies et al.,
405 2012).

406

407 *Assessing the value of individual tephra deposits: geochemical characterisation*

408 All of the layers identified here have been robustly geochemically characterised but several
409 deposits exhibit similar geochemical signatures. Their use as time-parallel marker horizons is
410 subject to careful scrutiny of geochemical results and, where possible, the stratigraphic
411 position of the tephra in question. In particular, ninety-four of the layers are basaltic in
412 composition with 43 originating from Grimsvötn, 17 deposits are from Kverkfjöll, which has
413 previously been suggested to form a single volcanic system with Grimsvötn (Grönvold and
414 Jóhannesson, 1984), and 19 are from Katla. Whilst several of these layers are geochemically
415 similar, 70 of the layers can be discriminated based either on small geochemical differences
416 or their stratigraphic position (provided this can be adequately resolved in other sequences).
417 Often, however, the small geochemical differences are between 0.2 and 0.5 wt% and, thus, it
418 is essential that geochemical analysis of any potential correlatives is bracketed by analysis of
419 international secondary standards.

420

421 This large number of basaltic horizons is in contrast to the number identified in the same
422 time-interval within the European INTIMATE tephra framework, where only 2 basaltic
423 horizons are identified (Blockley et al., 2012; Davies et al., 2012). This difference in the

424 number of basaltic tephra layers found in Greenland and in terrestrial European records could
425 be due to the preferential dispersal of basaltic eruptions from Iceland towards Greenland or
426 could reflect the fact that routine density separation techniques employed to detect
427 cryptotephra in terrestrial records does not allow detection of basaltic material (Turney, 1998;
428 Blockley et al., 2005; Larsen and Eiriksson., 2007). This situation may well change with the
429 wider application of a magnetic separation technique for the isolation of glass shards of
430 basaltic composition from sedimentary deposits (e.g. Mackie et al., 2002; Griggs et al., in
431 press).

432

433 *Assessing the value of individual tephra deposits: geographical extent*

434 As yet, the geographical extent of these tephra deposits outside of Greenland is currently
435 unknown, but by investigating the tephra record within the different ice-cores, we can
436 reconstruct the extent of ash deposition over the ice sheet. NGRIP and NEEM correlations
437 and NGRIP and GRIP correlations have been outlined previously by Rasmussen et al., (2013)
438 and Seierstad et al., (in review), respectively (Table 2, Figure 8). We advance this work by
439 highlighting 11 new correlations here giving particular attention to those tephras that can be
440 traced between more than 2 ice-cores. A summary of all ice-core correlations, both new and
441 published, is presented in Table 3. Statistical analyses (similarity coefficient and statistical
442 distance) support these correlations and none of the statistical distances exceed the critical
443 value, therefore no correlations are statistically different (Table 3).

444

445 Within Period 1, major element results indicate that the Katla-sourced deposit found within
446 NGRIP 1895.24 m and NEEM 1656.50 m by Rasmussen et al., (2013) can also be extended
447 to GRIP 2049.50 m (Figure 9A). Secondly GRIP 2060.85 m can be correlated to both NGRIP
448 1908.70 m and NEEM 1664.95 m, which were themselves correlated by Rasmussen et al.,

449 (2013) (Figure 9A). Finally, GRIP 2079.40 m can be correlated to NEEM 1677.60 m which
450 has already been correlated to NGRIP 1929.95 m by Seierstad et al., (in review) (Figure 9A).
451 In Period 2, NGRIP 1973.16 m and NEEM 1702.45 m can be correlated for the first time
452 (Figure 9B), their ages of $33,686 \pm 1207$ yrs b2k and $33,692 \pm 1208$ yrs b2k (Table 2) support
453 this correlation. This correlation also provides a further independent test for the volcanic
454 matching method used to transfer the GICC05 timescale to NEEM.

455

456 Within period 3, five correlations can now be made between GRIP tephra layers presented
457 here and the NEEM data published in Bourne et al., (2013). The GRIP to NGRIP
458 correlations are considered by Seierstad et al., (in review). GRIP 2195.45 m correlates to
459 NEEM 1755.60 m, GRIP 2197.45 m to NEEM 1757.10 m, GRIP 2201.50 m to NEEM
460 1759.85 m, GRIP 2207.00 m to NEEM 1764.25 m and finally GRIP 2227.15 m to NEEM
461 1780.20 m (Figure 9c). Each tephra correlation has a similarity coefficient greater than 0.97
462 and the statistical distance does not exceed the critical value (Table 3), supporting the
463 geochemical correlations (Figure 9C). Correlations to other GRIP layers in this period e.g.
464 GRIP 2200.75 m and GRIP 2202.40 m can be excluded based on their stratigraphic position
465 (Figures 2 and 8). Whilst the DYE-3 tephra layers appear to correlate with some of the GRIP
466 layers (Figure 6), it is clear from their FeO/MgO ratios that DYE-3 1901.80 m, DYE-3
467 1904.10 m and DYE-3 1940.15 m are offset from GRIP 2195.45 m and GRIP 2197.25 m
468 (Figure 9C). Therefore, whilst the DYE-3 layers in this time period reveal more
469 geochemically homogenous populations than in period 1, they do not correlate with layers in
470 the other ice-cores.

471

472 The correlation of GRIP 2197.45 m to NEEM 1757.10 m also implies a correlation to NGRIP
473 2066.95 m as the NEEM and NGRIP layer were correlated by Rasmussen et al., (2013). This

474 is supported by the geochemical data (Figure 9C, black triangles), however, the correlation of
475 NGRIP 2066.95 m and GRIP 2197.45 m is stratigraphically inconsistent with the recent
476 synchronisation of the NGRIP and the GRIP cores based on chemo-stratigraphic records
477 (Seierstad et al. submitted). According to the chemo-stratigraphic matching the two tephra
478 layers are separated by 5 to 13 years (according to the actual stratigraphic position of the
479 tephra deposit within the 15 cm ice sample) (Seierstad et al., in review). Thus, the
480 geochemical signatures support a tephra correlation, but the inconsistencies with the chemo-
481 stratigraphic matching prevents a firm correlation (dashed red line, Figure 8).

482

483 With the new tephra correlations outlined here, 8 tephra horizons are common to GRIP,
484 NGRIP and NEEM (Figure 8). A further four correlations are present between NGRIP and
485 NEEM (green lines) and one additional correlation links NGRIP and GRIP (orange line)
486 (Figure 8). The layers that only correlate between NGRIP and NEEM are found in period 2
487 and late in period 1, where the GRIP core was not sampled, indicating that with further
488 targeted sampling of GRIP more correlations between all three cores may be identified.
489 Single age estimates for these correlating tephra deposits are shown in Figure 8. These ages
490 represent the basal age of the NGRIP sample, as the glacial part of the GICC05 chronology
491 was based on NGRIP annual layer counting. If these layers are traced beyond Greenland then
492 the ages presented in Figure 8 represent the age of the tephra deposit. No correlations were
493 possible with the DYE-3 deposits due to their geochemical heterogeneity (especially in
494 period 1) and geochemical offsets with period 3 deposits (Figures 3 and 9C). Glass shards
495 from the same samples in DYE-3 show affinities to Katla, Grimsvötn and Veidivötn (Figure
496 3E). Many deposits are found within consecutive samples and the heterogeneous
497 geochemical signatures suggest mixing of different tephra deposits. It is possible, that the
498 lower temporal resolution of DYE-3 during the glacial is too low (~ 100 m) to isolate

499 deposits from individual eruptions as seen in the other cores. Secondly, Ram and Gayley
500 (1991) discuss whether the aggregates from the Z2 eruption (which is spread over 78 cm in
501 DYE-3) were caused by melt and refreeze, which could also be the cause of the geochemical
502 heterogeneity observed here. Alternatively, as DYE-3 is the most southerly ice-core (Figure
503 1) it is possible that storms from Iceland could redeposit tephra on the ice-sheet surface
504 (Arnalds et al., 2013; Prospero et al., 2012).

505

506 The deposits that correlate between GRIP, NGRIP and NEEM suggest a northerly dispersal
507 of ash from Iceland, and they represent the most widespread deposits identified to date
508 (Figure 8). The decrease in tephra grain size with increasing northerly latitude supports this
509 direct transport route in a north westerly direction from Iceland to the Greenland core sites
510 (Figure 10). This decrease appears to be a consequence of increasing distance from Iceland,
511 which holds true when considering the distance of the ice-cores from Katla (Figure 10A),
512 however GRIP is actually closer to Grimsvötn (the most common source of tephra layers
513 here) than DYE-3 (Figure 10B). DYE-3 is at a lower altitude (2480 m above sea level (asl),
514 than GRIP (3230 m asl) (Vinther et al., 2006) (Figure 1) suggesting that ash travelling to the
515 GRIP drill site still has further to travel than that reaching DYE-3. The length of a typical air
516 mass trajectory from Iceland to the drill sites may be the cause of the grain size decrease.
517 This is very much dependent on the pathway of cyclones over the North Atlantic as
518 retrograde transport relative to the overall westerly flow is required. On the whole, the
519 decreasing grain-size trend with increasing latitude suggests ash is transported directly to the
520 core sites from source. Whether or not these deposits are widespread beyond Greenland
521 remains to be seen, and will require high-resolution investigations of sequences in the North
522 Atlantic region and north and west of Greenland (if available). Until then, it is unknown
523 whether volcanic ash from these eruptions was also dispersed eastwards towards Europe.

524

525 *Assessing the value of individual tephra deposits: tephra constraints on rapid climate events*

526 In order to optimise the application of tephrochronology to establish the phase relationships
527 between different proxy records, the most valuable tephtras are those that fall close to rapid
528 climatic events. In this case a tephra layer is considered to fall close to a rapid climate event
529 if its age is within 100 years of the age of the stratigraphic boundaries defined by Rasmussen
530 et al., (in review). Of the most widespread deposits only two, NGRIP 2066.95/NEEM
531 1757.10 m/GRIP 2197.45 m (no. 11 on Figure 8) and NGRIP 2071.50 m/NEEM 1759.85
532 m/GRIP 2201.50 m (no. 12 on Figure 8), constrains a rapid climate event (GI-8c onset)
533 evident in the NGRIP $\delta^{18}\text{O}$ curve (Figure 11).

534

535 However, five deposits common to both NGRIP and NEEM were deposited during the
536 warming and cooling transitions of GI-3, GI-4, GI 5.2 and GI-6 (Figure 11). Finally a further
537 5 tephra deposits present only in NGRIP, 3 deposits found only in NEEM and 4 found only in
538 GRIP are also found on rapid warming or cooling transitions (Figure 11). Thus, 19 tephra
539 deposits within this framework (but 28 tephtras from different cores) constrain rapid climate
540 events of interest within the Greenland ice cores. Despite their useful stratigraphic position,
541 their potential value to link different records rest on their distinct geochemical composition
542 relative to other tephtras close in age. Of these 19 tephra deposits NEEM 1636.65 m, NGRIP
543 1861.55 m (Figure 12A), NGRIP 1882.50 m, NGRIP 1888.10 m (Figure 12B), NGRIP
544 1950.50/NEEM 1689.25 m, NGRIP 1952.15 m/NEEM 1690.35 m (Figure 12D), NGRIP
545 1973.16 m/NEEM 1702.45 m, NGRIP 2009.15 m (Figure 12E), NGRIP 2100.65 m and
546 NEEM 1784.46 m (Figure 12G) are all compositionally unique from the other layers that sit
547 in similar stratigraphic positions to these key layers, making these the most useful layers for
548 tracing into other archives. NGRIP 1882.10 m/NEEM 1648.9 m cannot be chemically

549 distinguished from NGRIP 1895.24 m/NEEM 1656.50 m/GRIP2-49.50 m, however they fall
550 in GS-4 and GS-5.1, respectively, so could be distinguished from one another if found in
551 other well-resolved archives. However, NGRIP1915.50 m/NGRIP1915.63 m/NEEM
552 1669.25 m, GRIP 2067.85 m, GRIP 2070.20 m, GRIP 2070.60 m, NEEM 1671.85 m,
553 NGRIP2066.95 m/NEEM1757.10 m/GRIP2197.25 m, GRIP2200.75 m and NGRIP 2071.50
554 m/NEEM1759.85 m/GRIP2201.50 m are all geochemically indistinguishable from other
555 tephra deposits within the same stratigraphic unit (Figure 12C and F) suggesting that robust
556 correlations of these deposits to other sequences will be more challenging.

557

558 Therefore the 10 layers that fall close to rapid climate events and are geochemically
559 distinctive (layers in bold italic, Figure 11) are the most useful layers for establishing the
560 phase relationships between different proxy records. NGRIP2066.95 m/NEEM1757.10 m
561 /GRIP2197.25 m, GRIP2200.75 m and NGRIP 2071.50 m/NEEM1759.85 m/GRIP2201.50 m
562 (numbers 11 and 12 Figure 8) are valuable as they are widespread and fall on a rapid
563 transition, however they are geochemically similar to other tephra layers of a similar age and
564 therefore care is needed if these are correlated to other archives. Likewise the other
565 widespread deposits that are found across Greenland (Figure 8) may also represent valuable
566 isochrons for future correlations to other disparate sequences.

567

568 *Chemical indicators of volcanism in ice-cores and their relationship to tephra deposits*

569 Initial work on tephra deposits preserved in ice cores focussed on horizons that were visible
570 in the record or sections of ice where large sulphate spikes were present (Grönvold et al.,
571 1995; Mortensen et al., 2005). However recent research has begun to question whether using
572 the sulphate record is a reliable method for locating tephra deposits (e.g. Coulter et al., 2012).
573 Our investigation of more than 400 m of ice allows a detailed insight into the imprint of

574 volcanic aerosol fallout (especially sulphate) in the ice alongside the stratigraphic position of
575 volcanic glass shards. The position of tephra deposits are considered relative to the NGRIP
576 electrical conductivity measurements (ECM, Fig. 13 a) (Dahl-Jensen et al., 2002) and NGRIP
577 concentrations of sulphate, dust and calcium as well as the conductivity of the liquid phase
578 (Bigler, 2004) with a particular focus on the continuously sampled portion of ice spanning
579 GI-9, GS-9 and GI-8 (Figure 13b and c).

580 Isolating and detecting volcanic sulphate spikes above a fluctuating and climatically-driven
581 background level is complex (Figure 13). Sulphate concentrations in ice cores have a
582 complex origin, sea salt, mineral dust, biogenic $\text{H}_2\text{S}/\text{SO}_2$ and volcanism contribute to the
583 sulphate concentrations observed and a volcanic eruption will often give rise to sulphate
584 spikes with concentrations of 3 - 10 micro equivalent/kg (6 - 20 micro moles/kg) (Steffensen,
585 1995). When climate variability is low (e.g. during the Holocene and interstadial periods) the
586 background level of sulphate in the ice is around 1-2 micro-equivalent/kg and natural
587 variability in sulphate concentration is also low (1-2 micro-equivalent/kg inter annually)
588 (Figure 13A and B). During these warm episodes the large sulphate spikes often stand out
589 above background signals and are clearly detectable (Figure 13B and Ci). However during
590 GI-8, NGRIP 2065.65 m is the only tephra layer to coincide with one of these large sulphate
591 peaks that are discernible from the background signal (Figure 13Ci). The oldest tephra in this
592 interval (NGRIP 2066.95 m) is accompanied by a small peak in sulphate of 200 ppbw and no
593 discernible signal is evident for NGRIP 2064.55 m (younger tephra). Large sulphate spikes in
594 this interval e.g. at 2067.36 m do not coincide with glass shard deposition.

595

596 On the other hand, during the cold stadials of the last glacial, the amount of continental dust
597 and other impurities in the ice are 10-20 times higher (see Figure 13B), giving a slightly
598 alkaline signal to the ice. Most of the sulphate present in the ice during these times is derived

599 from CaSO₄ resulting from increased calcium ions caused by increased dustiness, which
600 reacts with the naturally occurring sulphate in the ice. The background levels of sulphate are
601 about 5 times higher than in interstadials, as is the natural variability. As such the increased
602 input of sulphate from a volcanic eruption is masked and is not always a useful indicator.
603 Therefore during cold stadial periods elevated sulphate values due to increased dustiness or
604 volcanism are difficult to be discerned (Steffensen, 1995; Svensson et al., 2013). During the
605 cold stadial of GS-9, the four tephra deposits present coincide with a peak in sulphate and
606 other chemical indicators (Figure 13Cii). However, the sulphate is no greater than the general
607 background level, which suggests that, whilst there is a link between the tephra deposition
608 and the sulphate record, this would not be diagnostic if considering the sulphate record alone.

609

610 Our results suggest that the relationship between tephra deposition and coeval volcanic
611 aerosol fallout is complex and it is unclear whether or not it is solely controlled by prevailing
612 climatic conditions, which supports similar findings reported in Davies et al., (in press).
613 Tephra deposits do fall in association with increased levels of the chemical indicators but the
614 records are so variable that it is difficult to know whether or not they are related to each other
615 or whether it's just coincidental. Therefore it is recommended that future tephra sampling be
616 guided by time periods of interest and not peaks in the chemical records.

617

618 **Conclusions**

619 A detailed Greenland ice-core tephrochronological framework for GS-3 to GI-12 (25,000-
620 45,000 yrs b2k) has been outlined. This framework builds on the work of Davies et al.,
621 (2010) and Bourne et al., (2013) and includes 99 geochemically characterised tephra deposits
622 identified within the NGRIP, NEEM, GRIP and DYE-3 ice-cores. An examination of the
623 relationship between tephra shards and chemical composition of the ice shows that, whilst

624 tephra deposits do occur with small peaks in sulphate, this is not a sufficient diagnostic to use
625 as an indicator of the presence of tephra deposits.

626

627 This study improves our understanding of Icelandic volcanic history and is a crucial first step
628 to facilitate the synchronisation of the Greenland ice-cores with other palaeoclimatic
629 archives. In particular, fourteen tephra deposits are traced in at least 2 ice-cores (Figure 8)
630 and their extensive nature adds value as potential isochrons. In addition, 19 tephra deposits
631 constrain the rapid warming and cooling transitions that characterise this time period and 10
632 of these are geochemically distinct (Figures 11 and 12) also revealing their value as
633 isochrons. Therefore tephra deposits outlined in both Figure 8 and 11 should be an important
634 focus for tracing these cryptotephra deposits in distal and high sedimentation areas of the
635 North Atlantic region, where some of the Greenland tephra layers may also be preserved.

636

637 Given the large number of basaltic tephra layers present in the Greenland ice-cores it would
638 be particularly beneficial to employ extraction methods such as the magnetic separation
639 technique that also allow the identification of basaltic cryptotephra deposits within mineral-
640 rich marine and terrestrial sediments (e.g. Griggs et al., in press). Once identified, potential
641 correlatives to the tephra deposits described here also require robust geochemical
642 characterisation for rigorous comparison to the ice-core tephra deposits, to ensure that they
643 are correlated robustly.

644

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673 uncertainty modelling an INQUA INTAV-led project (International Focus Group on
674 Tephrochronology and Volcanism, project no. 0907).

675

676 **Figure Captions**

677

678 Figure 1 - Location of NGRIP, NEEM, GRIP and DYE-3 ice cores relative to the Katla and
679 Grimsvötn volcanoes. The altitude of each core site is also given.

680

681 Figure 2 - Stratigraphic position of tephra horizons identified within four different Greenland
682 ice-cores. Each tephra deposit is represented by a red line and plotted against the oxygen
683 isotope stratigraphy for each core. Also shown is the NGRIP $\delta^{18}\text{O}$ curve plotted on the
684 GICC05 timescale for 25-45 ka b2k (Andersen *et al.*, 2006). The four time periods used to
685 discuss the tephra deposits in the text are also shown. The Greenland event stratigraphy is
686 shown alongside the oxygen isotope record with GI (interstadial) events shown according to
687 Rasmussen *et al.* (in review).

688

689 Figure 3 Geochemical results for the new tephra deposits identified in period 1 (25-32 ka
690 b2k) A) Total Alkalis vs. Silica diagram (Le Bas *et al.*, 1986). SiO_2 vs. FeO/TiO_2 biplots for
691 the B) NGRIP deposits, C) NEEM deposits, D) GRIP deposits and E) DYE-3 deposits.
692 Geochemical fields for Icelandic source volcanoes are based on data presented in Jakobsson
693 (1979; 2008), Boyle (1994), Hunt *et al.* (1995), Dugmore and Newton (1998), Haflidason *et al.*
694 (2000) and references within, Davies *et al.* (2001), Wastegård *et al.* (2001, 2006), Larsen
695 *et al.* (2002), Andrews *et al.* (2002), Mortensen *et al.* (2005), Óladóttir *et al.* (2008, 2011a and
696 b). F) FeO vs. CaO biplot for the transitional alkali layers in GRIP, NGRIP and NEEM. G)
697 FeO/MgO vs. TiO_2 for the tholeiitic layers in GRIP, NGRIP and NEEM. Data shown are

698 normalised values. Error bars represent 2 standard deviations of replicate analyses of the
699 BCR2G reference glass.

700

701 Figure 4 – Geochemical results for glass shard analyses from mafic and silicic deposits.
702 Geochemical fields are adapted from Meara (2012). Data shown are normalised values.
703 Error bars represent 2 standard deviations of replicate analyses of the Lipari Obsidian
704 reference glass.

705

706 Figure 5 — Major element biplots for all tephra deposits identified during period 2 (32-37 ka
707 b2k). A) Total Alkalis vs. Silica diagram (Le Bas et al., 1986). B) Al_2O_3 vs. TiO_2 biplot C)
708 K_2O vs. TiO_2 biplot and D) SiO_2 vs. FeO/TiO_2 biplot. Geochemical fields for Icelandic
709 source volcanoes are based on data presented in Jakobsson (1979; 2008), Boygle (1994),
710 Hunt et al. (1995), Dugmore and Newton (1998), Haflidason et al. (2000) and references
711 within, Davies et al. (2001), Wastegård et al. (2001, 2006), Larsen et al. (2002), Andrews et
712 al. (2002), Mortensen et al. (2005), Óladottir et al. (2008, 2011a and b). Data shown are
713 normalised values. Error bars represent 2 standard deviations of replicate analyses of the
714 BCR2G reference glass.

715

716 Figure 6 Major element biplots for all new tephra deposits identified during period 3 (37-41
717 ka b2k). A) Total Alkalis vs. Silica diagram (Le Bas et al., 1986). B) SiO_2 vs. FeO/TiO_2
718 biplot. C) Al_2O_3 vs. TiO_2 biplot with an inset to show variation within the Grimsvötn field.
719 Geochemical fields for Icelandic source volcanoes are based on data presented in Jakobsson
720 (1979; 2008), Boygle (1994), Hunt et al. (1995), Dugmore and Newton (1998), Haflidason et
721 al. (2000) and references within, Davies et al. (2001), Wastegård et al. (2001, 2006), Larsen
722 et al. (2002), Andrews et al. (2002), Mortensen et al. (2005), Óladottir et al. (2008, 2011a and

723 b). Data shown are normalised values. Error bars represent 2 standard deviations of replicate
724 analyses of the BCR2G reference glass.

725

726 Figure 7 Major element biplots for all tephra deposits identified during period 4 (41-46ka
727 b2k) A) Total Alkalis vs. Silica diagram (Le Bas et al., 1986) B) Al_2O_3 vs. TiO_2 biplot, C)
728 SiO_2 vs. FeO/TiO_2 biplot and D) CaO vs. FeO . Geochemical fields for Icelandic source
729 volcanoes are based on data presented in Jakobsson (1979; 2008), Boygle (1994), Hunt et al.
730 (1995), Dugmore and Newton (1998), Haflidason et al. (2000) and references within, Davies
731 et al. (2001), Wastegård et al. (2001,2006), Larsen et al. (2002), Andrews et al. (2002),
732 Mortensen et al. (2005), Óladottir et al. (2008, 2011a and b). Data shown are normalised
733 values. Error bars represent 2 standard deviations of replicate analyses of the BCR2G
734 reference glass.

735

736 Figure 8 The Greenland tephra lattice highlighting the most extensive deposits that can be
737 traced in at least two cores. The deposits shown in red can be traced between all 3 cores,
738 those in green correlate between NGRIP and NEEM and those in orange correlate between
739 NGRIP and GRIP. The positions of other tephra deposits found in just one core are also
740 shown. Tephra correlations are based on results outlined in this study, Rasmussen et al (in
741 press) and Seierstad et al., (in review) (see Table 3). The Greenland event stratigraphy and
742 NGRIP $\delta^{18}\text{O}$ curve plotted on the GICC05 timescale (Andersen et al., 2006) GI (interstadial)
743 and GS (stadial) events are shown according to Rasmussen et al. (in review).

744

745 Figure 9 Geochemical biplots that support the new tephra correlations between ice-cores
746 shown in Figure 8 and Table 3. A) Period 1 correlations: i) TiO_2 vs. CaO biplot and ii) MgO
747 vs. CaO biplot for GRIP 2049.50 m (this study), NEEM 1656.50 m (this study) and NGRIP

748 1895.24 m of Davies et al., (2010); GRIP 2060.85 m (this study) to NGRIP 1908.70 m and
749 NEEM 1664.95 (Rasmussen et al., 2013) and GRIP 2079.40 (this study) to NEEM 1677.60 m
750 and NGRIP 1929.95 m (Rasmussen et al., 2013). B) Period 2 correlation between NEEM
751 1702.45 m and NGRIP 1873.16 m (this study) Bi) Al₂O₃ vs. TiO₂ biplot and Bii) FeO/MgO
752 vs. TiO₂ biplot. C) Period 3 correlations Ci) CaO vs. TiO₂ and Cii) FeO/MgO vs. TiO₂.
753 NEEM and NGRIP data presented are from Bourne et al., (2013). Data shown are normalised
754 values. Error bars represent 2 standard deviations of replicate analyses of the BCR2G
755 reference glass.

756

757 Figure 10 A) Tephra grain-size data for each deposit relative to distance of core locations
758 from Katla. B) Tephra grain-size data for each deposit relative to distance of core locations
759 from Grimsvötn.

760

761 Figure 11 – A Greenland tephrochronology framework for 25-45 ka b2k highlighting those
762 tephtras that are geochemically distinct (bold italic type) from other deposits of similar age
763 and those that fall close to rapid climatic events. Tephra layers are highlighted that lie on a
764 sharp transition in the Greenland event stratigraphy of Rasmussen et al. (in review).

765

766 Figure 12 – Geochemical comparisons of tephra deposits that fall on climatic transitions
767 relative to other tephra layers of similar age (see stratigraphic positions in Figure 11). A)
768 Tephra deposits in GS-3 and GI-3; B) Tephra deposits in GI-4 and GS-5.1; C) Tephra
769 deposits in GS-5.1 and GI-5.1; D) Tephra deposits in GI-5.2 and GS-6, E) Tephra deposits in
770 GI-6 and GI-7; F) Tephra deposits in GI-8 and GS-9 and G) Tephra deposits in GI-9, GS-10
771 and GI-10. Data shown are normalised values. Error bars represent 2 standard deviations of

772 replicate analyses of the BCR2G reference glass for basaltic layers and of the Lipari Obsidian
773 for silicic layers.

774

775 Figure 13 – NGRIP cryptotephra positions plotted alongside chemostratigraphical data. A)
776 Electrical Conductivity measurement (ECM) for 25-45 ka b2k. B) Dust, Calcium, Sulphate
777 and Conductivity measurements between GI-8 and GI-9 and C) Expanded interstadial (Ci)
778 and stadial section (Cii). Sulphate, calcium, electrolytic meltwater conductivity and dust
779 analyses have been measured by the continuous flow analysis (CFA) system. Tephra
780 positions are shown by the red lines and shading and ice sections sampled for tephra content
781 in A) are shown by grey shading.

782

783 Table 1: Summary table of tephra deposits identified in each ice-core within the different
784 time periods. Number of tephra deposits already published and noted in parentheses are from
785 Period 1: Davies et al., (2010) and Period 3: Bourne et al., (2013). Grain size was measured
786 using a graticule in the eye-piece of a transmitted light microscope.

787

788 Table 2: Tephra framework for the Greenland ice-cores spanning 25-45 ka b2k. For each
789 tephra the following information is provided: depth interval of ice sampled, shard numbers
790 identified per sample, climatic event within which tephra was deposited (according to
791 Rasmussen et al., in review), age, grain-size data, geochemical composition and most likely
792 volcanic source. Shard numbers are given for each sample but are not directly comparable
793 with one another due to differences in sample volume. Shading highlights 2 or 3 layers from
794 different ice-cores that have been correlated according to Rasmussen et al., (2013) (denoted
795 by #) and Seierstad et al., (in review) (denoted by ^), any unmarked shaded layers represent
796 new correlations discussed here and outlined in Table 3 and Figure 8. The climatic event is

797 defined based upon the event stratigraphy presented in Rasmussen et al., (in review). Ages
798 are in b2k (before 2000 CE) and represent the age of the basal depth of the ice sample
799 containing the glass shards. The ages are obtained from the GICC05 timescale in steps of 20
800 years for the NGRIP core (Andersen et al., 2006, Svensson et al., 2006, 2008) and the GRIP
801 core (Seierstad et al., in review) and in steps of 0.55 cm for the NEEM core (Rasmussen et
802 al., 2013). DYE-3 ages are approximate ages based on wiggle-matching of $\delta^{18}\text{O}$ to NGRIP.
803 MCE = maximum counting error; in a standard deviation context, the maximum counting
804 error should be regarded as 2 sigma (Andersen et al., 2006; Rasmussen et al., 2006). Ages
805 for deposits which have been traced and correlated between cores (indicated by shading) may
806 differ because 1) they are basal ages for the sample within which glass shards were identified,
807 2) that the exact position of the tephra horizon within the ice sample is unknown, 3) the depth
808 range of the sample is different from core to core, and 4) due to the uncertainty on the
809 timescale transfer from NGRIP to GRIP and NEEM. Chemical composition after Le Bas et
810 al., (1986): TB = Tholeiitic Basalt, TAB = Transitional Alkali Basalt, B = Basalt, R =
811 Rhyolite, Da = Dacite, TRDA = Trachydacite. † = Tephra deposits published by Davies et al.,
812 (2010) and * = Tephra deposits published by Bourne et al., (2013).

813

814 Table 3: Summary of tephra horizons that have been correlated between different ice-core
815 records from Rasmussen et al., (2013), Seierstad et al. (submitted) and this study. Similarity
816 coefficient (SC) and statistical distance (SD) comparisons for tephra horizon correlations are
817 presented. The similarity coefficient method is from Borchardt et al., (1972) and Hunt et al.,
818 (1995) and 7 major elements were used in the comparisons. The statistical distance method is
819 from Perkins et al., (1998; 1995) and 10 major elements were used in the comparisons.
820 Critical value for testing the statistical distance values at the 99% confidence interval is 23.21

821 (10 degrees of freedom). Correlations highlighted in **bold** were used as time-scale transfer
822 points in the respective studies.

823

824 Supplementary Data: Major element data for each tephra deposit analysed in this study. Data
825 are separated into four worksheets according to periods 1-4 outlined in the main text. The
826 date of analysis and beam size are given. EPMA operating conditions are adapted from
827 Hayward (2012) and vary by beam size and are as follows: 5 µm beam diameter –
828 Accelerating voltage: 15 kV Beam Current: 2 nA for Na, K, Si, Al, Mg, Fe, Ca and 80 nA for
829 Mn, Ti, P. 3 µm beam diameter – Accelerating voltage: 15 kV Beam Current: 0.5 nA for Na,
830 Al, 2 nA for K, Si, Mg, Fe, Ca and 60 nA for Mn, Ti, P. Analyses of the reference standard
831 glasses BCR2G and Lipari are given in the Standard data sheet. They are ordered by date of
832 analysis and were conducted throughout the analytical period. Recommended values for the
833 Lipari from Kuehn *et al.* (2011) and for BCR2G from
834 http://minerals.cr.usgs.gov/geo_chem_stand/basaltbcr2.html (accessed 12/06/13) are given.

835

836

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Table1

	Ice Core	Total number of tephra deposits (published)	Ice Sampled (m)	% of Total Ice in Period	Tephra average grain size (μm)
PERIOD 1 25-32 ka b2k	NEEM	9	30.80	44	27.1
	NGRIP	8 (6)	55.55	46	33.5
	GRIP	12	52.80	48	36.8
	DYE-3	8	15.40	40	57.3
PERIOD 2 32-37 ka b2k	NEEM	4	26.40	37	18.6
	NGRIP	5	50.05	40	30.6
	GRIP	0	0	0	N/A
	DYE-3	0	0.00	0	N/A
PERIOD 3 37-41 ka b2k	NEEM	2 (5)	26.40	86	25.0
	NGRIP	0 (15)	45.65	91	29.3
	GRIP	10	45.10	100	45.6
	DYE-3	7	18.70	100	54.6
PERIOD 4 41-46 ka b2k	NEEM	0	8.80	18	N/A
	NGRIP	8	30.25	34	28.1
	GRIP	0	0.00	0	N/A
	DYE-3	0	0.00	0	N/A

Table2

	Tephra layer	Depth Range (m)	Shards per Sample	Climatic Event	Age of base ± MCE (yr b2k)	Average Grain Size (µm)	Max Grain Size (µm)	M
PERIOD 1	NEEM 1626.15 m	1626.1-1626.15	1055	GS-3	26439 ± 766	22.0	80.0	
	GRIP 2002.20 m	2002.00-2002.20	361	GS-3	26544 ± 768	50.9	67.5	
	NGRIP 1848.05 m [†]	1848.00-1848.05	Visible	GS-3	26743 ± 780	38.8	62.5	
	NGRIP 1855.80 m [†]	1855.70-1855.80	24	GS-3	27198 ± 804	45.5	62.5	
	NEEM 1636.45 m	1636.25-1636.45	19	GS-3	27510 ± 820	28.0	60.0	
	NEEM 1636.65 m	1636.45-1636.65	56	GS-3	27528 ± 821	30.0	60.0	
	NGRIP 1861.55 m [†]	1861.45-1861.55	103	GS-3	27534 ± 821	37.3	52.5	
	NGRIP 1882.10 m [#]	1881.95-1882.10	51	GS-4	28575 ± 886	23.3	32.5	
	NEEM 1648.90 m [#]	1648.75-1648.90	214	GS-4	28578 ± 885	36.0	70.0	
	NGRIP 1882.50 m	1882.30-1882.50	48	GS-4	28594 ± 887	33.5	50.0	
	NGRIP 1888.10 m	1888.05-1888.10	70	GI-4	28789 ± 893	29.5	47.5	
	NGRIP 1894.05 m	1893.85-1894.05	25	GS-5.1	29048 ± 905	46.6	80.0	
	NGRIP 1895.24 m ^{†#}	1895.23-1895.24	Visible	GS-5.1	29132 ± 912	44.0	75.0	
	NEEM 1656.50 m [#]	1656.45-1656.50	2004	GS-5.1	29135 ± 911	40.0	105.0	
	GRIP 2049.50 m	2049.30-2049.50	1747	GS-5.1	29147 ± 912	50.8	80.0	
	GRIP 2060.85 m	2060.70-2060.85	429	GS-5.1	30066 ± 976	33.3	45.0	
	NGRIP 1908.70 m [#]	1908.50-1908.70	250	GS-5.1	30082 ± 977	26.3	47.5	
	NEEM 1664.95 m [#]	1664.85-1664.95	80	GS-5.1	30083 ± 977	16.3	30.0	
	GRIP 2061.40 m	2061.25-2061.40	11	GS-5.1	30111 ± 978	33.3	45.0	
	GRIP 2064.35 m	2064.15-2064.35		GS-5.1	30353 ± 993	23.0	35.0	
NGRIP 1913.10 m	1912.90-1913.10	1028	GS-5.1	30394 ± 995	11.3	20.0		
GRIP 2066.75 m	2066.55-2066.75	288	GS-5.1	30551 ± 1005	36.2	65.0		
NGRIP 1915.50 m ^{†#}	1915.10-1915.50	92	GS-5.1	30565 ± 1006	42.0	60.0		

	Tephra layer	Depth Range (m)	Shards per Sample	Climatic Event	Age of base ± MCE (yr b2k)	Average Grain Size (µm)	Max Grain Size (µm)	M Si
PERIOD 1	NGRIP 1915.63 m ^{†#}	1915.50-1915.63	84	GS-5.1	30573 ± 1007	47.0	77.5	
	NEEM 1669.25 m [#]	1669.10-1669.25	188	GS-5.1	30590 ± 1007	25.1	40.0	
	GRIP 2067.85 m	2067.65-2067.85	55	GI-5.1	30628 ± 1010	32.2	47.5	
	GRIP 2070.20 m	2070.05-2070.20	74	GI-5.1	30779 ± 1022	28.5	50.0	
	GRIP 2070.75 m	2070.60-2070.75	6	GI-5.1	30813 ± 1023	53.2	80.0	
	NEEM 1671.85 m	1671.65-1671.85	135	GI-5.1	30825 ± 1023	25.7	55.0	
	GRIP 2079.40 m [^]	2079.00-2079.40	1499	GS-5.2	31414 ± 1066	51.3	87.5	
	NGRIP 1929.95 m ^{^#}	1929.80-1929.95	130	GS-5.2	31432 ± 1067	32.1	45.0	
	NEEM 1677.60 m [#]	1677.50-1677.60	69	GS-5.2	31433 ± 1067	21.3	35.0	
	NGRIP 1931.60 m [^]	1931.45-1931.60	45	GS-5.2	31543 ± 1074	22.5	30.0	
	GRIP 2081.05 m [^]	2080.85-2081.05	11	GS-5.2	31555 ± 1076	30.0	45.0	
	GRIP 2081.40 m	2081.20-2081.40	183	GS-5.2	31581 ± 1078	34.3	57.5	
	DYE-3 1865.70 m	1865.60-1865.70	114	GI-4	28700 ± 1000	59.0	80.0	
	DYE-3 1865.80 m	1865.70-1865.80	38	GI-4	28720 ± 1000	66.9	115.0	
	DYE-3 1865.90 m	1865.80-1865.90	17	GI-4	28740 ± 1000	64.6	77.50	
	DYE-3 1866.00 m	1865.90-1866.00	12	GI-4	28760 ± 1000	56.0	92.5	
	DYE-3 1866.10 m	1866.00-1866.10	12	GI-4	28780 ± 1000	47.5	65.0	
	DYE-3 1866.40 m	1866.30-1866.40	8	GI-4	28800 ± 1000	29.2	52.5	
	DYE-3 1866.60 m	1866.50-1866.60	10	GI-4	28820 ± 1000	73.3	90.0	
DYE-3 1869.15 m	1869.05-1869.15	9	GI-5.1	29400 ± 1000	54.2	137.5		
PERIOD 2	NEEM 1689.25 m [#]	1689.05-1689.25	409	GI-5.2	32459 ± 1130	18.7	37.5	
	NGRIP 1950.50 m [#]	1950.30-1950.50	119	GI-5.2	32463 ± 1130	28.8	47.5	
	NGRIP 1952.15 m [#]	1951.95-1952.15	29	GS-6	32522 ± 1132	23.5	40.0	

	Tephra layer	Depth Range (m)	Shards per Sample	Climatic Event	Age of base ± MCE (yr b2k)	Average Grain Size (µm)	Max Grain Size (µm)	M Si
PERIOD 2	NEEM 1690.35 m [#]	1690.15-1690.35	93	GS-6	32534 ± 1133	14.1	25.0	
	NGRIP 1954.70 m	1954.55-1954.70	125	GS-6	32690 ± 1144	37.1	55.0	
	NEEM 1693.45 m	1693.30-1693.45	15	GS-6	32890 ± 1165	15.8	40.0	
	NGRIP 1973.16 m	1973.12-1973.16	583	GI-6	33686 ± 1207	38.7	75.0	
	NEEM 1702.40 m	1702.35-1702.40	57	GI-6	33692 ± 1208	19.0	37.5	
	NGRIP 2009.15 m	2009.00-2009.15	189	GS-8	35470 ± 1320	24.8	37.5	
PERIOD 3	NEEM 1747.10 m	1746.90-1747.10	317	GI-8c	37548 ± 1429	29.1	57.5	
	GRIP 2190.65 m	2190.50-2190.65	6	GI-8c	37864 ± 1435	32.0	45.0	
	NEEM 1755.60 m [#]	1755.45-1755.60	6	GI-8c	38040 ± 1441	20.0	25.0	
	NGRIP 2064.35 m ^{*#^}	2064.15-2064.35	116	GI-8c	38041 ± 1441	28.9	50.0	
	GRIP 2195.45 m [^]	2195.25-2195.45	9	GI-8c	38043 ± 1441	51.0	72.5	
	NGRIP 2065.65 m [*]	2065.45-2065.65	74	GI-8c	38081 ± 1441	21.1	42.5	
	NGRIP 2065.80 m [*]	2065.65-2065.80	785	GI-8c	38086 ± 1442	21.9	47.5	
	GRIP 2197.45 m	2197.25-2197.45	258	GI-8c	38115 ± 1445	47.8	65.0	
	NEEM 1757.10 m ^{*#}	1756.90-1757.10	19	GI-8c	38119 ± 1445	28.8	50.0	
	NGRIP 2066.95 m ^{*†#}	2066.93-2066.95	Visible	GI-8c	38121 ± 1445			
	GRIP 2200.75 m	2200.55-2200.75	177	GS-9	38249 ± 1450	49.8	97.5	
	GRIP 2201.50 m [^]	2201.10-2201.50	200	GS-9	38307 ± 1452	45.5	90.0	
	NGRIP 2071.50 m ^{*^#}	2071.30-2071.50	1138	GS-9	38309 ± 1452	44.8	72.5	
	NEEM 1759.85 m ^{*#}	1759.65-1759.85	550	GS-9	38311 ± 1452	25.8	50.0	
	GRIP 2202.40 m	2202.20-2202.40	6	GS-9	38371 ± 1456	46.0	90.0	
	NGRIP 2073.15 m [*]	2072.95-2073.15	10	GS-9	38411 ± 1461	17.3	30.0	
NGRIP 2078.01 m ^{*^#}	2077.90-2078.01	32	GS-9	38735 ± 1476	26.2	37.5		

	Tephra layer	Depth Range (m)	Shards per Sample	Climatic Event	Age of base ± MCE (yr b2k)	Average Grain Size (µm)	Max Grain Size (µm)	M Si
PERIOD 3	GRIP 2207.00 m [^]	2206.60-2207.00	194	GS-9	38748 ± 1477	43.3	62.5	
	NEEM 1764.25 m ^{*#}	1764.05-1764.25	12	GS-9	38763 ± 1477	30.8	80.0	
	NGRIP 2078.37 m [*]	2078.30-2078.37	561	GS-9	38759 ± 1478	22.3	42.5	
	NGRIP 2078.97 m [*]	2078.85-2078.97	126	GS-9	38796 ± 1479	29.5	62.5	
	NGRIP 2079.40 m [*]	2079.25-2079.40	115	GS-9	38826 ± 1479	21.7	37.5	
	NGRIP 2081.95 m [*]	2081.75-2081.95	9	GS-9	38993 ± 1491	40.4	60.0	
	NGRIP 2085.80 m [*]	2085.60-2085.80	5421	GS-9	39258 ± 1510	28.1	35.0	
	GRIP 2213.05 m	2212.85-2213.05	176	GS-9	39274 ± 1511	66.9	115.0	
	NGRIP 2100.65 m [*]	2100.45-2100.65	790	GS-10	40218 ± 1583	64.6	67.5	
	NGRIP 2101.55 m [*]	2101.45-2101.55	11	GS-10	40275 ± 1587	35.4	50.0	
	NGRIP 2103.98 m ^{*^#}	2103.92-2103.98	64	GS-10	40428 ± 1595	30.8	50.0	
	GRIP 2227.15 m [^]	2226.95-2227.15	44	GS-10	40433 ± 1596	59.3	107.5	
	NEEM 1780.20 m ^{*#}	1780.00-1780.20	12	GS-10	40449 ± 1596	26.9	35.0	
	GRIP 2227.90 m	2227.70-2227.90	167	GS-10	40498 ± 1599	15.3	37.5	
	NEEM 1784.46 m	1784.45-1784.46	1178	GI-10	40915 ± 1619	13.4	25.0	
	DYE-3 1895.55 m	1895.45-1895.55	6	GI-8	37600 ± 1450	47.5	80.0	
	DYE-3 1898.65 m	1898.60-1898.65	3	GS-9	38500 ± 1480	53.8	107.5	
	DYE-3 1900.80 m	1900.70-1900.80	7	GS-9	39000 ± 1500	40.4	110.0	
	DYE-3 1901.80 m	1901.70-1901.80	10	GS-9	39200 ± 1510	54.8	70.0	
	DYE-3 1904.10 m	1904.00-1904.10	28	GS-9	39700 ± 1550	50.8	80.0	
DYE-3 1904.15 m	1904.10-1904.15	50	GS-9	39800 ± 1560	45.8	70.0		
DYE-3 1912.35 m	1912.25-1912.35	4	GI-11	42300 ± 1700	36.7	70.0		

	Tephra layer	Depth Range (m)	Shards per Sample	Climatic Event	Age of base \pm MCE (yr b2k)	Average Grain Size (μm)	Max Grain Size (μm)	M
PERIOD 4	NGRIP 2150.90 m	2150.70-2150.90	78	GI-11	43066 \pm 1727	26.9	42.5	
	NGRIP 2162.05 m	2161.90-2162.05	40	GS-12	43683 \pm 1753	34.3	50.0	
	NGRIP 2162.60 m	2162.45-2162.60	21	GS-12	43726 \pm 1755	22.5	35.0	
	NGRIP 2163.35 m	2163.15-2163.35	73	GS-12	43783 \pm 1757	30.5	50.0	
	NGRIP 2164.10 m	2163.90-2164.10	61	GS-12	43840 \pm 1761	35.7	55.0	
	NGRIP 2185.70 m	2185.50-2185.70	827	GI-12c	45221 \pm 1827	28.1	50.0	
	NGRIP 2186.80 m	2186.60-2186.80	175	GI-12c	45285 \pm 1830	21.1	30.0	
	NGRIP 2188.25 m	2188.05-2188.25	382	GI-12c	45368 \pm 1836	25.6	45.0	

Table3

	NGRIP depth (m)	NEEM depth (m)	GRIP depth (m)	SC	SD	Period	Age of ba MCE (yr)
1	1881.95 - 1882.10	1648.75 - 1648.90		0.981	0.453	GS-4	28575 ± 1
2	1895.23 - 1895.24	1656.45 - 1656.50		0.979	3.569	GS-5.1	29132 ± 1
	1895.23 - 1895.24		2049.30-2049.50	0.991	2.108	GS-5.1	29132 ± 1
3		1656.45 - 1656.50	2049.30-2049.50	0.983	2.822	GS-5.1	29132 ± 1
	1908.50 - 1908.70	1664.85 - 1664.95		0.985	1.887	GS-5.1	30082 ± 1
	1908.50 - 1908.70		2060.70-2060.85	0.978	2.688	GS-5.1	30082 ± 1
4		1664.85-1664.95	2060.70-2060.85	0.976	2.106	GS-5.1	30082 ± 1
	1915.10-1915.50 1915.50-1915.63	1669.10-1669.25		0.977	2.681	GS-5.1	30565 ± 1
5	1929.80 - 1929.95	1677.50 - 1677.60		0.973	2.771	GS-5.2	31432 ± 1
	1929.80 - 1929.95		2079.00 - 2079.40	0.976	2.941	GS-5.2	31432 ± 1
		1677.50 - 1677.60	2079.00 - 2079.40	0.982	0.555	GS-5.2	31432 ± 1
6	1931.45 - 1931.60		2080.85 - 2081.05	0.941	2.174	GS-5.2	31543 ± 1
7	1950.30 - 1950.50	1689.05 - 1689.25		0.985	0.868	GI-5.2	32463 ± 1
8	1951.95 - 1952.15	1690.15 - 1690.35		0.974	1.691	GS-6	32522 ± 1
9	1973.12-1973.16	1702.35-1702.40		0.987	2.857	GI-6	33686 ± 1
10	2064.15 - 2064.35	1755.45 - 1755.60		0.985	2.014	GI-8c	38041 ± 1
	2064.15 - 2064.35		2195.25 - 2195.45	0.986	1.746	GI-8c	38041 ± 1
		1755.45 - 1755.60	2195.25 - 2195.45	0.982	1.448	GI-8c	38041 ± 1
11	2066.95 - 2066.95	1756.90 - 1757.10		0.995	4.151	GI-8c	38121 ± 1
		1756.90 - 1757.10	2197.25-2197.45	0.988	1.255	GI-8c	38121 ± 1
12	2071.30 - 2071.50	1759.65 - 1759.85		0.982	1.814	GS-9	38309 ± 1
	2071.30 - 2071.50		2201.10 - 2201.50	0.972	1.103	GS-9	38309 ± 1
		1759.65 - 1759.85	2201.10 - 2201.50	0.982	0.521	GS-9	38309 ± 1
13	2077.90 - 2078.01	1764.05 - 1764.25		0.985	0.870	GS-9	38735 ± 1
	2077.90 - 2078.01		2206.60 - 2207.00	0.985	1.971	GS-9	38735 ± 1
		1764.05 - 1764.25	2206.60 - 2207.00	0.977	1.063	GS-9	38735 ± 1
14	2103.92 - 2103.98	1780.00 - 1780.20		0.954	6.600	GS-10	40428 ± 1
	2103.92 - 2103.98		2227.05 - 2227.10	0.975	0.818	GS-10	40428 ± 1
		1780.00 - 1780.20	2227.05 - 2227.15	0.977	1.614	GS-10	40428 ± 1

Figure1

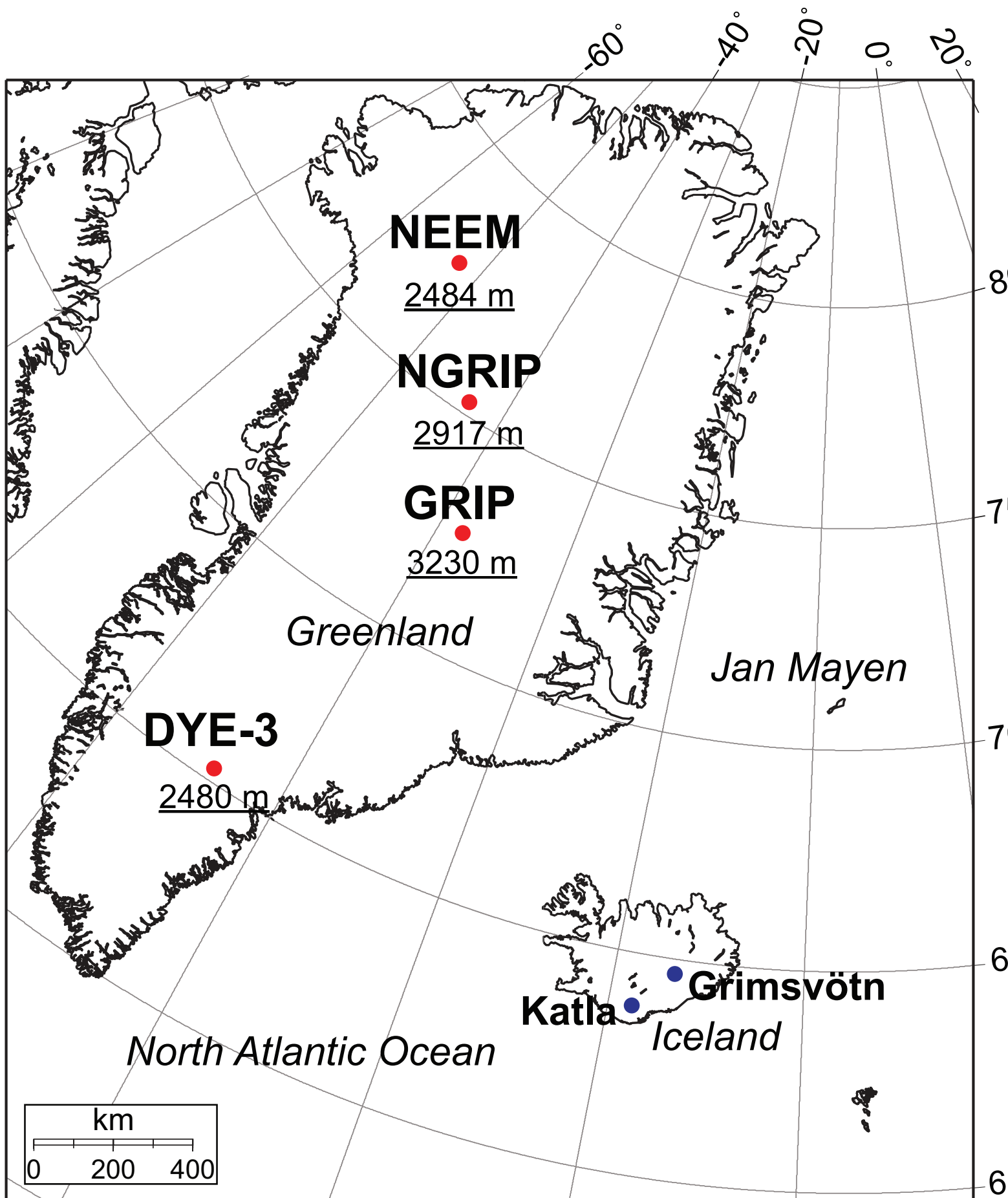
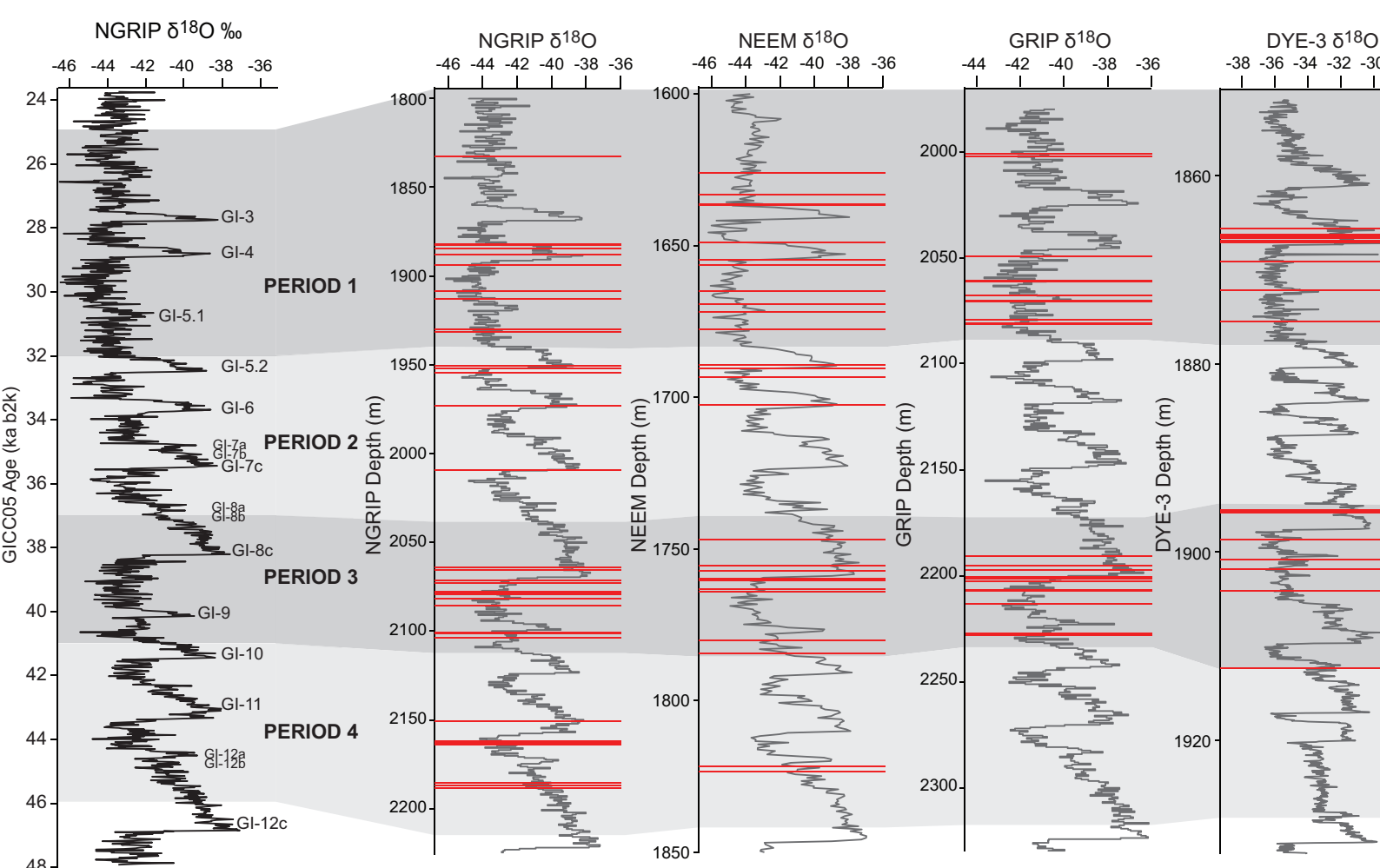


Figure2



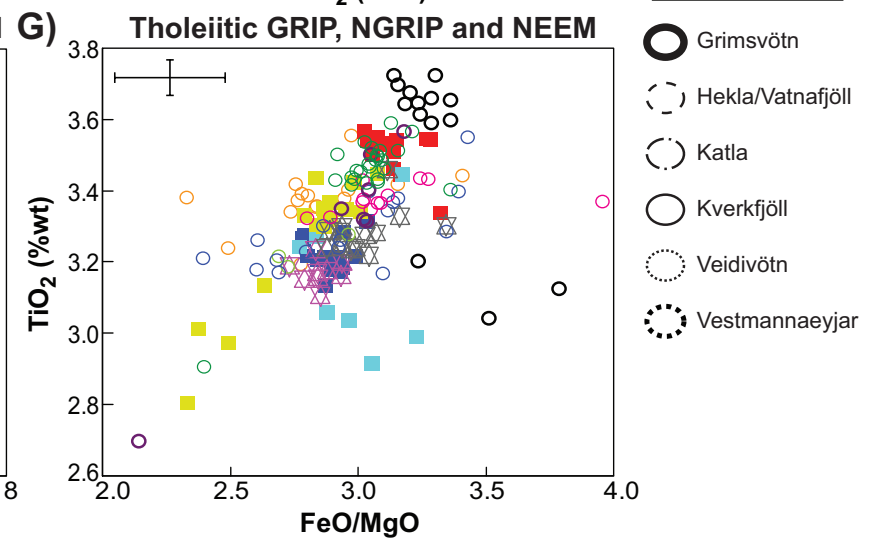
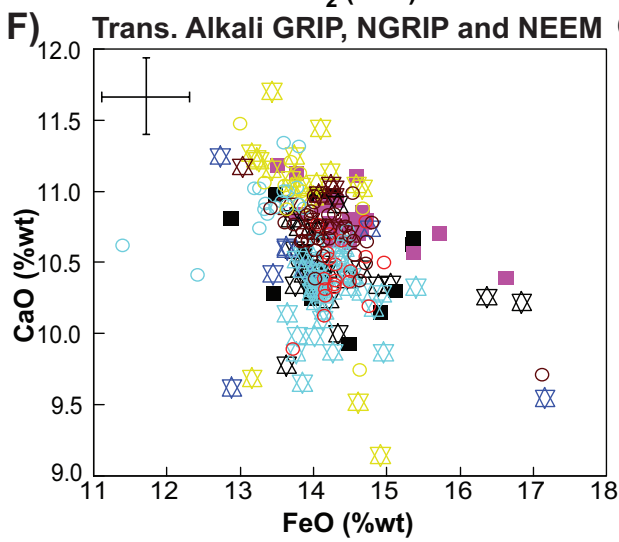
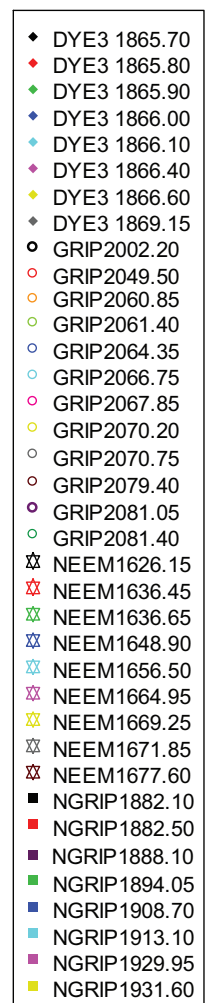
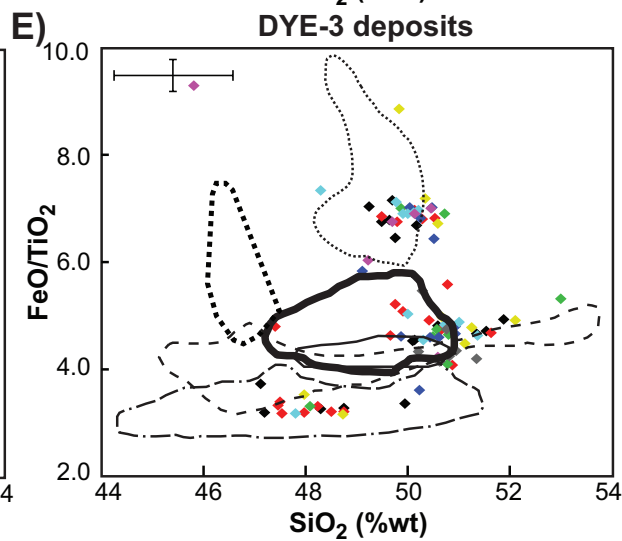
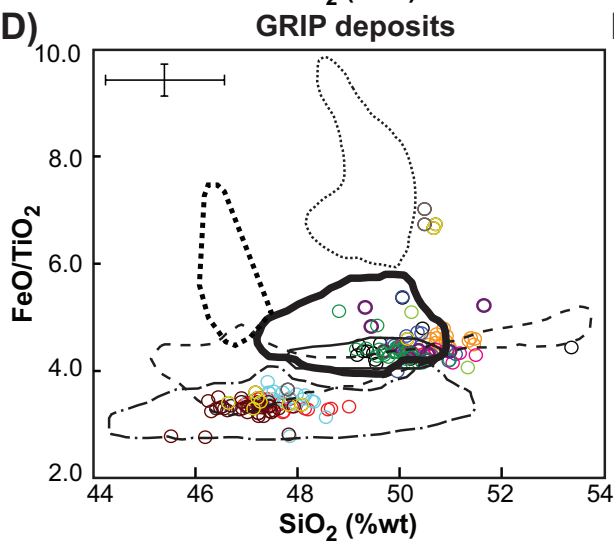
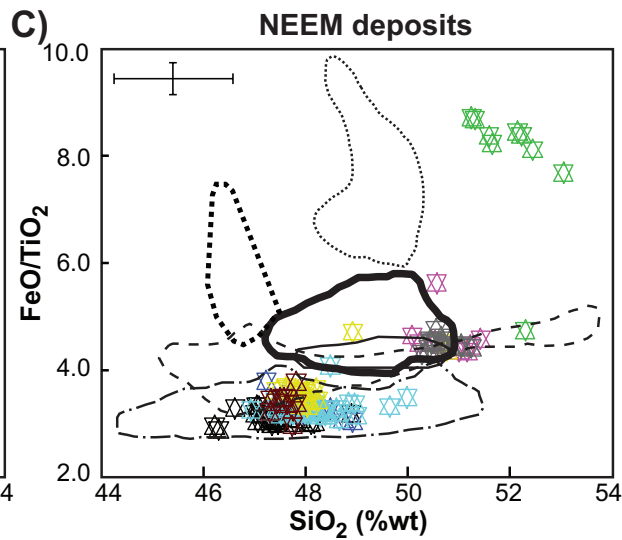
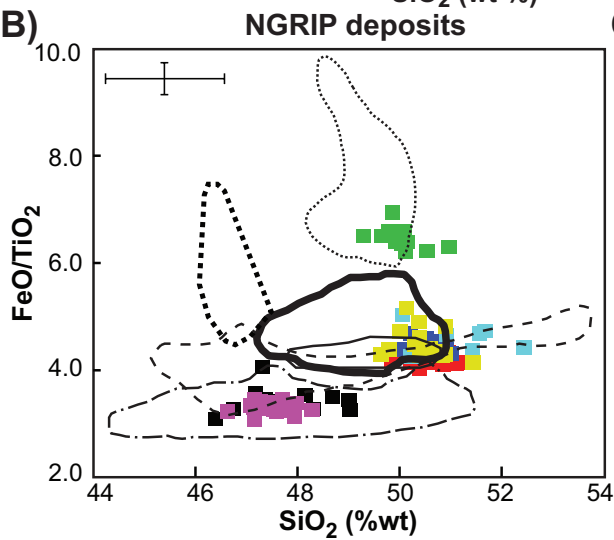
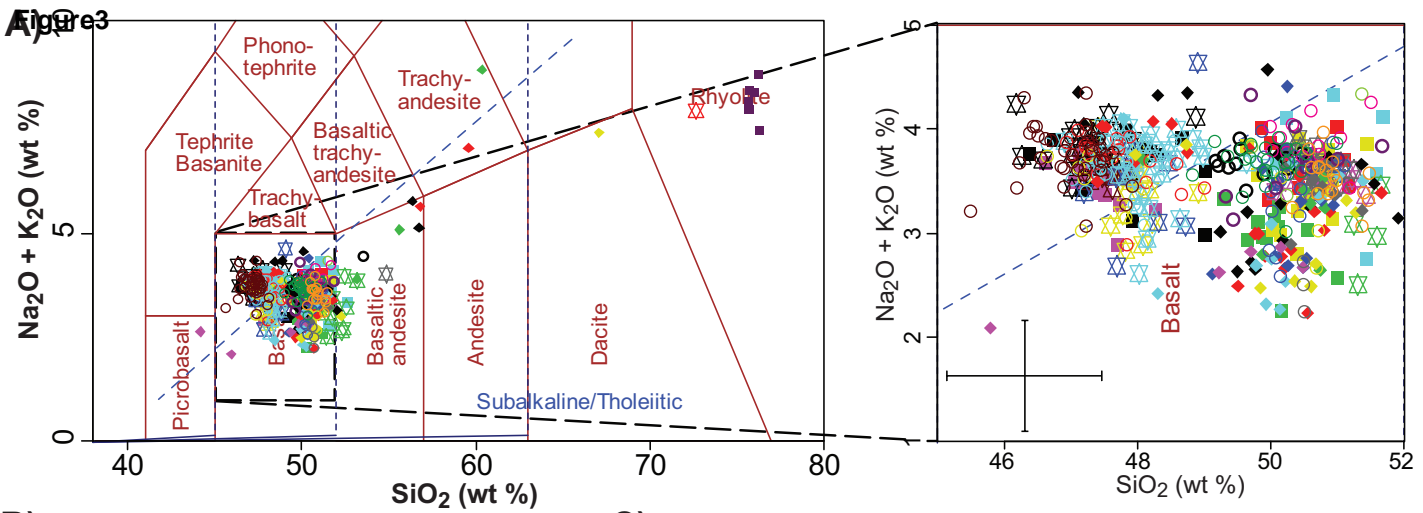


Figure4

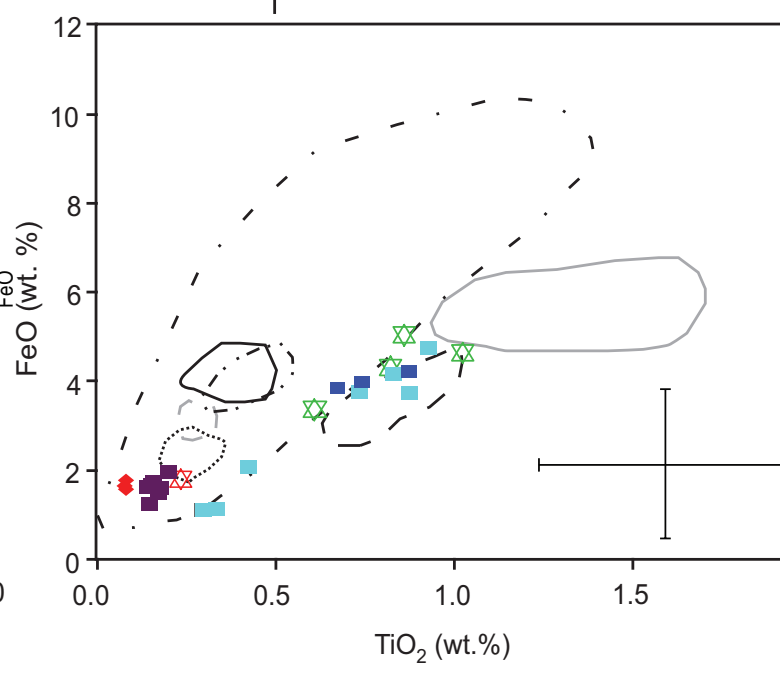
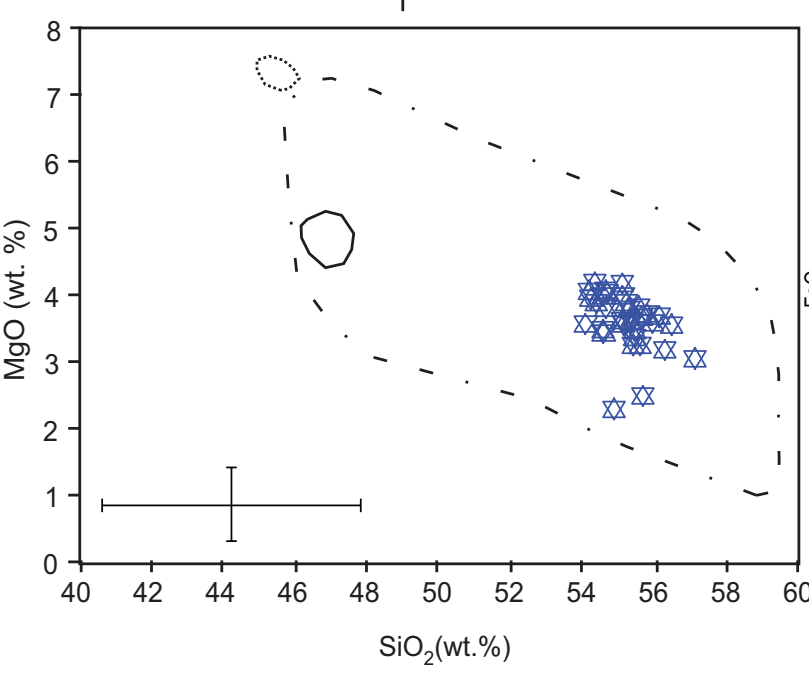
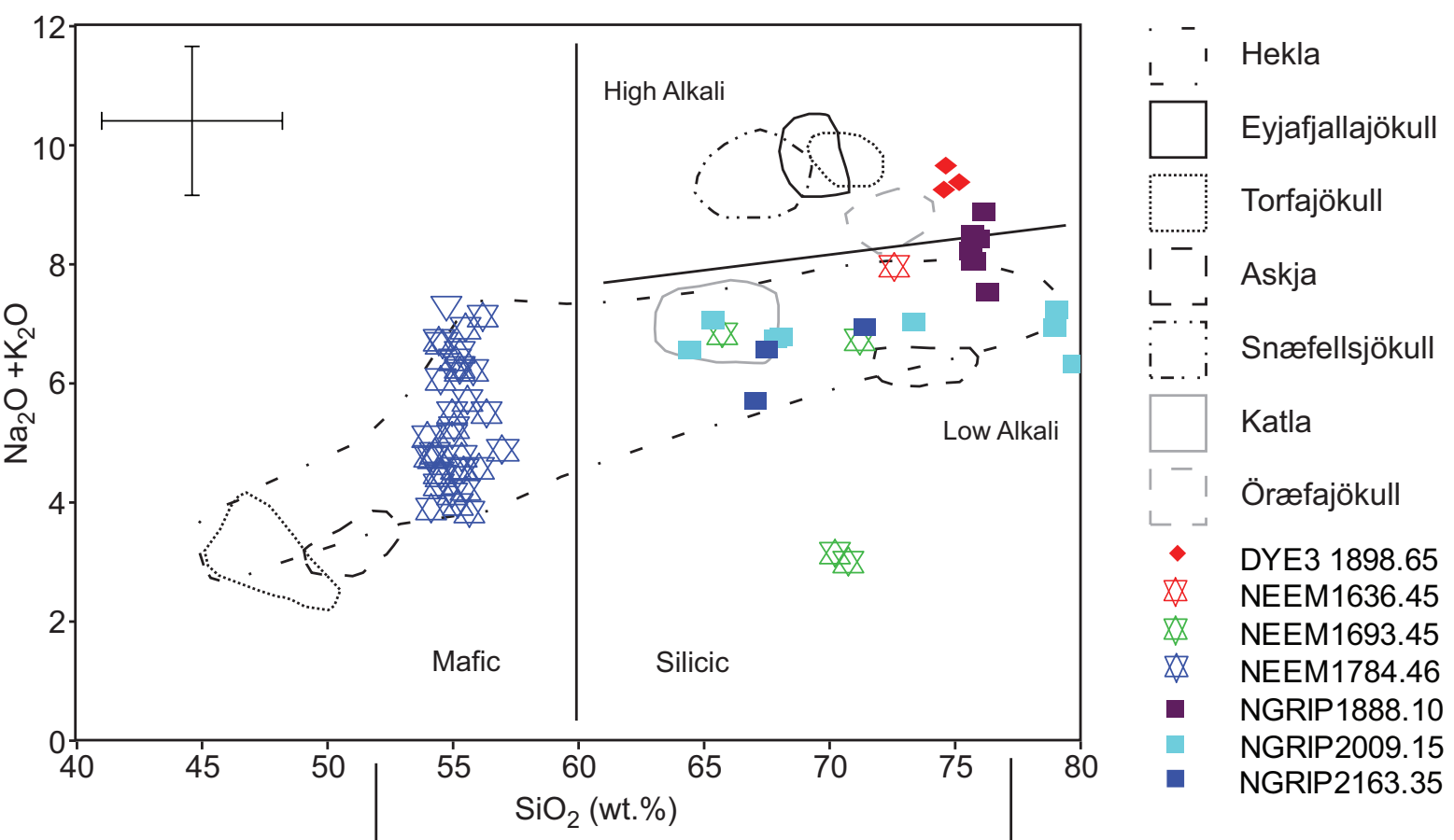


Figure 5

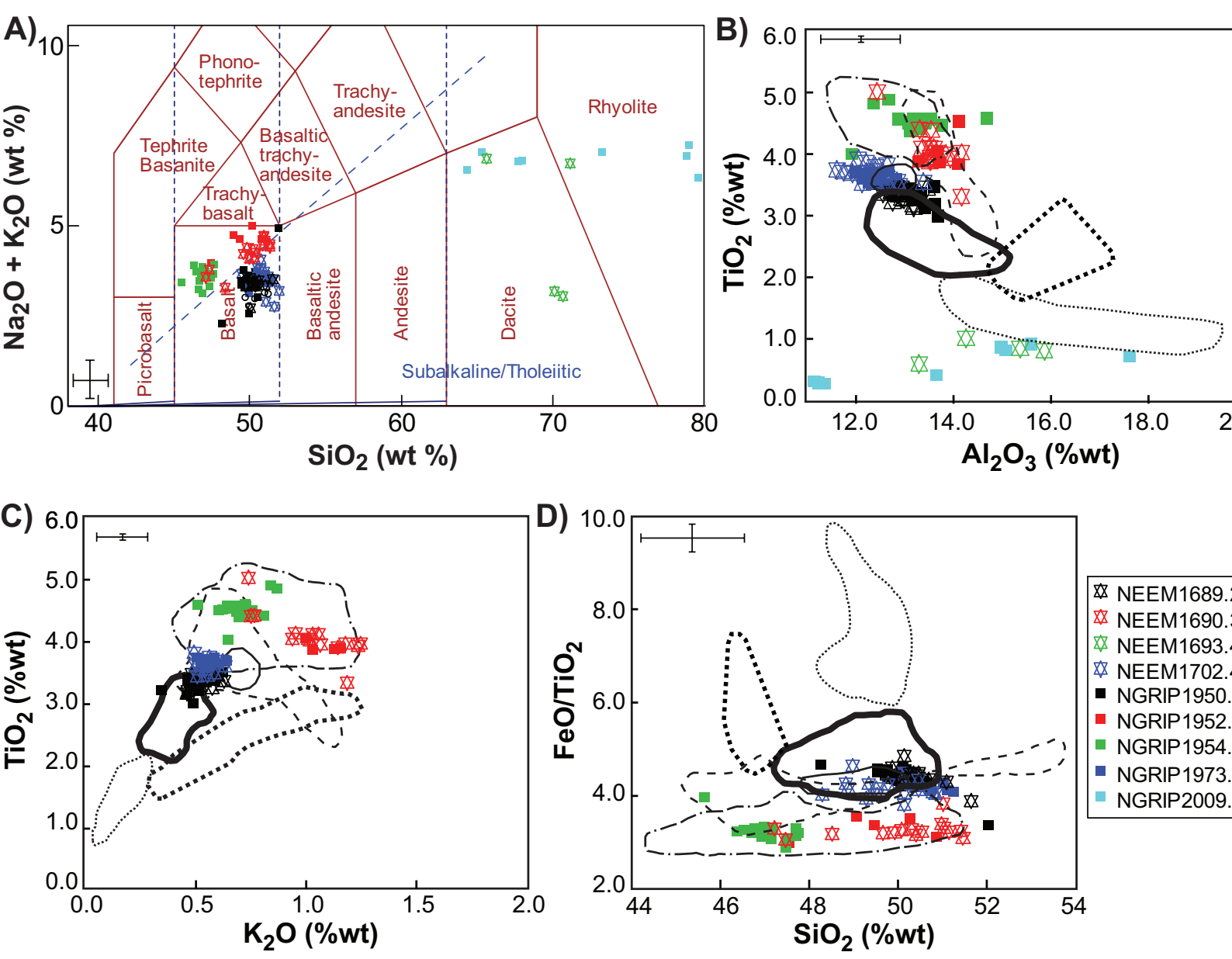


Figure 6

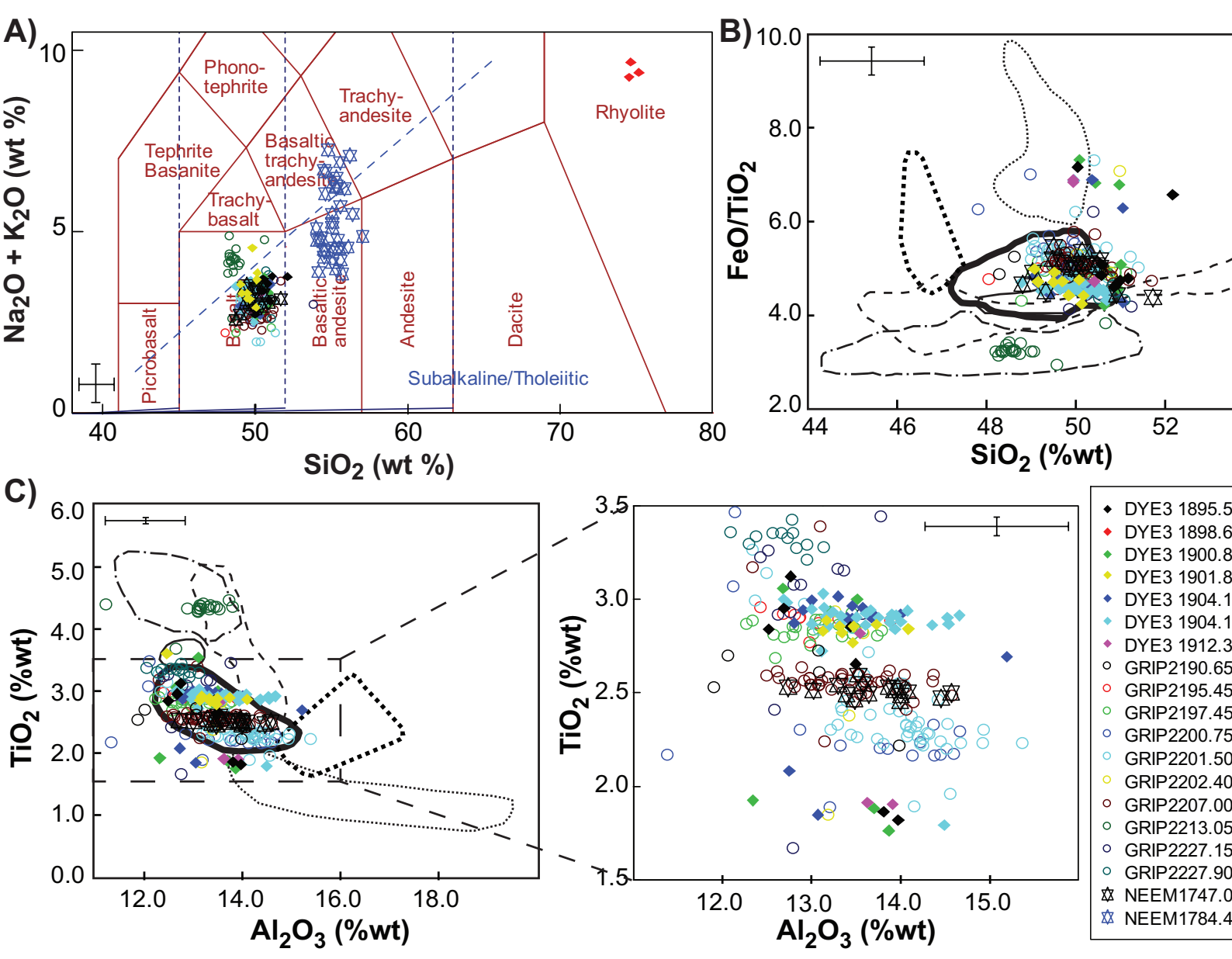


Figure 7

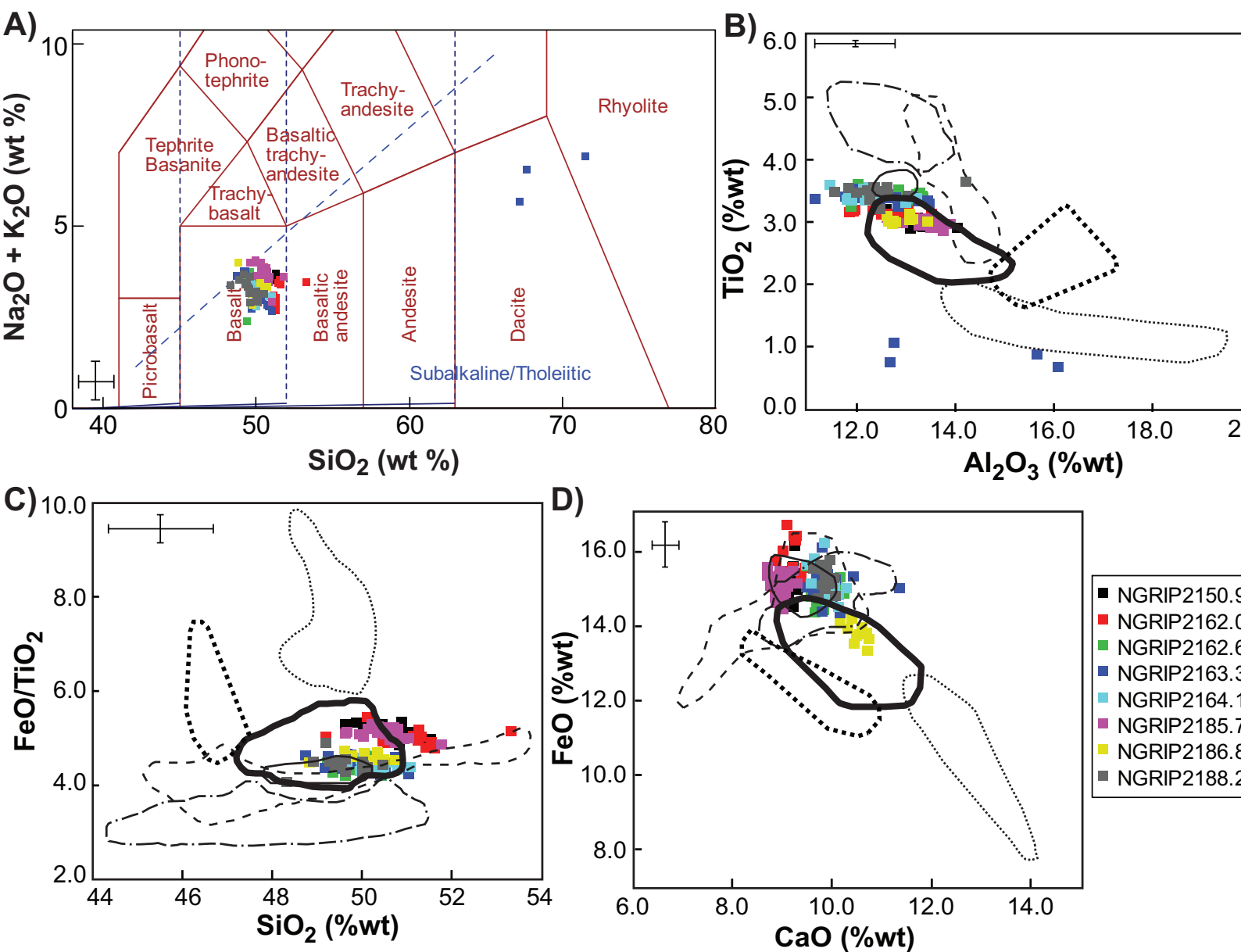
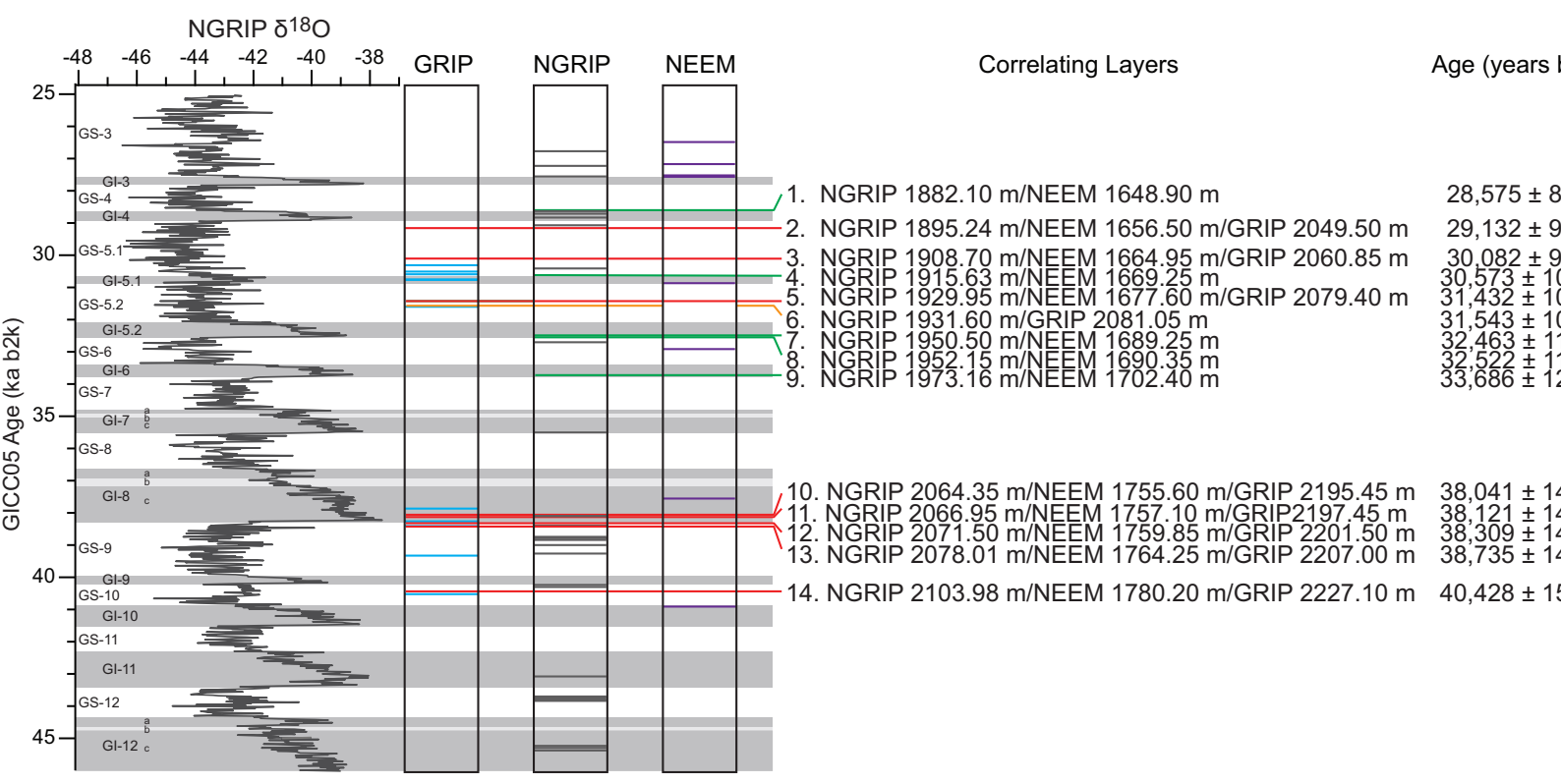
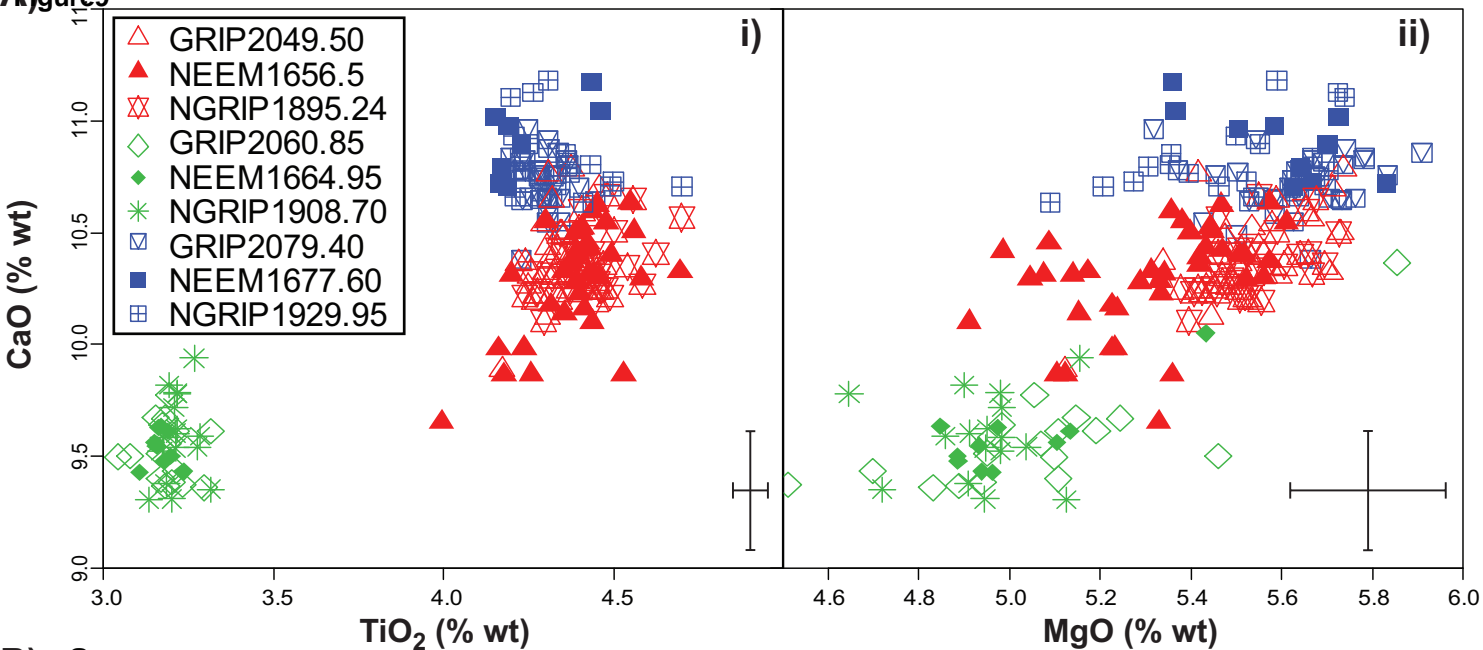


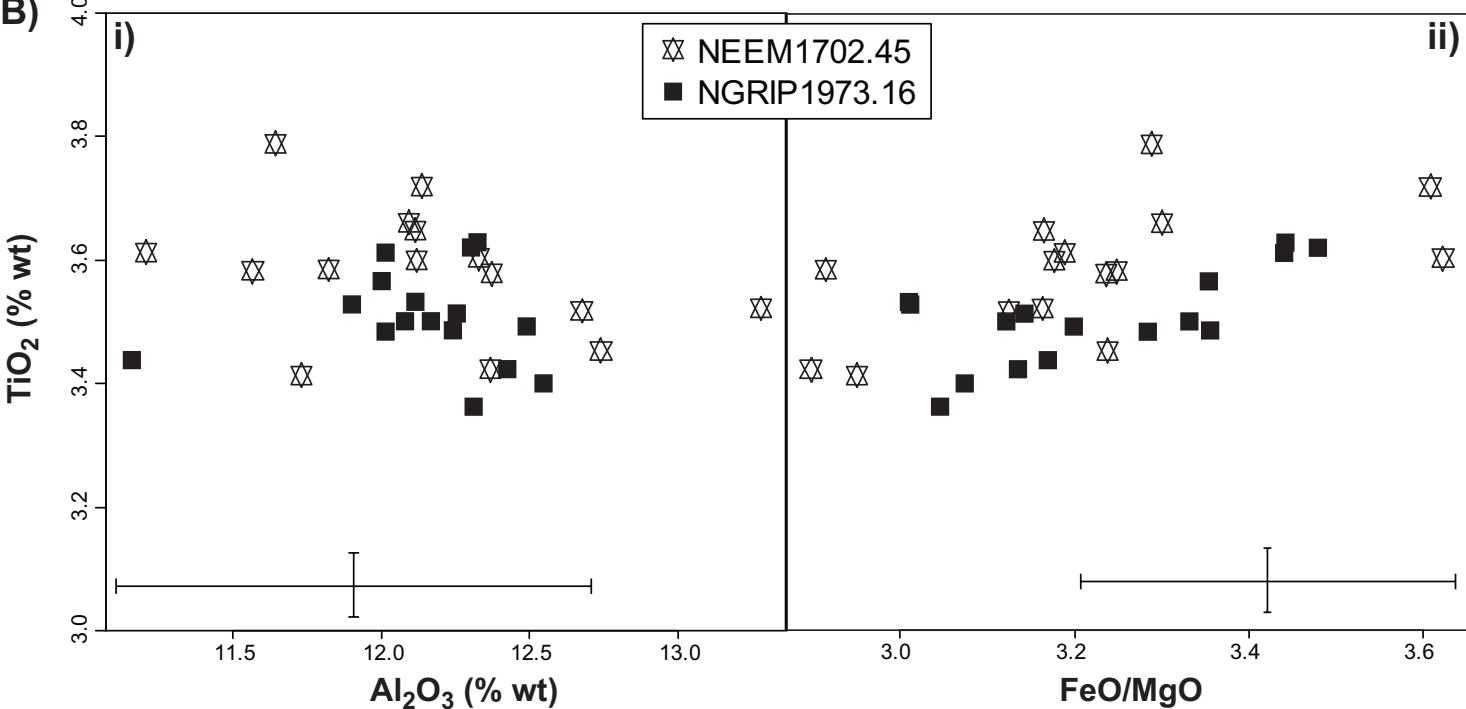
Figure 8



A) Figure 9



B)



C)

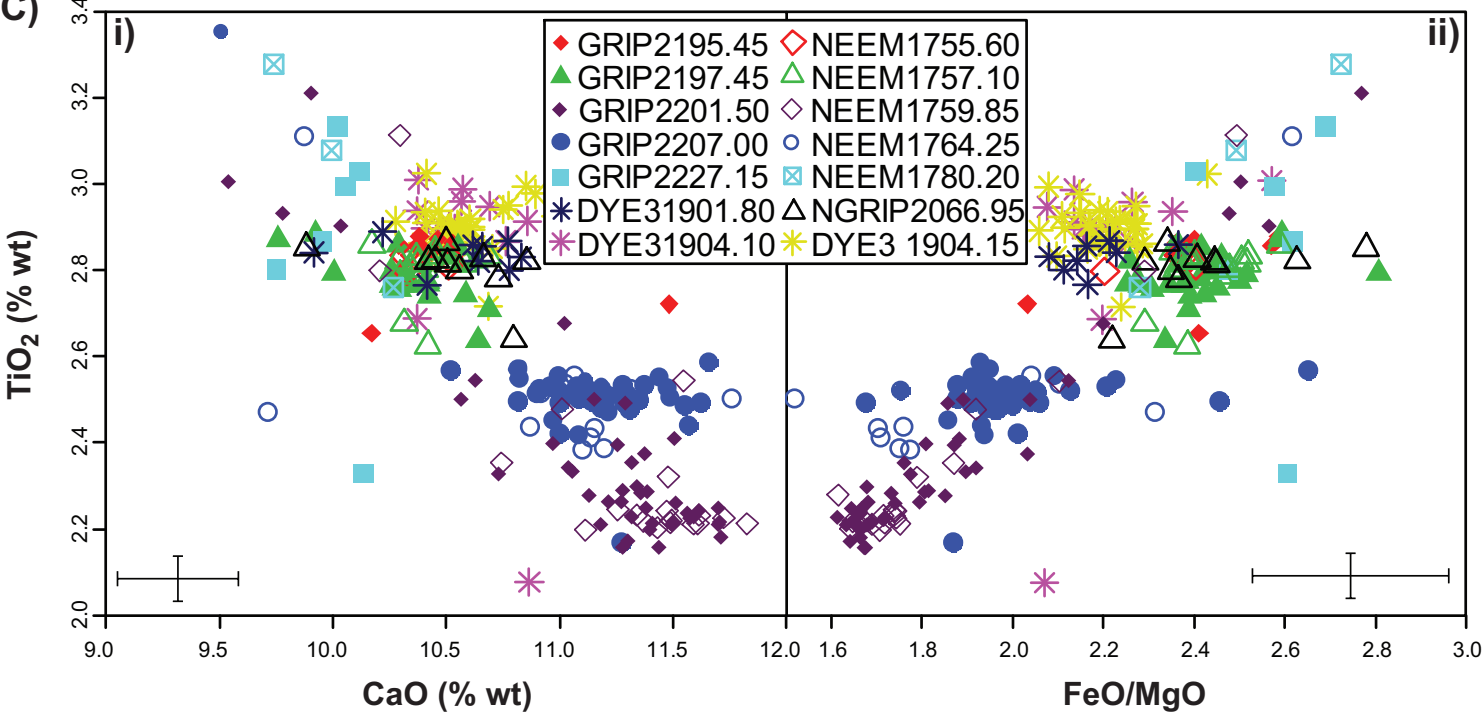


Figure10

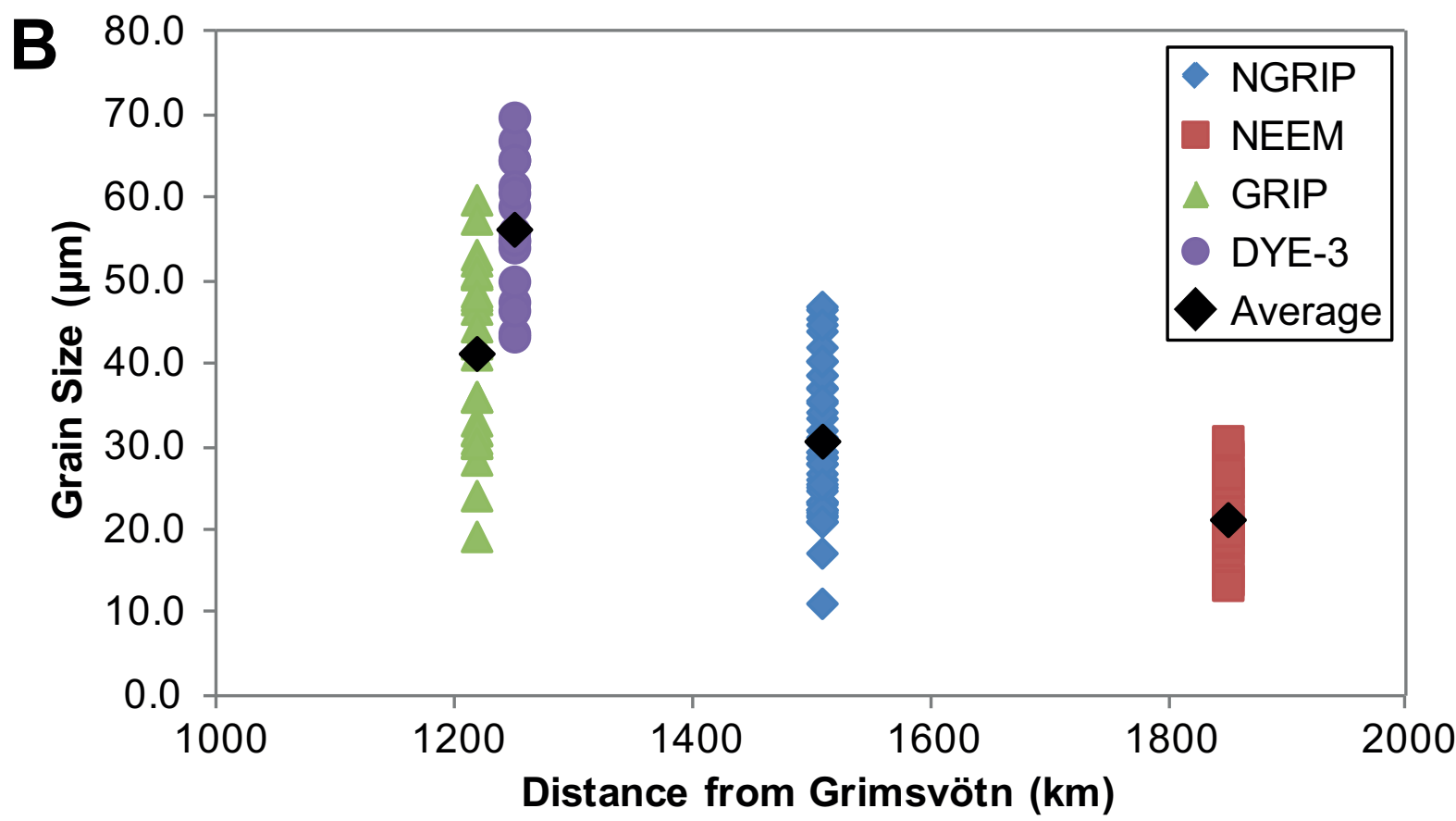
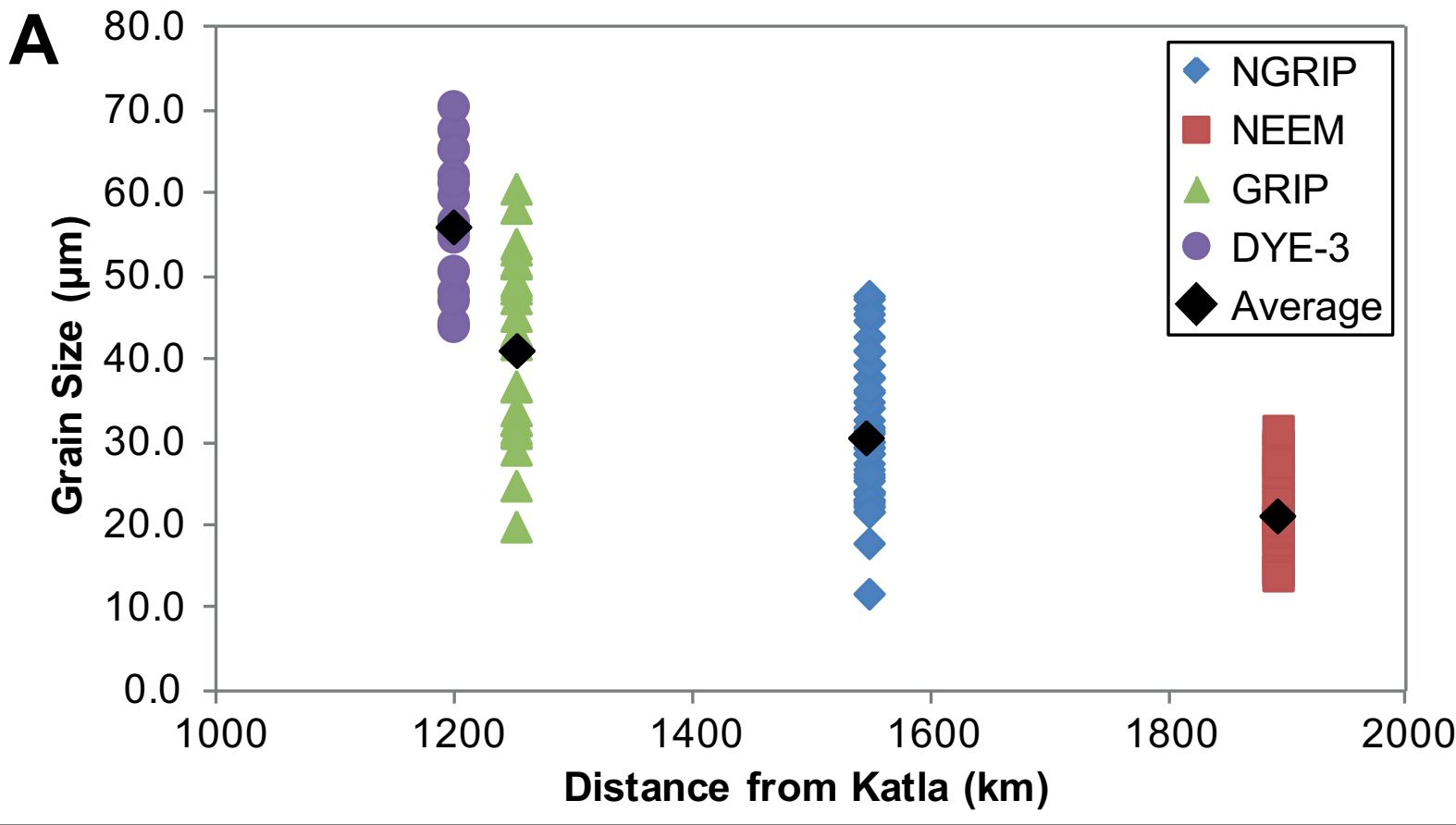
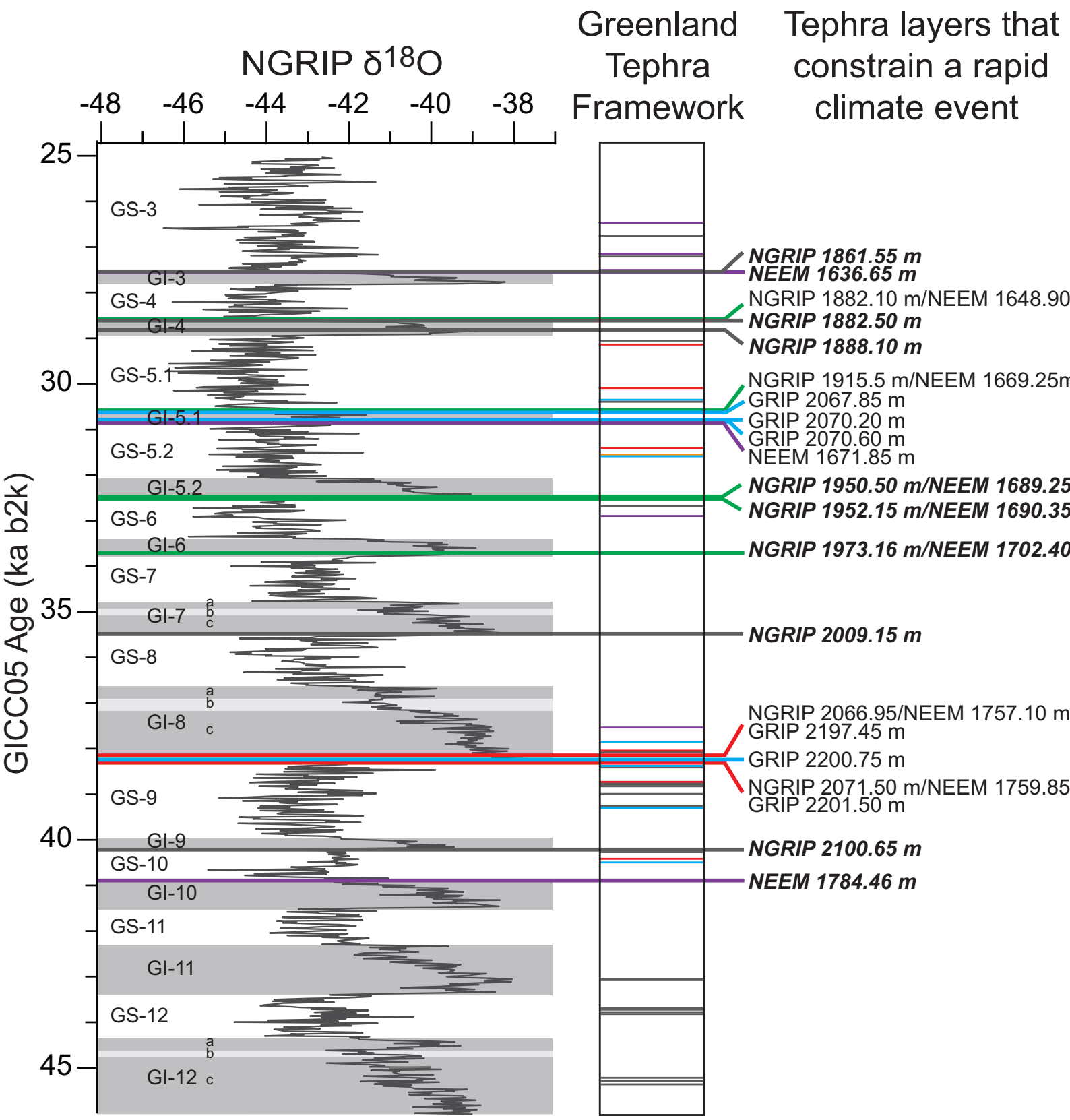


Figure11



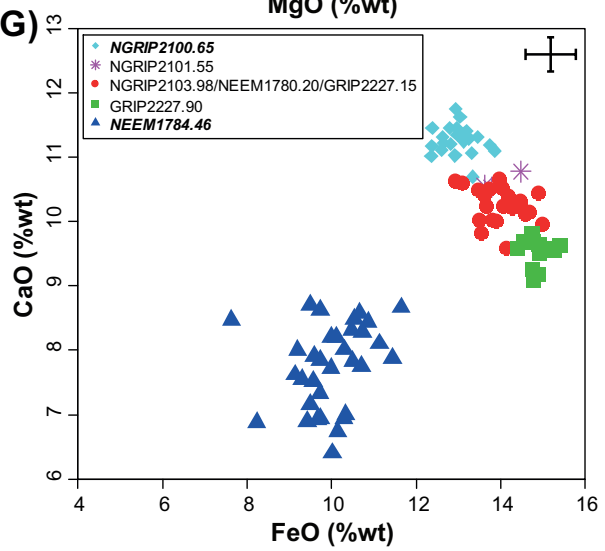
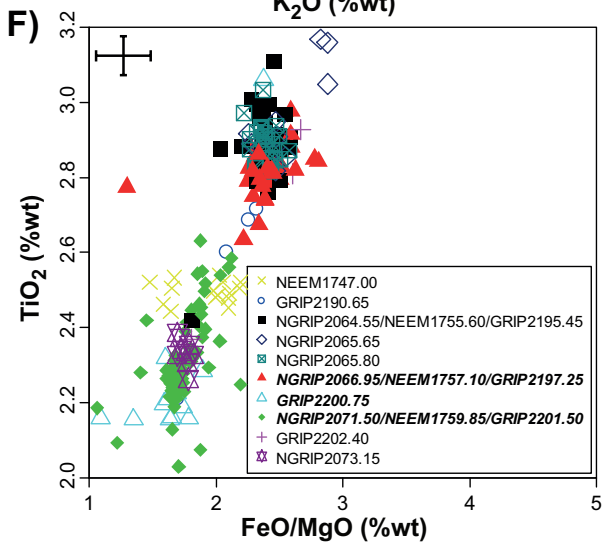
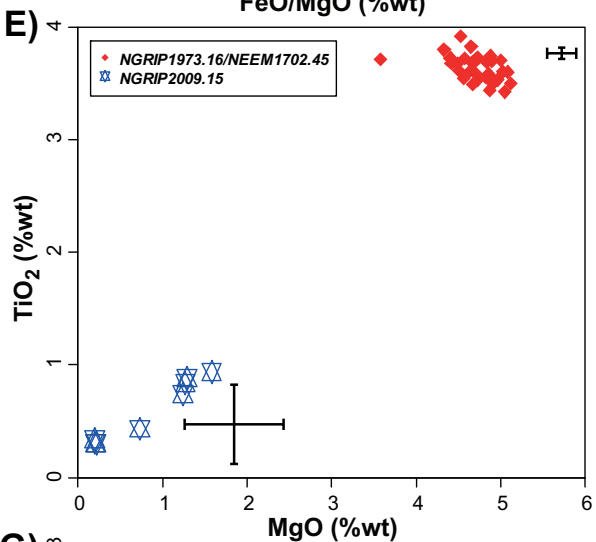
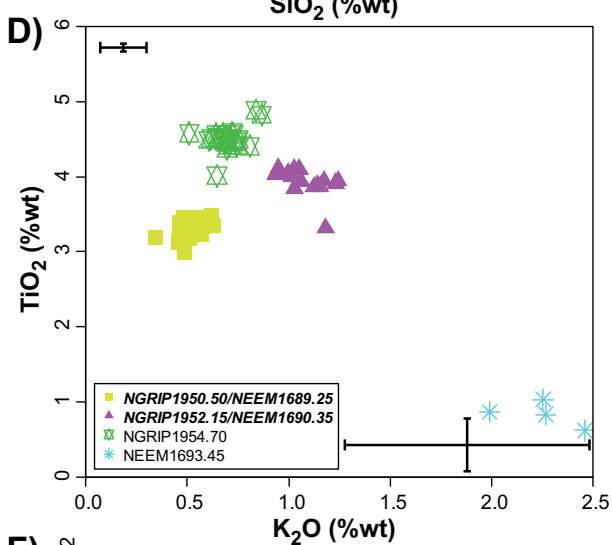
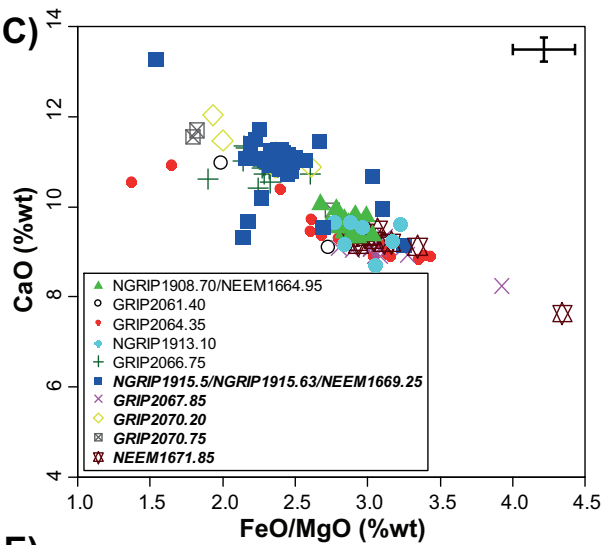
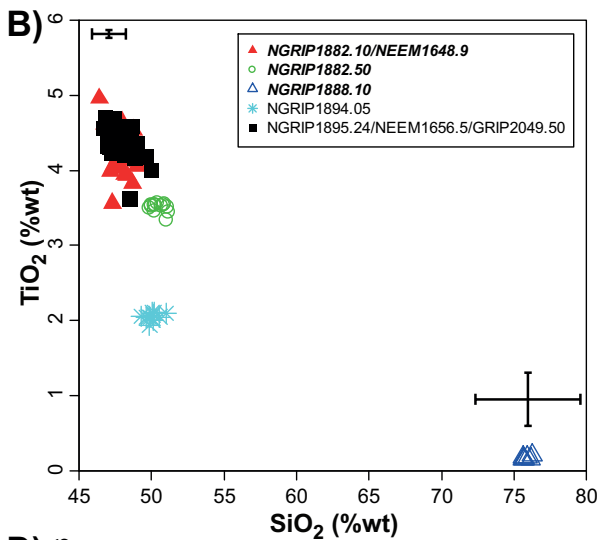
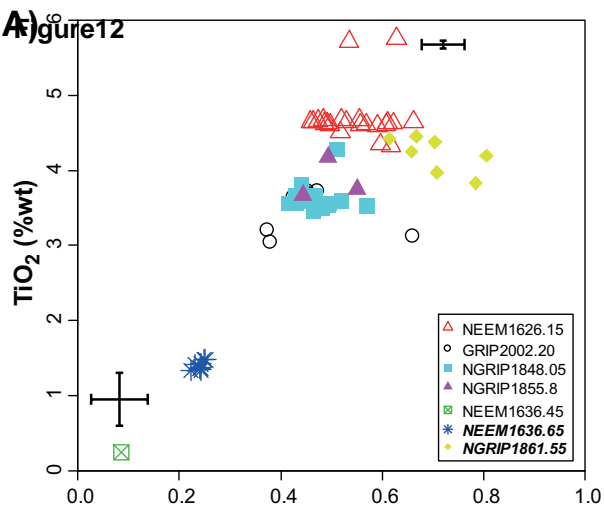
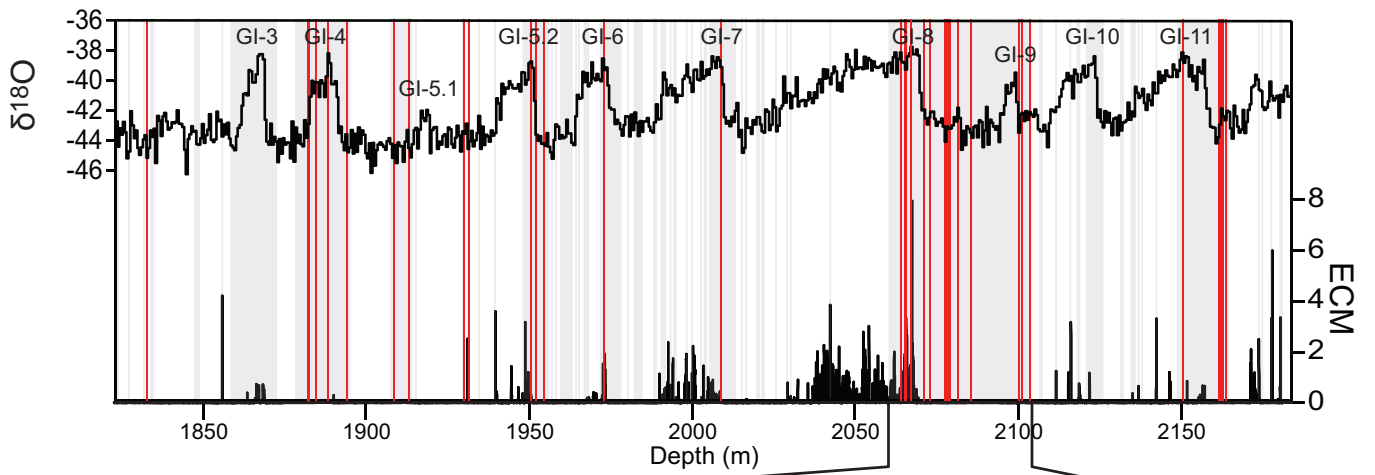
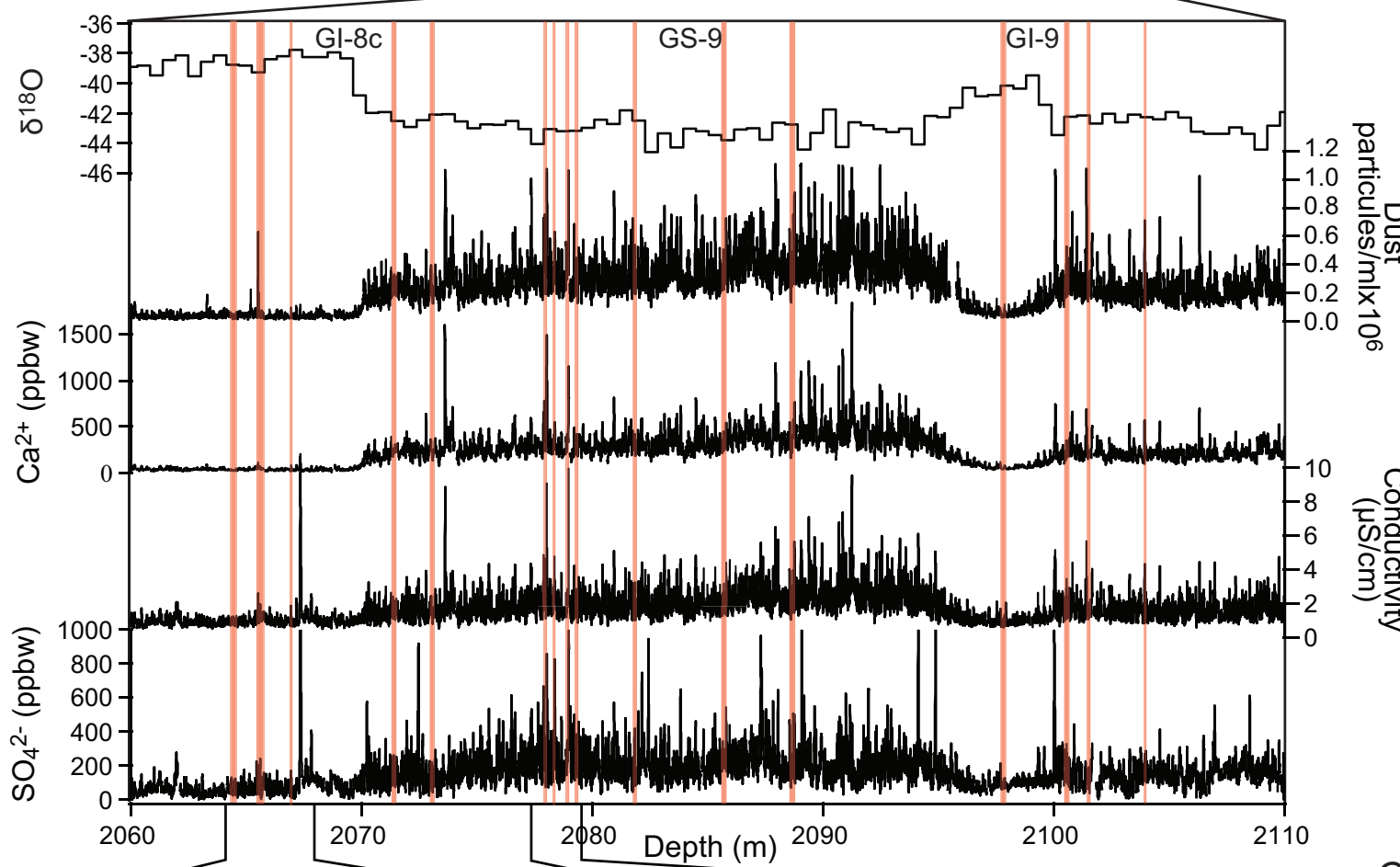


Figure 13

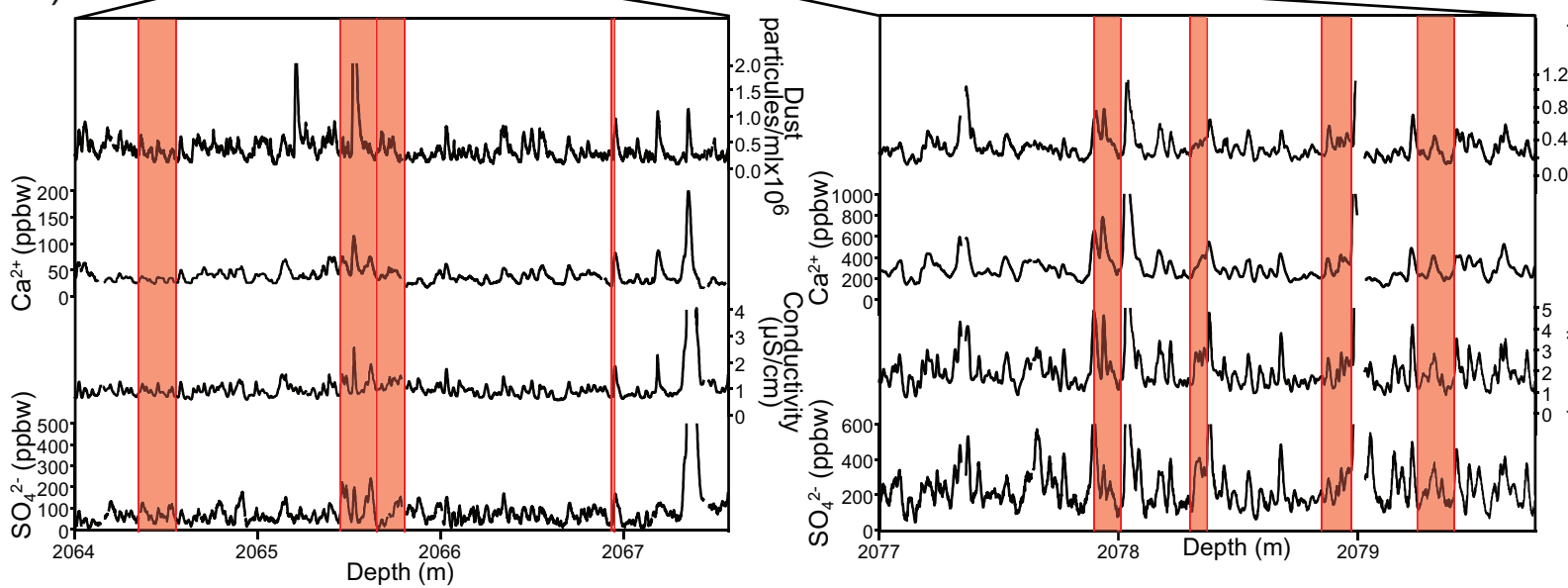
A)



B)



Ci)



Supplementary Data

[Click here to download Supplementary Data: Supplimentary Data.xls](#)