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Near conductive cooling rates in the upper-plutonic section of crust formed at the East Pacific

Kathrin Faak, Laurence A. Coogan, Sumit Chakraborty

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28 shallowest samples to $0.0003^{\circ}\text{C}\text{y}^{-1}$ for the deepest samples). Both the absolute
29 cooling rates, and the rate of decrease of cooling rate with depth, are consistent with
30 conductive thermal models. In contrast, the absolute cooling rates determined from
31 the deeper samples (>300 m below DGB), and the large decrease in cooling rate with
32 depth are inconsistent with thermal models that include substantial cooling by off-axis
33 hydrothermal circulation within the upper plutonic section of the crust.

34

35 **Keywords:** *cooling rate, oceanic crust, mid-ocean ridge, diffusion modeling, Mg-in-*
36 *plagioclase*

37

38 **1. Introduction**

39 The observation that only a small (~1 km wide and ~50 m deep) axial magma
40 lens (AML) overlies a zone of low seismic velocities (LVZ), that is interpreted to be a
41 crystal mush zone (e.g. Detrick et al., 1987; Dunn et al., 2000), at fast spreading
42 oceanic ridges has posed a conundrum: how is the large volume of the plutonic
43 section of the oceanic crust generated from such a small magma chamber? Two end-
44 member models – the gabbro-glacier model (Quick and Denlinger, 1993; Henstock et
45 al., 1993; Phipps Morgan and Chen, 1993; Fig. 1a) and the sheeted sill model
46 (Kelemen et al., 1997; Korenaga and Kelemen, 1997; Fig. 1b) – were proposed early
47 on to try to address this problem. Each model has its own set of geochemical (e.g.
48 fractionation trends), structural (e.g. pattern of distribution of fabric) and geophysical
49 (e.g. seismic or thermal) implications.

50 The thermal structure of mid-ocean ridges has long been known to provide
51 important insights into both the magmatic processes involved in crustal accretion and
52 the circulation of seawater-derived hydrothermal fluids through the crust (Morton and

53 Sleep, 1985; Lister, 1974). On-axis hydrothermal circulation through the sheeted dike
54 complex above the AML extracts heat from the roof of this magma body and drives
55 crystallization within it. What fraction of the lower crustal plutonic rocks crystallize
56 in the AML (losing their latent heat of crystallization into the overlying hydrothermal
57 system) is debated. The gabbro glacier model of crustal accretion (Fig. 1a) suggests
58 the vast majority of the latent heat is lost in this body with crystal subsidence to form
59 the lower crust (Quick and Denlinger, 1993; Henstock et al., 1993; Phipps Morgan
60 and Chen, 1993). The sheeted sill model (Fig. 1b), in contrast, suggests that most of
61 the plutonic section of oceanic crust formed at fast-spreading ridges crystallizes in
62 place (Kelemen et al., 1997; Korenaga and Kelemen, 1997); i.e. the latent heat of
63 crystallization of the plutonic section must be removed from throughout the lower
64 crust. For a given distribution of melt in the lower crust the sheeted sill model thus
65 requires less efficient heat extraction by on-axis hydrothermal circulation above the
66 AML and more efficient off-axis hydrothermal heat extraction from the sides of the
67 LVZ. As a consequence, the different end-member models predict different thermal
68 structures and different variations of cooling rate as a function of depth (Fig. 1). Note
69 that neither of these end-member models, or any hybrid between the two, excludes
70 hydrothermal circulation at any given depth or distance from the ridge axis – the
71 distinction between the models lies in the efficiency (amount and rate) of heat
72 removal by hydrothermal circulation at different locations. Our understanding of
73 crustal accretion mechanisms and the distribution and thermal consequences of
74 hydrothermal fluid flow would be substantially improved by better understanding the
75 thermal structure of the region surrounding the ridge axis.

76 The temperature structure in the lower oceanic crust at fast-spreading ridges
77 away from the ridge axis is currently poorly constrained. Seismic velocity anomalies

78 at 9°30'N on the East Pacific Rise (EPR), relative to a reference model, were
79 interpreted by Dunn et al (2000) in terms of temperature (and melt fraction)
80 anomalies. This approach led them to suggest that the isotherms in the lower crust are
81 steep, with temperatures <400°C throughout the upper 2 km of the lower crust, within
82 4 km of the ridge axis. As discussed by Dunn et al. (2000) and Webb (2008), there are
83 considerable uncertainties in the calculation of the thermal structure from seismic
84 velocity variations. For example, close to the ridge, the effect of anelasticity – a
85 relatively poorly known parameter (Dunn et al., 2000) – on seismic wave velocities is
86 prominent. Among other quantities, the activation energy of anelasticity plays a role
87 in the calculations and it is assumed to be the same as the activation energy of creep.
88 Dunn et al. (2000) used a value of 276 kJmol⁻¹ (Caristan, 1982) for their inversion of
89 seismic velocities to thermal structure, but subsequently this value has been re-
90 determined to be much higher (~ 485 kJmol⁻¹; Mackwell et al., 1998). As activation
91 energy controls the temperature dependence of properties, a thermal structure
92 calculated using this revised value is likely to be quite different. There are additional
93 sources of uncertainty such as in our knowledge of the power law exponent for the
94 frequency dependence of attenuation and what reference model the seismic structure
95 should be compared to. Notwithstanding such uncertainties, the thermal structure of
96 Dunn et al. (2000) has significantly influenced thinking about processes operating at
97 mid-ocean ridges. For example, this was the starting point for a recent model of
98 hydrothermal circulation developed by Hasenclever et al. (2014). However,
99 compliance data suggest that the crust is partially molten to >5 km off-axis (Crawford
100 and Webb, 2002) in the region studied by Dunn et al. (2000), inconsistent with the
101 thermal structure proposed by Dunn et al. (2000). Likewise, Han et al. (2014) show
102 that at 9°37-40 N on the EPR there are off-axis melt bodies that exist up to 10 km off-

103 axis. Such bodies would rapidly freeze if the off-axis crust was cooled and hence their
104 presence suggests a relatively warm off-axis thermal structure (Han et al., 2014;
105 Coogan, 2014).

106 The discussion above indicates that there is a need to determine the thermal
107 structure directly, for example by determining cooling rates at different locations. One
108 quantitative approach for determining the thermal structure of the lower crust comes
109 from the petrological tool of *geospeedometry* (Lasaga, 1983). This allows the cooling
110 rate of rocks to be determined from the compositional zoning of minerals. Given a
111 spreading rate, a subsolidus cooling rate can be directly inverted into isotherm
112 separation or compared to cooling rates predicted by thermal models. However, the
113 only attempts to-date to apply such tools to the lower oceanic crust have used the *Ca-*
114 *in-olivine geospeedometer* (Coogan et al., 2002; Coogan et al., 2007, VanTongeren et
115 al., 2008). Due to paucity of olivine in many evolved gabbros, and the susceptibility
116 of olivine to alteration, this approach has not been widely applied to *in situ* crust
117 formed at fast-spreading ridges. Here, we apply a new approach specifically
118 developed to be suited to such rocks.

119 Clinopyroxene and plagioclase are virtually omnipresent in the lower oceanic
120 crust and thus a geospeedometric method based on exchange of Mg between these
121 phases was developed (Faak et al., 2014). Natural rock samples from the oceanic crust
122 show much higher concentrations of MgO in plagioclase phenocrysts in mid ocean
123 ridge basalts (MORBs) than in the cogenetic, but more slowly cooled, gabbroic rocks
124 of the lower oceanic crust (Fig. 13f in Coogan, 2014). This difference in plagioclase
125 Mg-content is explained by diffusion out of the plagioclase into the clinopyroxene
126 during cooling. The partition coefficient of Mg between plagioclase and
127 clinopyroxene, $K_{Mg}^{P/Cp}$, decreases with temperature (Faak et al., 2013), i.e., during

128 cooling an exchange of Mg between plagioclase and clinopyroxene rims is required
129 for these to remain in equilibrium. The exchange process leads to the development of
130 a Mg concentration gradient within the plagioclase as Mg diffuses out of plagioclase
131 and into clinopyroxene. Diffusion is a thermally activated process and becomes
132 slower with decreasing temperatures. Hence, there will be a temperature, T_c , at which
133 diffusion becomes too slow for measurable Mg loss from the plagioclase. This closure
134 temperature (T_c) depends on the distance from the interface (Dodson, 1986; Onorato
135 et al., 1981), such that the rims of a plagioclase crystal will be able to maintain
136 equilibrium Mg-concentrations down to lower temperatures than the core of the
137 crystal, leading to the development of a closure profile that is convex upwards for a
138 continuous cooling history. Faak et al. (2014) show how the evolution of the resulting
139 concentration profile of Mg in plagioclase depends on the cooling history. For
140 example, a slow cooling rate will allow extensive diffusive exchange of Mg from
141 plagioclase into clinopyroxene leading to low Mg contents of plagioclase. In contrast,
142 fast cooling rates will lead to high plagioclase Mg contents if all other relevant
143 parameters (grain size, grain shape, anorthite content) remain unchanged (for a
144 detailed discussion on the evolution of diffusion profiles for different cooling histories
145 and additional factors influencing the resulting shape of the profile see Faak et al.,
146 2014). Here we apply the *Mg-in-plagioclase geospeedometer* (Faak et al., 2014) to
147 three suites of oceanic gabbros that formed at the East Pacific Rise to determine their
148 cooling history and infer the thermal state of the ridge axis region.

149

150 **2. The Sample Suites**

151

152 There are only three locations in the world where well-located samples of the
153 upper plutonic section of oceanic crust formed at modern fast-spreading ridges have
154 been collected. These are the *Hess Deep Rift* in the equatorial Pacific, the *Pito Deep*
155 in the southern Pacific, and *IODP (International Ocean Discovery Program) Site*
156 *I256D* in the eastern Pacific (Fig. 2a). For this study we investigated samples from all
157 three locations and from different depths within the lower oceanic crust.

158 The largest and best constrained sample suite is from the *Hess Deep Rift*,
159 where ~1 Ma old crust that formed at the equatorial East Pacific Rise (full spreading
160 rate ~135 mmy⁻¹), is rifted apart due to the westward propagation of the Cocos-Nazca
161 spreading center (Lonsdale, 1988; Francheteau et al., 1990). The rifting has created a
162 tectonic window (Fig. 2b) that exposes the entire upper crust (lavas and dikes,
163 ~1200 m) as well as the upper part (~1000 m) of the gabbros (Karson et al., 2002).
164 This study mainly focuses on samples collected by submersible from the North wall
165 of the Hess Deep Rift (Lonsdale, 1988; Karson et al., 2002, Fig. 2d and e). Because
166 the dike/gabbro boundary (DGB) was mapped on multiple dives (Figure 2d), and the
167 depth below sea level is known for each sample, it is possible to reconstruct the depth
168 below the DGB for each sample (Table 1).

169 The second sample suite comes from *Pito Deep*, where ~3 Ma old crust
170 formed at the EPR (full spreading rate ~140 mmy⁻¹) is rifted apart due to a
171 propagating rift tip at the northeastern corner of the Easter Microplate (Francheteau et
172 al., 1988; Hey, 1995, Fig. 2c). Continuous sections of the oceanic crust including
173 lavas, sheeted dikes and the upper gabbroic rocks are exposed (Constantin et al.,
174 1995; Hekinian et al., 1996, Constantin et al., 1996; Perk et al., 2007). Gabbroic rocks
175 investigated in this study were collected during the *Jason II* and *Alvin* dive programs
176 during cruise AT11-33 of the R/V *Atlantis* (e.g., Perk et al., 2007; Figure 2f and g),

177 The DGB was mapped during the dives (Figure 2f), which allows us to reconstruct the
178 depth below the DGB for each sample (Table 1).

179 The final place that a short section of plutonic rocks formed at a modern fast-
180 spreading ridge have been sampled at is *IODP Site 1256D*, which drilled into ~15 Ma
181 old intact oceanic crust of the Cocos Plate that formed at the superfast spreading EPR
182 (full spreading rate ~220 mmy⁻¹). In this drilling project ~1250 m of oceanic crust has
183 been sampled to-date, providing a section from extrusive lavas, through sheeted dikes
184 and into the top of the plutonic section (Wilson et al., 2006). Hole 1256D penetrated
185 the top of the plutonic section where gabbros and granoblastic dikes are intermixed
186 (Wilson et al., 2006; Koepke et al., 2008; France et al., 2009; Sano et al., 2011).

187

188 **3. Mg-concentration profiles in plagioclase**

189

190 In order to apply the *Mg-in-plagioclase geospeedometer* to the most suitable
191 rocks from these sample suites, gabbroic rocks with coexisting plagioclase and
192 clinopyroxene that appear fresh (i.e. nearly unaltered) were chosen. In addition,
193 plagioclase crystals with nearly idiomorphic grain shapes and high aspect ratios were
194 selected, and a full profile (rim-core-rim) was measured along the short dimension of
195 the crystal to minimize effects from diffusion in three dimensions (Faak et al., 2014).
196 The composition of the clinopyroxene adjacent to each profile was analyzed when
197 possible. If no clinopyroxene was directly adjacent to a profile, then a clinopyroxene
198 in close proximity to the respective plagioclase was analyzed. Some Mg-profiles were
199 measured along a profile, where plagioclase was in direct contact with clinopyroxene
200 on one side, but with another plagioclase crystal on the other side (but with
201 clinopyroxene in close proximity) to test if grain boundaries acted as fast pathways

202 for diffusion. If they did then plagioclase is expected to exchange Mg with a
203 neighboring clinopyroxene, even when the grains are not in direct contact, and the
204 measured Mg-profile should show approximately the same Mg-concentration on both
205 rims (for similar X_{An} on both rims; see Dohmen and Chakraborty, 2003 for a
206 quantitative explanation). As discussed below this allows us to evaluate whether
207 transport along the grain boundary is infinitely fast and efficient or if transport along
208 grain boundaries limits Mg exchange between plagioclase and clinopyroxene grains if
209 they are separated by too long a distance.

210

211 ***3.1 Measurement of Mg-concentration profiles in plagioclase***

212 The concentration of Mg and major elements in plagioclase was measured
213 using a *Cameca SX-50* electron microprobe (EMP) fitted with four wavelength-
214 dispersive spectrometers at the Ruhr-Universitaet in Bochum. Natural and synthetic
215 mineral standards were used for calibration and an on-line $\phi(\rho z)/PAP$ correction
216 procedure was used to correct for absorption, fluorescence and atomic number. To
217 attain high precision analysis of Mg in plagioclase, the measurement conditions
218 outlined by Faak et al. (2014) were applied (also see Supplemental Table S1 for
219 details). Briefly, these are: beam current = 40 nA, accelerating voltage = 15 kV, and
220 long counting times for Mg (90 s on the peak and 45 s on each background, with the
221 background positions selected specifically for the measurement of Mg in plagioclase),
222 which allows a 3σ detection limit of ~ 75 ppm to be achieved. Using these conditions
223 the precision on Mg in plagioclase at a concentration of 0.05 wt% MgO is ~ 0.0025
224 wt% on this instrument. The distance between analyzed spots along the profile was
225 $5 \mu\text{m}$ for shorter profiles and $10 \mu\text{m}$ for longer profiles. The first and last

226 measurements at the rims of the plagioclase were approximately 2 to 5 μm away from
227 the interface.

228

229 ***3.2 Mg-concentrations and profile shapes as function of depth in the***
230 ***plutonics***

231 The measured concentrations of MgO in plagioclase from all sample suites
232 vary between 0.01 wt% (at some of the rims) and 0.13 wt% (at the cores of the
233 shallowest samples). In the same plagioclase grain, Mg-concentrations at the rim are
234 similar for plagioclase-clinopyroxene contacts and plagioclase-plagioclase contacts,
235 implying that the grain boundaries indeed act as fast pathways for the Mg-exchange
236 (see above).

237 Samples from the Hess Deep and Pito Deep sample suites studied here, cover
238 a depth range from 0 to 520 m and 42 to 836 m below the DGB, respectively, and
239 allow variation of the plagioclase Mg-concentration profiles as a function of depth to
240 be observed. Both sample suites show a systematic decrease of MgO in the cores of
241 plagioclase crystals with increasing sample depth (Fig. 3). The plagioclase rims show
242 MgO-concentrations between 0.01 to 0.05 wt%, with no systematic variation with the
243 sampling depth. This leads to Mg-profiles with stronger curvature in the shallower
244 samples (i.e., larger difference in Mg content between the cores and rims, e.g., Fig. 3a
245 and e) and rather homogeneous Mg-profiles for the deeper samples (i.e., equally low
246 Mg-concentrations at the core and the rims, e.g., Fig. 3d and h).

247 Many samples from the IODP Site 1256D show MgO-concentrations below
248 the detection limit of the EMP, and the plagioclase appears to be hydrothermally
249 altered – we interpret this as indicating that Mg-loss occurred during dissolution-
250 reprecipitation reactions rather than via solid-state diffusion. Only two shallow

251 samples (12.1 and 12.4 m below the DGB) from this location were suitable for
252 modeling using the approach of this study (Table 1).

253

254 **4. Modeling diffusion profiles of Mg in plagioclase and fitting the natural data**

255

256 ***4.1 Diffusion model and parameters***

257 Cooling rates are obtained from diffusion modeling of Mg in plagioclase, using
258 the *Mg-in-plagioclase geospeedometer* of Faak et al. (2014). This method is based on
259 the diffusive exchange of Mg between plagioclase and clinopyroxene under the
260 assumptions of: (i) instantaneous equilibrium at the interface between the two phases,
261 and (ii) clinopyroxene acting as an infinite reservoir. As described above (Section 1),
262 during cooling, the Mg-concentration at the interface changes, and the resulting
263 concentration gradient provides a driving force for diffusion out of the plagioclase
264 into the clinopyroxene. As diffusion of Mg in plagioclase is coupled with the
265 anorthite content, X_{An} , in plagioclase, this has to be accounted for in the diffusion
266 equation (Costa et al., 2003; Appendix A). Additionally, the diffusion coefficient of
267 Mg in plagioclase, D_{Mg}^{Pl} , and the partition coefficient $K_{Mg}^{Pl/Cpx}$ depend on the silica
268 activity, a_{SiO_2} , of the system (Faak et al., 2013; Appendix A), which therefore needs
269 to be constrained.

270 Faak et al. (2014) show how the general *Mg-in-plagioclase geospeedometer* may
271 be specifically applied to rocks from the lower oceanic crust, and here we follow the
272 approach outlined in their example. In detail, (i) the diffusion coefficient D_{Mg}^{Pl} and the
273 partition coefficient $K_{Mg}^{Pl/Cpx}$ are taken from Faak et al. (2013), as those were
274 specifically determined in the compositional range of plagioclase and clinopyroxene

275 found in the lower oceanic crust; (ii) an initial profile for Mg in plagioclase is
276 calculated from $K_{Mg}^{Pl/Cpx}$ at temperatures around 1200°C, where the exact starting
277 temperature, T_{start} , depends on the grain size of the plagioclase; (iii) the silica activity,
278 a_{SiO_2} , is assumed to be constrained by the assemblage olivine + orthopyroxene
279 (Appendix A), since this assemblage is found in many of the samples investigated
280 here (see Faak et al., 2014 for details and discussion). The change in plagioclase Mg-
281 concentration due to diffusion is calculated iteratively along a given cooling path
282 using a finite difference approach (Faak et al., 2014; Appendix A). As a first
283 approximation, a linear cooling path over the modeled temperature interval is
284 assumed, and the cooling rate dT/dt is iteratively refined until the best visual match
285 between the measured and the modeled Mg-profile is found.

286

287 ***4.2 The temperature interval***

288 Under the assumption of instantaneous equilibrium at the interface between
289 plagioclase and clinopyroxene, diffusion along a linear cooling path produces Mg-
290 closure-profiles that are convex upwards (Faak et al., 2014), i.e., have lower Mg-
291 concentrations at the rims than at the core. The measured Mg-concentration profiles
292 from the oceanic crust commonly show a significantly smaller degree of convex
293 curvature, than would be expected from the model of Faak et al. (2014) for a cooling
294 history in which temperature changes as a linear function of time (i.e., a constant
295 cooling rate). The lack of a strong curvature of the Mg-profiles can be explained in
296 two ways, either: (A) the approximation of a linear cooling history is only valid up to
297 a certain temperature, and the cooling rate increases below this temperature. A
298 cooling history with increasing cooling rates towards lower temperatures produces
299 more homogeneous Mg-profiles (Faak et al, 2014); or, (B) the assumption of

300 instantaneous equilibration at the crystal interface is only valid above a certain
301 temperature (here referred to as T_{closed}). This means, Mg is not effectively exchanged
302 between plagioclase and clinopyroxene below T_{closed} ; however, diffusion of Mg in
303 plagioclase continues in a closed system and tends to homogenize the Mg-
304 concentration (for a homogeneous X_{An} -content), leading to less curvature in the
305 measured Mg-profile (for details see Appendix B). Faak et al. (2014) briefly discuss
306 this issue and model two examples from the oceanic crust. They defined a
307 temperature, T_{rim} , that is calculated from the measured Mg-concentration at the rim
308 (defined as 2-5 μm from the interface) of a plagioclase crystal with a clinopyroxene
309 (Faak et al., 2014). This temperature is interpreted as the closure temperature below
310 which diffusion ceased to change the Mg-concentration significantly in a region 5 μm
311 away from the interface. Hence, Faak et al. (2014) end the modeling procedure for a
312 linear cooling rate at this temperature T_{rim} , implying that diffusion effectively stops
313 below this temperature (i.e., they model scenario (A)). They show that it is possible to
314 fit strongly curved and homogeneous Mg-profiles using this approach.

315 For this study, we also investigated the evolution of diffusion profiles of Mg in
316 plagioclase for the case of exchange of Mg between plagioclase and clinopyroxene
317 ceasing below a temperature T_{closed} , and the continuation of Mg diffusion in
318 plagioclase in a closed system (i.e., scenario (B), see Appendix B). In this model the
319 bulk Mg content of the plagioclase does not change below T_{closed} and the diffusive
320 fluxes within the plagioclase below this temperature have to be sufficient to
321 homogenize the Mg-distribution. These combined constraints tightly constrain the
322 maximum cooling rates. Both modeling approaches were applied to three hypothetical
323 plagioclase crystals (P1: weakly curved Mg-profile with 0.04 wt% MgO at the rims
324 and 0.06 wt% MgO at the core; P2: homogeneous Mg-profile at 0.04 wt% MgO; P3:

325 homogeneous Mg-profile at 0.02 wt% MgO; all with homogeneous $X_{An} = 0.6$). The
326 direct comparison of the two scenarios for identical plagioclase crystals shows that
327 T_{closed} is always slightly lower than T_{Crim} (25°C for P1, and 20°C for P2 and P3;
328 Appendix B). The cooling rates dT/dt determined for the temperature interval above
329 T_{closed} are slightly faster, when scenario B is modeled (e.g., P1: $dT/dt = 0.008^{\circ}\text{Cy}^{-1}$ for
330 scenario (A) and $0.01^{\circ}\text{Cy}^{-1}$ for scenario (B); P2: $dT/dt = 0.0003^{\circ}\text{Cy}^{-1}$ for scenario (A)
331 and $0.001^{\circ}\text{Cy}^{-1}$ for scenario (B); Appendix B). Thus, for homogeneous Mg-profiles,
332 the levels of variation in dT/dt and T_{Crim} / T_{closed} arising from the choice of the
333 modeling approach (A vs. B) are in the range of other uncertainties of the *Mg-in-*
334 *plagioclase geospeedometer* (e.g., uncertainties in cooling rate arising from the choice
335 of D_{Mg}^{Pl} can lead to a factor of 3 difference in cooling rate; Faak et al., 2014). For Mg-
336 profiles with some degree of curvature, the differences are even smaller. The cooling
337 rates presented below were determined using method A.

338

339 **5. Cooling rates as a function of depth in the plutonic section of EPR crust**

340

341 Cooling rates were obtained from 41 individual plagioclase crystals (51
342 profiles) in 23 samples from the three sample suites (32 profiles in 13 samples from
343 Hess Deep, 16 profiles in 8 samples from Pito Deep, 3 profiles in 2 samples from
344 IODP Site 1256D; Table 1; Supplemental Material), spanning a total depth interval
345 below the DGB of 0 to 836 m. Cooling rates systematically decrease with increasing
346 depth, and cover almost 4 orders of magnitude, ranging from $\sim 1^{\circ}\text{Cy}^{-1}$ for the
347 shallowest samples to $0.0003^{\circ}\text{Cy}^{-1}$ for the deepest samples (Table 1, Fig. 4). Cooling
348 rates from all three sample suites agree very well with each other (with respect to their
349 individual depth below the DGB) and fall on the same trend (Fig. 4).

350 The uncertainty for each data point (i.e. the uncertainty of the cooling rate
351 obtained from each Mg-profile) cannot be expressed by a simple analytical expression
352 for propagation of errors. This is because, in addition to uncertainties arising from the
353 error associated with each individual parameter in the diffusion equation, the overall
354 uncertainty in the determination of cooling rates also depends on factors such as the
355 X_{An} -gradient in the plagioclase crystal. However, Faak et al. (2014) evaluated the
356 overall uncertainty in the procedure using Monte-Carlo simulations and simulated
357 profiles with unzoned X_{An} and known thermal histories. They also evaluated the
358 uncertainty that results from the use of different sets of diffusion coefficients (i.e. Van
359 Orman et al., 2014 and Faak et al., 2013). They found that cooling rates can be
360 determined to better than half an order of magnitude. Additionally, the difference in
361 the obtained cooling rates from multiple Mg-profiles in multiple plagioclase crystals
362 in the same sample (i.e. the scatter in the data for one sample) may be taken as a
363 measure for the precision of the cooling rate estimate for a single sample. Following
364 this approach, the precision on the obtained cooling rate is also found to be better than
365 half an order of magnitude. therefore, the uncertainties are much smaller than the
366 observed decrease in cooling rate of almost four orders of magnitude.

367 The cooling rates obtained here are similar to those derived from a small
368 subset of the samples using *Ca-in-olivine geospeedometry* (Coogan et al., 2007) and
369 from a larger sample suite from the Oman ophiolite (Fig. 4). The fact that two
370 different methods (*Mg-in-plagioclase* and *Ca-in-olivine*), which are based on different
371 diffusion and partition coefficients, yield similar values for the cooling rate at a given
372 depth below the DGB implies that not only is the relative variation in cooling rate
373 with depth well constrained, but also that the absolute values of cooling rates are
374 robust (i.e., the cooling rates are accurate as well as precise). Importantly, the very

375 similar cooling rates extracted from four different crustal sections indicates that
376 cooling rates in the uppermost plutonics formed at fast-spreading ridges are similar –
377 i.e., the thermal structure must be similar and close to steady-state. We expand on
378 these points below.

379 One sample from the Pito Deep suite yields cooling rates that are shifted from
380 the cooling rate versus depth trend defined by all the other samples to slower cooling
381 rates (sample 022005-1052, Table 1 and Fig. 4). The slow cooling rates obtained from
382 this sample are due to the low Mg-content (~0.015-0.03 wt% MgO) of plagioclase in
383 this sample. We do not have any explanation for this observation at this point of time
384 and treat this as an outlier from the general trend.

385

386 **6. Discussion - implications for the mode of cooling of the oceanic crust and** 387 **comparison to thermal models**

388

389 The observed variation in cooling rate with depth is very consistent for three
390 different sections of crust formed at the EPR (and the Oman ophiolite; Fig. 4),
391 implying a similar thermal structure in the near-axis region along the EPR. This
392 indicates that the thermal structure is near steady-state in the temperature window
393 recorded (~1100-700°C; Table 1). The significant decrease of cooling rate with
394 increasing depth in the uppermost part of the oceanic lower crust (~4 orders of
395 magnitude over 840 m below DGB) provides a fundamental constraint on models of
396 crustal accretion and hydrothermal circulation at fast-spreading ridges.

397 The cooling rates determined here are compared to those from modeling
398 studies, and the inversion of seismic velocity into temperature, in Fig. 5. Both the
399 absolute cooling rates determined from the deeper samples (>300 m below DGB) and

400 the large decrease in cooling rate with depth are inconsistent with thermal models that
401 include substantial cooling by off-axis hydrothermal circulation. For example,
402 Maclennan et al. (2005) report the results of three thermal models that have 50-71
403 kWm^{-1} of hydrothermal cooling off-axis. These models calculate little variation in
404 cooling rate with depth, faster cooling rates than those observed (except in the upper
405 ~100 m; blue lines in Fig. 5), and are inconsistent with the depth distribution of
406 cooling rates determined by geospeedometry. The same is true of the more
407 sophisticated thermal model of Hasenclever et al. (2014) and the cooling rates
408 predicted by the isotherm separation of Dunn et al. (2000) that this model is based on
409 (blue crosses in Fig. 5).

410 While the cooling rates determined here are inconsistent with models with
411 extensive off-axis hydrothermal heat extraction at depth (Maclennan et al., 2005;
412 Hasenclever et al., 2014; blue area in Fig. 5), they are broadly consistent with
413 conductive cooling models (green lines in Fig. 5). This does not mean that there is no
414 hydrothermal circulation in this region, or that it extracts no heat from the lower crust,
415 but just that the thermal effects of such circulation are minor. Near conductive cooling
416 rates in the upper part of the lower oceanic crust are consistent with the recent finding
417 of off-axis sills in the lower crust along the East Pacific Rise (Canales et al., 2009;
418 Han et al., 2014). The recent discovery of on-axis, sub-axial melt lenses (Marjanovic
419 et al., 2014) provides no new constraints on the thermal structure of the axial region
420 because it has long been thought that this region contains partial melt due to the low
421 seismic velocities (e.g. Harding et al., 1989). If the crust was efficiently cooled in the
422 off-axis by hydrothermal circulation, producing the kind of thermal structure
423 suggested by Dunn et al. (2000) and Hasenclever et al. (2014), then any magma
424 intruded off-axis would be emplaced into cold wall-rocks and hence would freeze

425 rapidly. For example, using Eq. (3) in Chapter 2.2 in Carslaw and Jaeger (1959), a
426 200 m high sill that intruded at 1200°C into country rock with a temperature of 600°C
427 would solidify within <1000 y even if conduction was the only heat transport
428 mechanism; since extensive hydrothermal circulation is required for the off-axis
429 region to be cool, this calculation over estimates the solidification time. Hence the
430 probability of observing such bodies in seismic studies would be very small (Coogan,
431 2014; Han et al., 2014). In contrast, the near conductive cooling rates determined here
432 imply warm off-axis upper plutonic crust, and hence longer off-axis sill lifetimes, and
433 a higher probability of identifying these bodies seismically.

434 The observed plagioclase Mg contents, the shapes of their Mg zoning profiles,
435 and their variation with depth provide information about: (a) the temperatures around
436 which exchange of Mg with surrounding clinopyroxene froze ($\sim T_{c_{rim}}$ in our
437 notation), and (b) the cooling rates around these temperatures as a function of depth.
438 These allow us to constrain the earliest point of time (equivalent to distance from the
439 axis in a ridge-spreading situation with a known spreading-rate) at which the
440 temperature $T_{c_{rim}}$ may have been reached at a given depth (Fig. 6). This is obtained by
441 assuming the physically unrealistic, but mathematically limiting, case where cooling
442 above ($T_{c_{rim}} + 100$ °C) was instantaneous. Any other more physically realistic cooling
443 scenario would lead to the observed $T_{c_{rim}}$ at that depth being attained at a later point
444 of time and hence greater distance from the ridge (marked as gray field in Fig. 6).
445 This information extracted from the compositional characteristics (concentration,
446 zoning pattern) of plagioclase crystals may be compared with those obtained from
447 studies where the thermal structure was obtained by thermal modeling or inverting
448 other kinds of data such as seismic velocity profiles (e.g., Hasenclever et al., 2014;
449 Dunn et al., 2000). As seen in Fig. 6, these predict much lower temperatures (~ 400

450 °C) at depths greater than 500 m below the dike / gabbro boundary at much earlier
451 times, and hence much closer to the ridge axis (due to the much faster cooling rates in
452 Fig. 5). These trends fall well outside the gray region in Fig. 6 that is permissible
453 based on the observations of the compositional characteristics of plagioclase. In other
454 words, the thermal structure of the upper part of the lower crust that we obtain from
455 our observations is inconsistent with the models of Hasenclever et al. (2014) or Dunn
456 et al. (2000).

457 We have discussed above (Section 1) how the inversion of seismic velocity
458 perturbations to yield thermal anomalies may be affected by uncertainties in our
459 knowledge of several material parameters. The hydrothermal circulation model of
460 Hasenclever et al. (2014) is anchored to the model of Dunn et al. (2000) and is subject
461 to the same sources of uncertainty. Additionally, in spite of the sophistication of their
462 calculations, there are aspects that could be improved – for example, consideration of
463 anisotropy of permeability and distinct permeabilities for gabbros and sheeted dike
464 complexes may yield more realistic hydrothermal circulation patterns. Therefore, the
465 mismatch between the results from these models and the directly determined cooling
466 rates of this study is not surprising. Indeed, the direct determination of the spatial
467 distribution of cooling rates as a function of depth in this study provides important
468 boundary conditions that may be used to develop the next generation of such models.

469

470 **7. Summary and conclusions**

471

472 A new geospeedometer, based on the Mg content of plagioclase co-existing
473 with clinopyroxene, was applied to natural gabbroic rocks from three locations where
474 samples of the plutonic section of oceanic crust formed at modern fast-spreading

475 ridges are available. Samples from each location originate from different depths
476 below the DGB and the total depth interval covers the upper 836 m of the plutonic
477 section of the oceanic crust. The obtained cooling rates decrease significantly with
478 increasing sample depth (covering almost 4 orders of magnitude, ranging from ~ 1
479 $^{\circ}\text{C}\text{y}^{-1}$ for the shallowest samples to $0.0003^{\circ}\text{C}\text{y}^{-1}$ for the deepest samples). The fact
480 that this observation is very robust for three different locations along the EPR implies
481 a very comparable, and near steady-state, thermal structure in the off-axis region
482 along the EPR. Two independent methods (*Mg-in-Pl* and *Ca-in-Ol*) give the same
483 cooling rates for a given depth below the DGB, which implies that not only is the
484 relative trend of cooling rate with depth well constrained, but also that the absolute
485 values obtained for the cooling rates are robust. Both, the absolute cooling rates
486 determined from the deeper samples (>300 m below DGB), and the large decrease in
487 cooling rate with depth, are inconsistent with thermal models that include substantial
488 cooling by off-axis hydrothermal circulation. Instead, our data is consistent with
489 thermal models in which the lower crust cools conductively in the off-axis, implying
490 that most of the latent heat is removed by hydrothermal circulation at the top of an
491 axial melt lens and heat conduction becomes the dominant process of heat removal
492 with increasing depth and away from the ridge axis. Our observational results provide
493 important additional boundary conditions for modeling calculations of thermal
494 structure of fast spreading ridge systems (e.g., MacLennan et al., 2005; Hasenclever et
495 al., 2014), and should help in the development of the next generation of models.

496

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498

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505

506 **Appendix A Model parameters and input conditions for the diffusion model**

507

508 A detailed description of the used diffusion model may be found in Faak et al. (2014),
 509 here we briefly summarize the modeling approach and the input parameters used.

510

511 *Diffusion equation and diffusion coefficient*

512 The diffusion of Mg in plagioclase is coupled with the anorthite content, X_{An} ,
 513 in plagioclase, which needs to be accounted for in the diffusion equation (Eq. (7) in
 514 Costa et al., (2003)):

$$515 \quad J_{Mg} = D_{Mg} \frac{\partial C_{Mg}}{\partial x} + L_{Mg} A \frac{\partial X_{An}}{\partial x} \quad (\text{Eq. A.1})$$

516 where D_{Mg} = diffusion coefficient of Mg in plagioclase, C_{Mg} = concentration of Mg in
 517 plagioclase, x = distance and A = factor to describe the dependence of the partition
 518 coefficient on X_{An} . Their factor L_{Mg} is a phenomenological coefficient, equivalent to
 519 $(D_{Mg} C_{Mg})/RT$ (see Costa et al. (2003) for details). The resulting diffusion equation to
 520 describe the flux of Mg has been presented by Costa et al. (2003):

$$521 \quad \frac{\partial C_{Mg}}{\partial t} = \left(D_{Mg} \frac{\partial^2 C_{Mg}}{\partial x^2} + \frac{\partial C_{Mg}}{\partial x} \frac{\partial D_{Mg}}{\partial x} \right) - \frac{A}{RT} \left(D_{Mg} \frac{\partial C_{Mg}}{\partial x} \frac{\partial X_{An}}{\partial x} + C_{Mg} \frac{\partial D_{Mg}}{\partial x} \frac{\partial X_{An}}{\partial x} + D_{Mg} C_{Mg} \frac{\partial^2 X_{An}}{\partial x^2} \right)$$

522 (Eq. A.2)

523 where $t = \text{time}$.

524 The differential equation (Eq. (A.2)) can be solved numerically for one-
 525 dimensional diffusion in a plate by applying the method of central finite differences
 526 (Crank, 1975, Costa et al., 2008):

$$\begin{aligned}
 C_{i,j+1} = C_{i,j} + \frac{\Delta t}{(2\Delta x^2)} & \left[D_{i,j} (C_{i+1,j} - 2C_{i,j} + C_{i-1,j}) + \frac{1}{2} (C_{i+1,j} - C_{i-1,j}) (D_{i+1,j} - D_{i-1,j}) \right] \\
 & - \frac{A}{RT} \frac{\Delta t}{(2\Delta x^2)} \left[\frac{1}{2} D_{i,j} (C_{i+1,j} - C_{i-1,j}) (X_{An_{i+1,j}} - X_{An_{i-1,j}}) \right. \\
 & \left. + \frac{1}{2} C_{i,j} (D_{i+1,j} - D_{i-1,j}) (X_{An_{i+1,j}} - X_{An_{i-1,j}}) \right. \\
 & \left. + D_{i,j} C_{i,j} (X_{An_{i+1,j}} - 2X_{An_{i,j}} + X_{An_{i-1,j}}) \right]
 \end{aligned}$$

528 (Eq. A.3)

529 where $i = \text{step in distance}$ and $j = \text{step in time}$. $C_{i,j}$ and $D_{i,j}$ are the concentration of Mg
 530 in plagioclase and the diffusion coefficient of Mg in plagioclase at given i and j ,
 531 respectively.

532 We used the diffusion coefficient of Mg in plagioclase from Faak et al. (2013)
 533 that was determined experimentally in the compositional range of the lower oceanic
 534 crust:

$$D_{Mg}^{Pl} [\text{m}^2\text{s}^{-1}] = 1.25 \cdot 10^{-4} [\text{m}^2\text{s}^{-1}] \exp(-320924 [\text{Jmol}^{-1}] / (RT)) (a_{SiO_2})^{2.6}.$$

536 (Eq. A.4)

537

538 *Initial and boundary conditions*

539 The partition coefficient, $K_{Mg}^{Pl/Cpx}$, has been determined experimentally by
 540 Faak et al. (2013) and is given by the relationship:

$$\ln K_{Mg}^{Pl/Cpx} = -9219 [K] \frac{1}{T} + 1.6 + \frac{16913 [\text{Jmol}^{-1}]}{RT} X_{An} + \ln a_{SiO_2} \quad (\text{Eq. A.5})$$

542 This can be re-arranged to give the interface plagioclase composition as a function of
 543 the measured clinopyroxene composition, temperature and the silica activity in the
 544 system:

$$545 \quad C_{Mg}^H = C_{Mg}^{Cpx} \exp \left[-9219 \left[\frac{K}{T} \right] + 1.6 + \frac{16913 \left[\text{Jmol}^{-1} \right]}{RT} X_{An} + \ln a_{SiO_2} \right] \quad (\text{Eq. A.6})$$

546

547 An initial profile is calculated based on Eq. (A.5) at temperatures around
 548 1200°C, (the exact starting temperature T_{start} depends on the grain size of the
 549 plagioclase and is given in Table 1).

550

551 **Silica activity**

552 Solving equations (A.4-A.6) requires knowledge of the silica activity, a_{SiO_2} , as
 553 a function of temperature. Here, we assume a_{SiO_2} is constrained by the assemblage
 554 olivine + orthopyroxene as these are commonly observed phases in the samples
 555 studied. Thus, a_{SiO_2} is given by the reaction:



557 and the relationship proposed by Carmichael et al. (1970):

$$558 \quad \log a_{SiO_2} = \frac{\Delta G_r}{2.303RT} \quad (\text{Eq. A.8})$$

559 where ΔG_r is the Gibbs Free energy of the reaction given in equation (A.7). The
 560 silica activity defined by Eq (A.7) at 1 bar was calculated for pure Mg-endmembers
 561 (Faak et al., 2013 report that the incorporation of a molar fraction of 0.1 of the
 562 respective Fe-endmembers increases a_{SiO_2} by about 10%) and for different
 563 temperatures using the data set of Ghiorso and Sack (1995) to determine ΔG_r , and the
 564 resulting dataset fitted by a 2nd-order polynomial to obtain:

565
$$a_{SiO_2} = -4.8690410^{-7} T^2 + 1.5157010^{-3} T - 0.618707$$
 (Eq. A.9)

566

567

568 **Appendix B**

569

570 Section 4.2 discusses how the lack of a strong curvature of the Mg-profiles and
571 the high Mg-concentrations at the rims, which is measured in plagioclase from the
572 lower oceanic crust, can be explained by two possible scenarios. This Appendix
573 illustrates the modeling approaches used to simulate these different scenarios, and
574 shows that this leads to only small differences in the extracted cooling rates.

575 We consider two scenarios. Scenario (A) assumes a linear cooling history up to a
576 certain temperature, T_{Crim} , which can be calculated from the measured Mg-
577 concentration at the rim of a plagioclase crystal (Eq. (4) in Faak et al., 2014). Below
578 T_{Crim} cooling continues with an increased cooling rate, fast enough such that the
579 diffusion profile, which was evolved up to T_{Crim} , is not significantly modified.

580 Scenario (B) assumes a cooling history that is linear down to temperatures low
581 enough that the diffusion profile is not effectively changed by diffusion anymore
582 (around 600°C for the diffusion of Mg in plagioclase and geological reasonable
583 cooling rates). However, in this scenario there is a certain temperature T_{closed} (>600°C)
584 below which the exchange of Mg between plagioclase and clinopyroxene ceases.
585 However, diffusion of Mg within the plagioclase continues and tends to homogenize
586 the concentration of Mg in plagioclase (or for plagioclase zoned in X_{An} , tends to
587 equilibrate the plagioclase Mg content with the An content). Thus, in model B, the
588 system continues cooling down to 600°C at the same cooling rate, but diffusion of Mg
589 below T_{closed} is simulated in a closed system (= plagioclase has closed boundary

590 conditions). An additional constraint on this scenario comes from the argument of
591 mass balance: if Mg cannot be exchanged below T_{closed} , the total amount of Mg in
592 plagioclase at this temperature (Mg_{tot}) *must* equal the final amount of Mg in
593 plagioclase, $Mg_{observed}$. Since we model diffusion in 1D, assuming a plane sheet
594 geometry, the amount of Mg in plagioclase is calculated along a Mg-profile as $Mg =$
595 $Mg(x)*dx$, where $Mg(x)$ is the concentration of MgO in wt% at a certain grid point
596 and dx is the distance between grid points. We define that mass balance is fulfilled if
597 $Mg_{tot} = Mg_{observed} \pm 1\%$.

598 The two different approaches were applied to three theoretical plagioclase
599 crystals: P1 has a weakly curved Mg-profile with 0.04 wt% MgO at the rims and 0.06
600 wt% MgO at the core (Fig. B1 a-c); P2 has a homogeneous Mg-profile at 0.04 wt%
601 MgO (Fig. B1 d-e) and P3 has a homogeneous Mg-profile at 0.02 wt% MgO (Fig. B1
602 g-i). When modeled as scenario (A), the best fit between modeled and observed data
603 is obtained for the following values: $dT/dt=0.008^{\circ}C_y^{-1}$ and $T_{Crim} = 910^{\circ}C$ for P1 (Fig.
604 B1a), $dT/dt = 0.0003^{\circ}C_y^{-1}$ and $T_{Crim} = 910^{\circ}C$ for P2 (Fig. B1d), and $dT/dt =$
605 $0.00002^{\circ}C_y^{-1}$ and $T_{Crim} = 810^{\circ}C$ for P3 (Fig. B1g). Note that for P2 and P3, where the
606 observed Mg-profiles are homogeneous, only maximum cooling rates can be
607 obtained. For homogeneous X_{An} , these homogeneous Mg-profiles represent the
608 equilibrium Mg-concentration at T_{Crim} and would not be changed by slower cooling.
609 When modeling scenario (B), once a cooling rate for a model run is fixed, there is
610 only limited variation for T_{closed} , i.e. for a given cooling rate, T_{closed} can only be
611 changed by a few $^{\circ}C$ without violating the mass balance criteria (in fact, if Mg_{tot}
612 exactly equals $Mg_{observed}$, then T_{closed} is fixed for a given cooling rate). Furthermore,
613 diffusion below T_{closed} will tend to homogenize the Mg-profile, i.e. decrease the Mg-
614 concentration at the core and increase the Mg-concentration at the rims. Thus, mass

615 balance can only be fulfilled if T_{closed} is lower than T_{Crim} and dT/dt is faster than dT/dt
616 in scenario (A). However, if dT/dt is too fast, mass balance cannot be attained for any
617 temperature T_{closed} , because Mg is not removed efficiently enough from the core. Even
618 for cases where mass balance can be attained, dT/dt may be too fast to allow for
619 sufficiently diffusion of Mg in the closed system to allow a good match between the
620 observed and the modeled profile (e.g. Fig. B1 c, f, and i). In the end, there is only a
621 small range of combinations of dT/dt and T_{closed} that lead to acceptable fits to the
622 observed Mg-profiles (e.g. Fig. B1 b, e, and h).

623 To obtain an estimate of the level of variation of dT/dt and T_{Crim} / T_{closed} that is
624 introduced by the choice of modeled scenario, we used dT/dt and T_{Crim} from modeling
625 scenario (A) (Fig. B1 a, d, and g) as a starting point for modeling scenario (B). Then
626 dT/dt was iteratively increased, and T_{closed} was adjusted for each run until mass
627 balance was fulfilled. For each plagioclase, the last fit that was still considered
628 ‘acceptable’ is shown in Fig. B1 b, e, and h. The next run with fits that were not
629 accepted any more is shown in Fig. B1 c, f and i. The difference between dT/dt and
630 T_{Crim} / T_{closed} from scenario (A) and those obtained from last accepted fit from scenario
631 (B) show the small difference in cooling rate extracted using these different
632 approaches as discussed in the main text.

633

634

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- 791

792 **Table Captions**

793

794 **Table 1.** Summary of the cooling rates that were obtained from each Mg-profile in
795 plagioclase from the three sample suites. The sample depth is given in m below the
796 DGB (mbDGB) of the individual sample suite. The given temperature range refers to
797 the *T*-interval over which the diffusion process was modeled (see Section 4.2 and
798 Appendix B for details). *italics* = the observed Mg-profile is very homogeneous, so
799 that only a maximum estimate of the cooling rate can be obtained.

800

801 **Figure Captions**

802

803 **Fig. 1.** Schematic sketch to illustrate how different modes of heat extraction and
804 crystallization in the lower oceanic crust result in different trends for cooling rates as
805 a function of depth. Panel (a) shows a gabbro glacier type model that assumes most of
806 the latent heat to be released in the AML and removed by hydrothermal circulation
807 above this body. To emphasize the potential for heat conduction to be the dominant
808 process of heat removal in the off-axis region in this model the green line in the
809 middle panel shows this scenario. It should be noted that the gabbro glacier crustal
810 accretion model is also permissive of efficient off-axis hydrothermal cooling (e.g.
811 Phipps Morgan and Chen, 1993; Hasenclever et al., 2014). Panel (b) shows a sheeted
812 sill type model that assumes less efficient heat extraction above the AML and more
813 efficient heat extraction by off-axis hydrothermal circulation in the lower crust. The
814 middle panel compares (schematically and not to scale) the expected trends for
815 vertical variation of cooling rate in these hypothetical models with different

816 efficiencies of hydrothermal cooling at depth. (For interpretation of the references to
817 color in this figure, the reader is referred to the online version of this article.)

818

819 **Fig. 2.** Geographical and structural maps with locations of the three sample sites. (a)
820 Overview of the location of the Hess Deep, the Pito Deep, and IODP drill Site 1256D
821 (red marks). (b) Map of the Galapagos triple junction in the eastern equatorial Pacific
822 Ocean showing major tectonic boundaries and the location of the Hess Deep Rift (red
823 box). (c) Map of the Easter Microplate in the southern Pacific Ocean showing major
824 tectonic boundaries and the location of the Pito Deep (red box). (d and e) Simplified
825 geological map of the Northern wall of the Hess Deep Rift showing the dive tracks
826 with general lithological units and sample locations (based on Gillis et al., 2001). The
827 blue dashed line represents the inferred location of the DGB in this area and the blue
828 shaded area represents a region where only gabbroic rocks were mapped along dive
829 tracks. Contour lines represent depth below sea level given in meters. (f and g)
830 Simplified geological map of the Pito Deep area B showing the dive tracks with
831 general lithological units and sample location (based on Perk et al., 2007). Base maps
832 in (a)-(c) were created with GeoMapApp. (For interpretation of the references to color
833 in this figure, the reader is referred to the online version of this article.)

834

835 **Fig. 3.** Examples of the different Mg-profile shapes observed in plagioclase in
836 samples from different depth below the DGB (in meters, gray numbers besides the
837 boxes) from Hess Deep (left column) and Pito Deep (right column). Open circles
838 show the Mg-concentrations that were measured along full profiles in plagioclase.
839 Solid pink lines show the modeled diffusion profile that matches best with the
840 measured profile. The inset in each box shows the X_{An} -content that was measured

841 along the same transect as the Mg-profile. (For interpretation of the references to
842 color in this figure, the reader is referred to the online version of this article.)

843

844 **Fig 4.** Diagram to show the obtained cooling rates (plotted as $\log dT/dt$) versus the
845 sample depth below the DGB. Arrows indicate that only maximum cooling rates
846 could be obtained, because the measured Mg-profiles were homogeneous (see text for
847 discussion). Light pink symbols show data from sample 022005-1052 that is regarded
848 as an outlier (see text for discussion). For comparison, cooling rates obtained from the
849 *Ca-in-olivine geospeedometer* (Coogan et al., 2007) are shown for the depth range,
850 where data from that study is available. (For interpretation of the references to color
851 in this figure, the reader is referred to the online version of this article.)

852

853 **Fig. 5.** Comparison of the cooling rates obtained from diffusion modeling of Mg in
854 plagioclase in the natural samples with different thermal models. M = cooling rates
855 from Maclennan et al. (2005) obtained from a thermal model with a thermal
856 diffusivity of $8 \times 10^{-7} \text{ m}^2\text{s}^{-1}$ and various amounts of hydrothermal cooling off-axis
857 ($50\text{-}71 \text{ kWm}^{-1}$; only their model 4 has no hydrothermal cooling off-axis). D = cooling
858 rates calculated from the isotherms separation of Dunn et al. (2000) for a cooling
859 interval of $1000\text{-}600^\circ\text{C}$. H = cooling rates calculated from the isotherm separation of
860 Hasenclever et al. (2014) for a cooling interval of $1000\text{-}600^\circ\text{C}$. T&S = simple
861 conductive half-space model based on Eq. 4.124 in Turcotte and Schubert (2002) with
862 a thermal diffusivity of $1 \times 10^{-6} \text{ m}^2\text{s}^{-1}$. The top of the lower crust (i.e. the dike/gabbro
863 boundary) is held at a constant temperature of 400°C , i.e. the temperature in the dikes
864 is assumed to be 400°C (based on temperature estimates for hydrothermal fluids (e.g.
865 Von Damm, 2000) and from petrology of alteration of dikes (e.g. Gillis, 1995) in the

866 area). The initial temperature of the entire crust and upper mantle is assumed to be
867 1300°C. Cooling rates are extracted for the T -interval of 900-600°C and 700-600°C.
868 Tini layered = conductive cooling model with a layered initial temperature
869 distribution. The temperature at the top (= seafloor and lava) is held constant at 4°C.
870 The initial temperature profile is set based on the expected profile at the edge of the
871 AML as follows: layer 1 (dikes): 1 km thick, 400°C; layer 2 (gabbros): 4 km thick,
872 1250°C; layer 3 (upper mantle): 15 km thick, 1300°C. Conductive cooling was
873 modeled from these initial and boundary conditions and cooling rates were extracted
874 at around 900°C for all depth and around the average obtained rim closure
875 temperature (T_{crim}) for each sample depth of the Hess Deep (HD) and Pito Deep (PD)
876 sample suite. Note that the difference between the conductive model 4 from
877 MacLennan et al., 2005, and the other conductive models results from the fact that
878 their model 4 is a 2D model that additionally considers horizontal heat transport, and
879 latent heat of crystallization. (For interpretation of the references to color in this
880 figure, the reader is referred to the online version of this article.)

881

882 **Fig. 6.** Diagram to illustrate a bounding constraint on the thermal structure in the
883 lower crust in the off-axis region. For each sample depth, the average cooling rate
884 obtained from the *Mg-in-plagioclase geospeedometer* was used to determine the time
885 needed to cool from $(T_{crim} + 100^\circ\text{C})$ to T_{crim} . The distance the crust would move in
886 this time was calculated assuming a half spreading rate of 65 mmy^{-1} to yield the
887 horizontal separation of two isotherms that are 100°C apart. In order to locate the
888 limiting distance from the ridge at which these temperatures may be attained, cooling
889 above $(T_{crim} + 100^\circ\text{C})$ was assumed to be instantaneous (i.e. $(T_{crim} + 100)^\circ\text{C}$ is
890 assumed to be attained at zero time at the ridge axis itself at the respective depths).

891 Thus, these values provide a minimum constraint on the time needed for the crust to
892 cool down to $T_{c_{rim}}$ at a given depth, and the distance from the ridge where this may be
893 attained for the known spreading velocity. The bold green and blue lines represent the
894 minimum time (and equivalently, distance) required to cool down to 750°C, 700°C
895 and 650°C respectively, at depths where information on cooling rates around each of
896 these temperatures has been frozen in the plagioclase record. An alternate, less
897 extreme case is where cooling occurs linearly from the solidus temperature of
898 $\sim 1200^\circ\text{C}$ to $T_{c_{rim}}$. $T_{c_{rim}}$ at any given depth is attained much later (at a greater distance
899 from the ridge) in this case and tips of horizontal arrows drawn at each depth mark
900 these points. Thus, for any conceivable cooling history, the inferred temperature at a
901 given depth has to lie within the gray region to be consistent with the thermal history
902 recorded in the plagioclase crystals. For comparison, the position of the 1000-400°C
903 isotherms from Hasenclever et al. (2014) (marked “H”) for the depth interval
904 corresponding to our study are shown. (For interpretation of the references to color in
905 this figure, the reader is referred to the online version of this article.)

906

907 **Fig B1.** Illustration of the effect of the choice of modeling scenario on the obtained
908 cooling rate. The diagrams show synthetic Mg-concentration profiles for three
909 different plagioclase crystals (P1 = weakly curved Mg-profile; P2 = homogeneous
910 Mg-profile at 0.4 wt% MgO and P3 = homogeneous Mg-profile at 0.2 wt% MgO).
911 For simplicity, all plagioclase crystals are assumed to have homogeneous $X_{An} = 0.6$
912 and to be in contact with Cpx with 14 wt% MgO. The silica activity of the system is
913 assumed to be constraint by the assemblage Ol+Opx. The pink bold lines show best
914 fit profiles for modeling scenario (A), i.e., $T_{c_{rim}}$ can be obtained using Eq. (4) from
915 Faak et al. (2014) with the Mg-concentration at the rim of the plagioclase. The light

916 pink bold lines show best fit profiles for modeling scenario (B), i.e., modeling is
917 continued to 600°C, but at temperature T_{closed} ($T_{rim} > T_{closed} > 600^{\circ}\text{C}$), the system was
918 modeled with closed boundaries (see text for discussion). (For interpretation of the
919 references to color in this figure, the reader is referred to the online version of this
920 article.)

Highlights

- We use diffusion modeling of Mg in plagioclase to determine cooling rates
- Cooling rates were obtained from natural samples of the lower oceanic crust
- Obtained cooling rates significantly decrease with increasing sample depth
- Our data is best explained by conductive thermal models
- Our data is inconsistent with substantial cooling by off-axis hydrothermal circulation

Figure 1

[Click here to download Figure: Fig1_Models_cooling_rate.pdf](#)

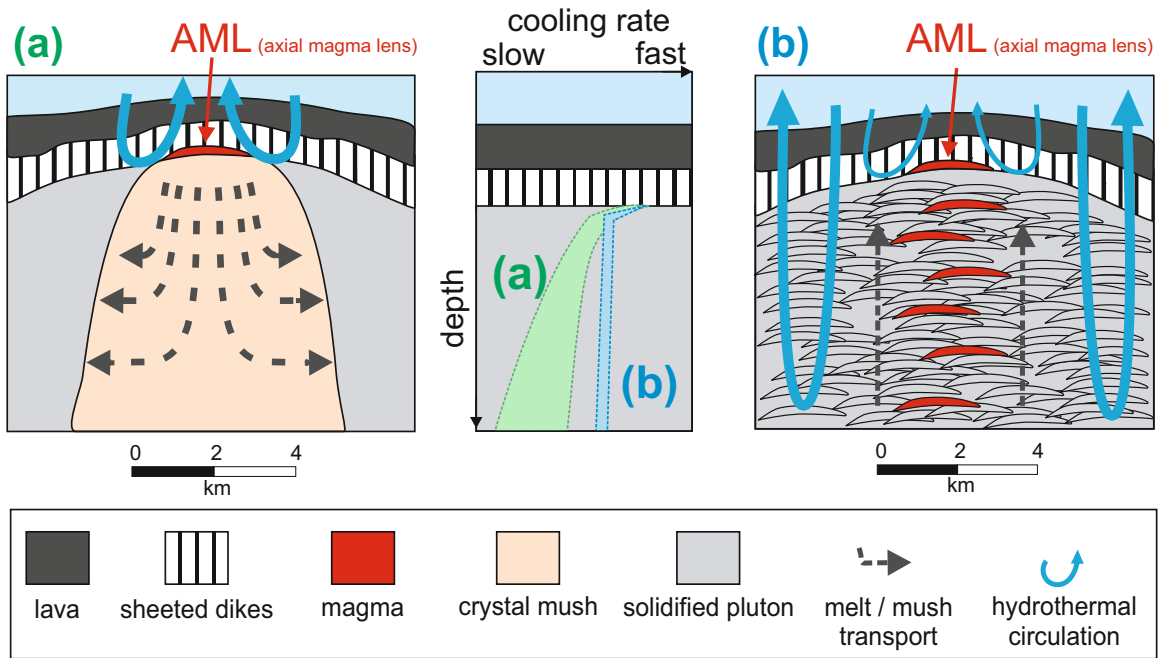


Figure 2

[Click here to download Figure: Fig2_Maps.eps](#)

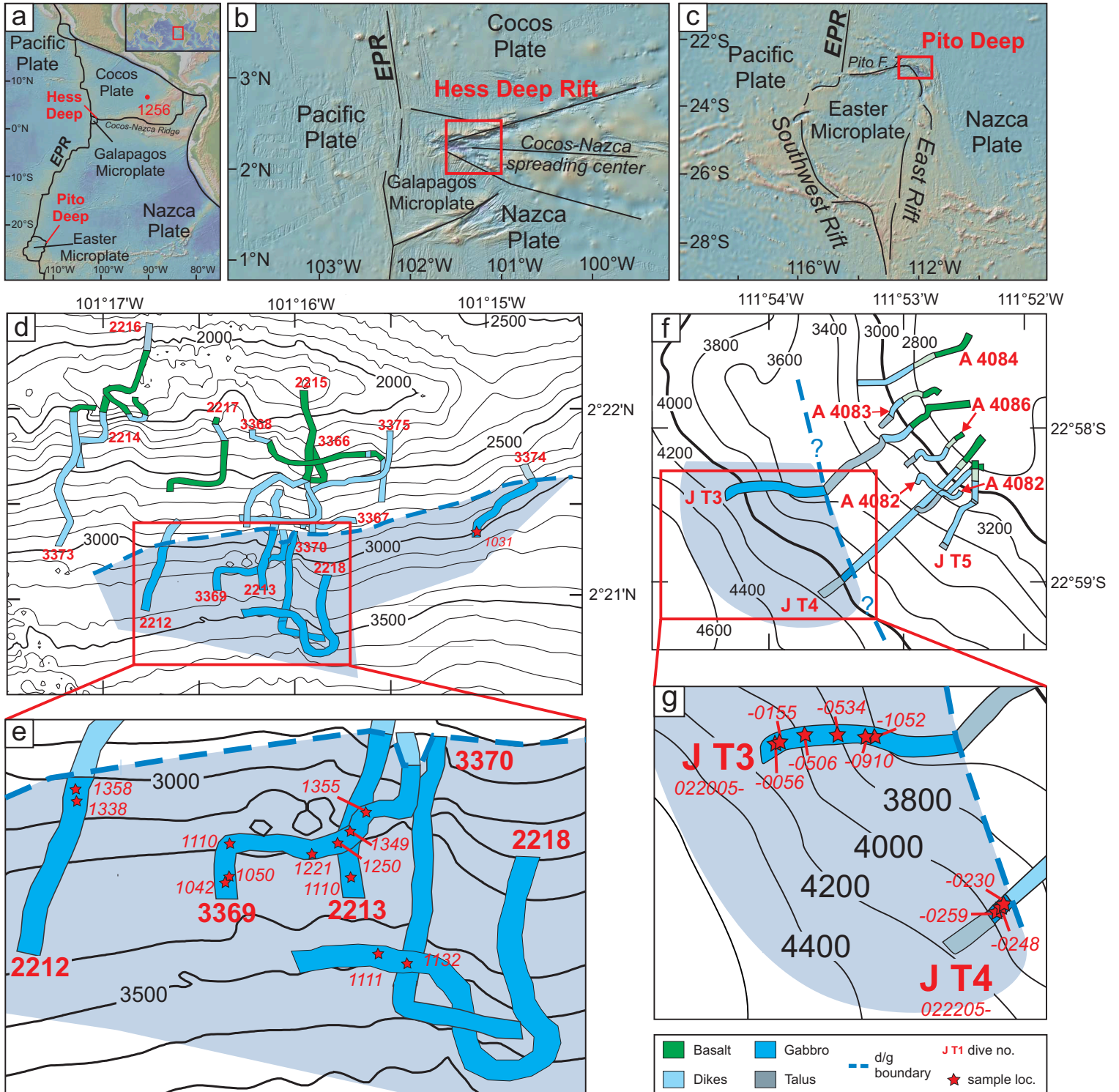


Figure 3

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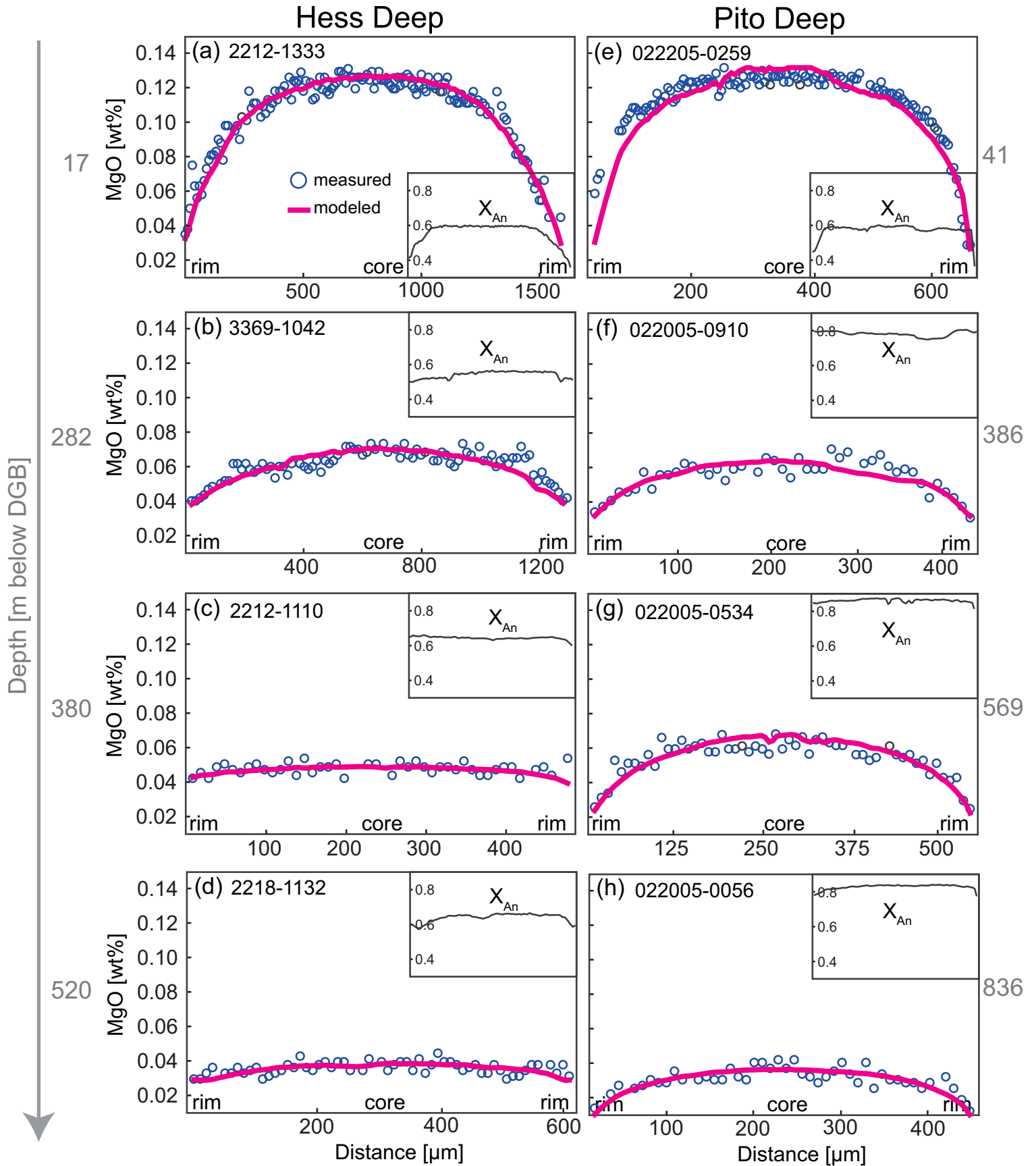


Figure 4

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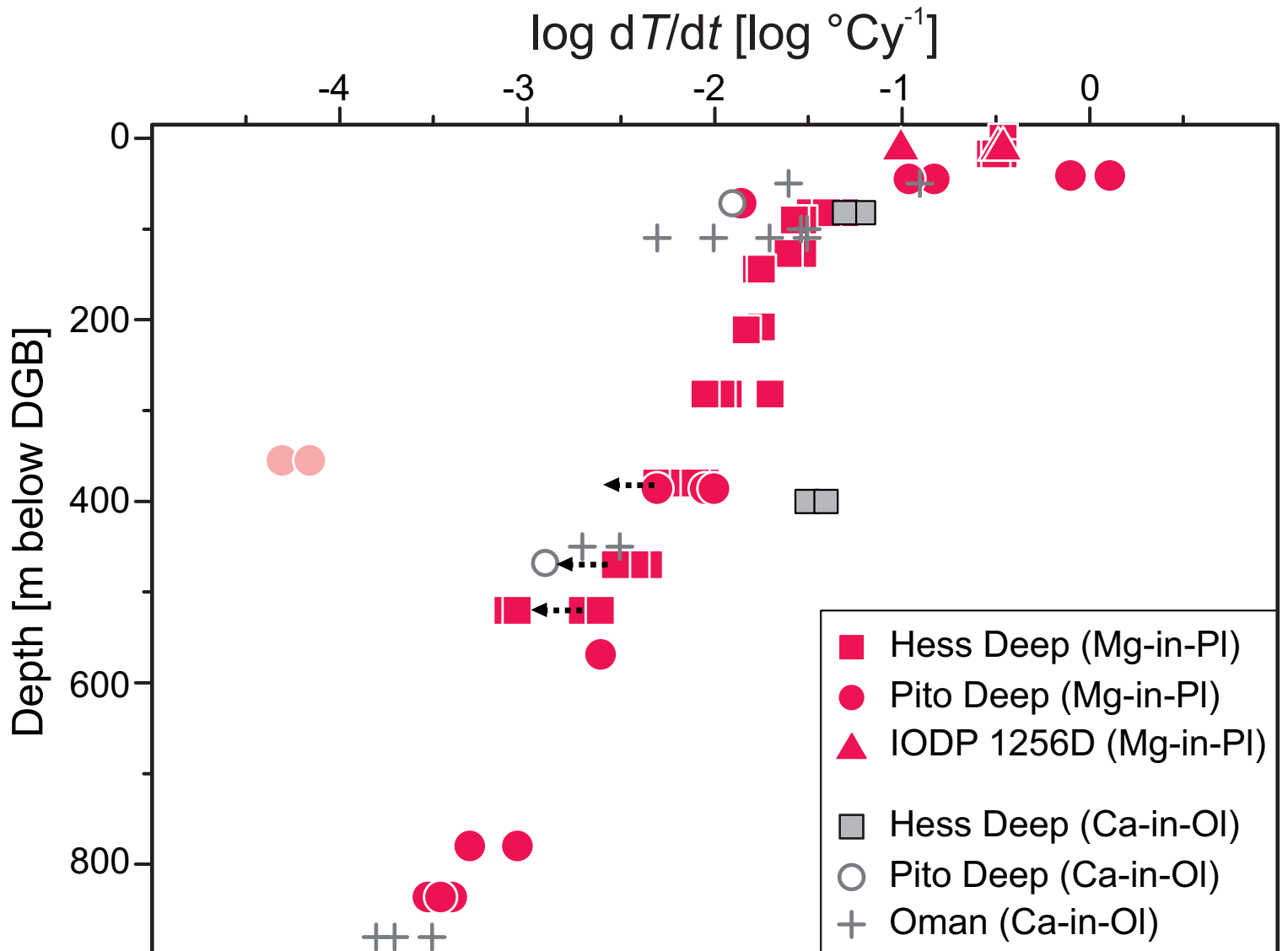


Figure 5

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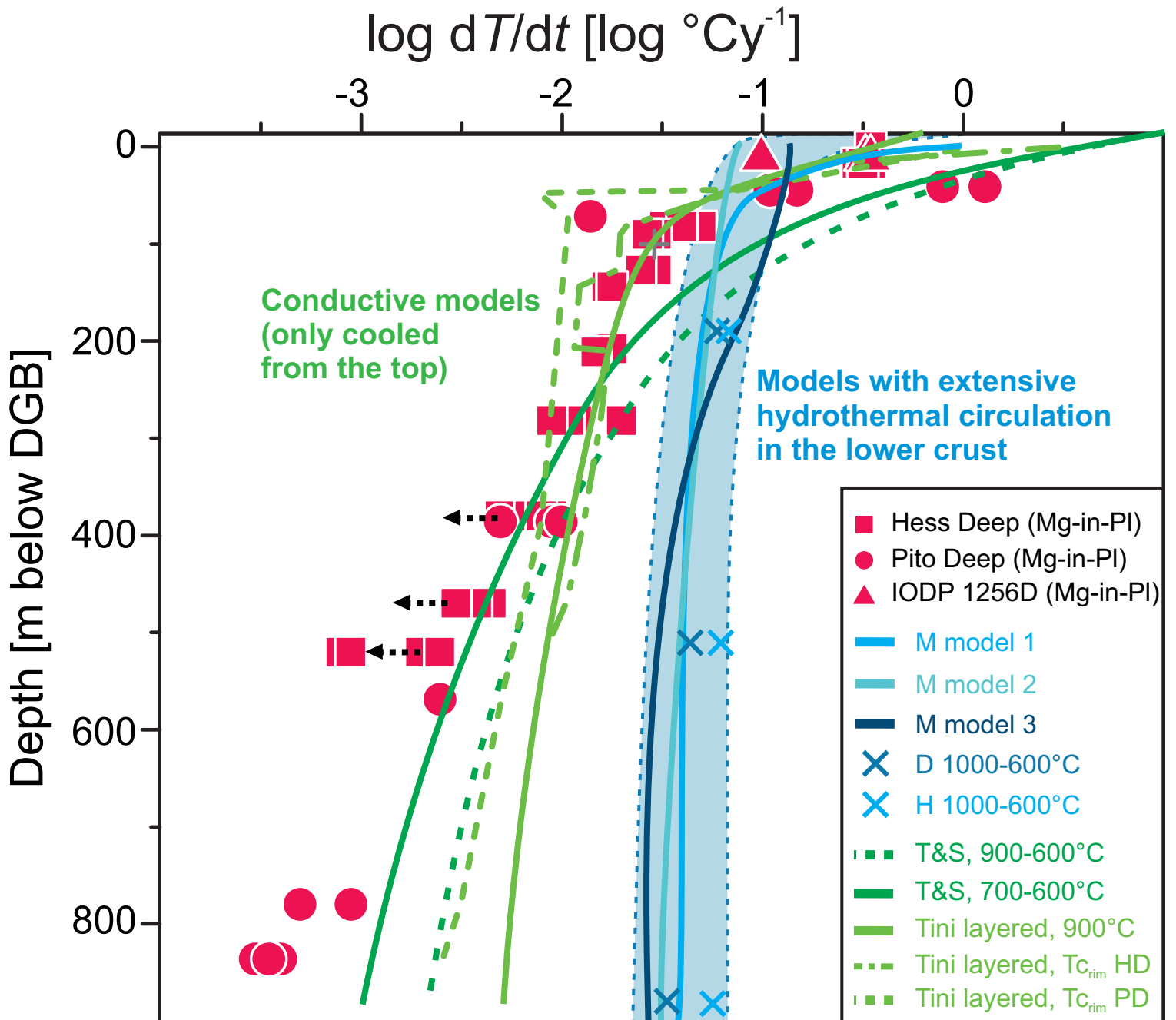


Figure 6

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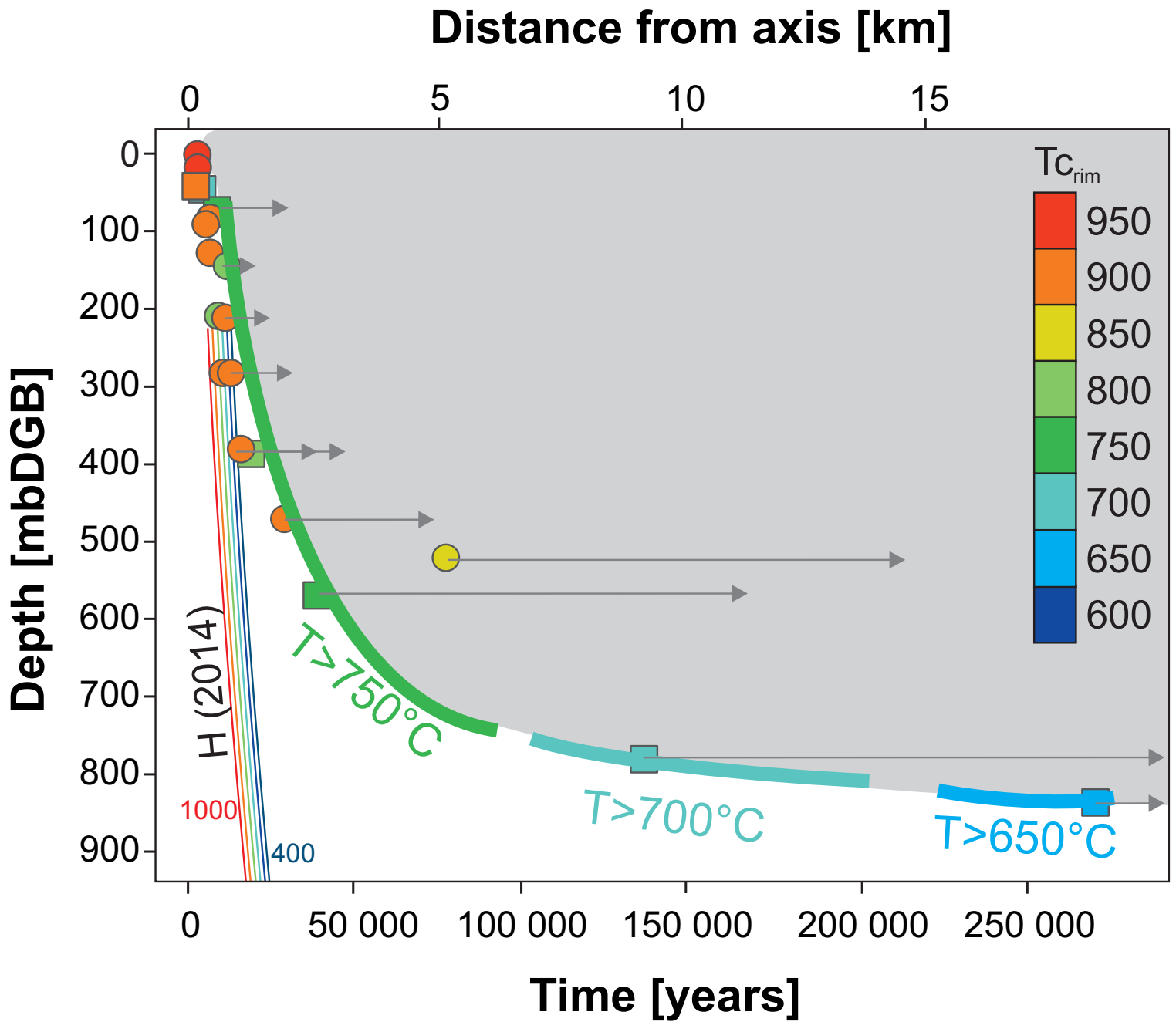


Figure A1

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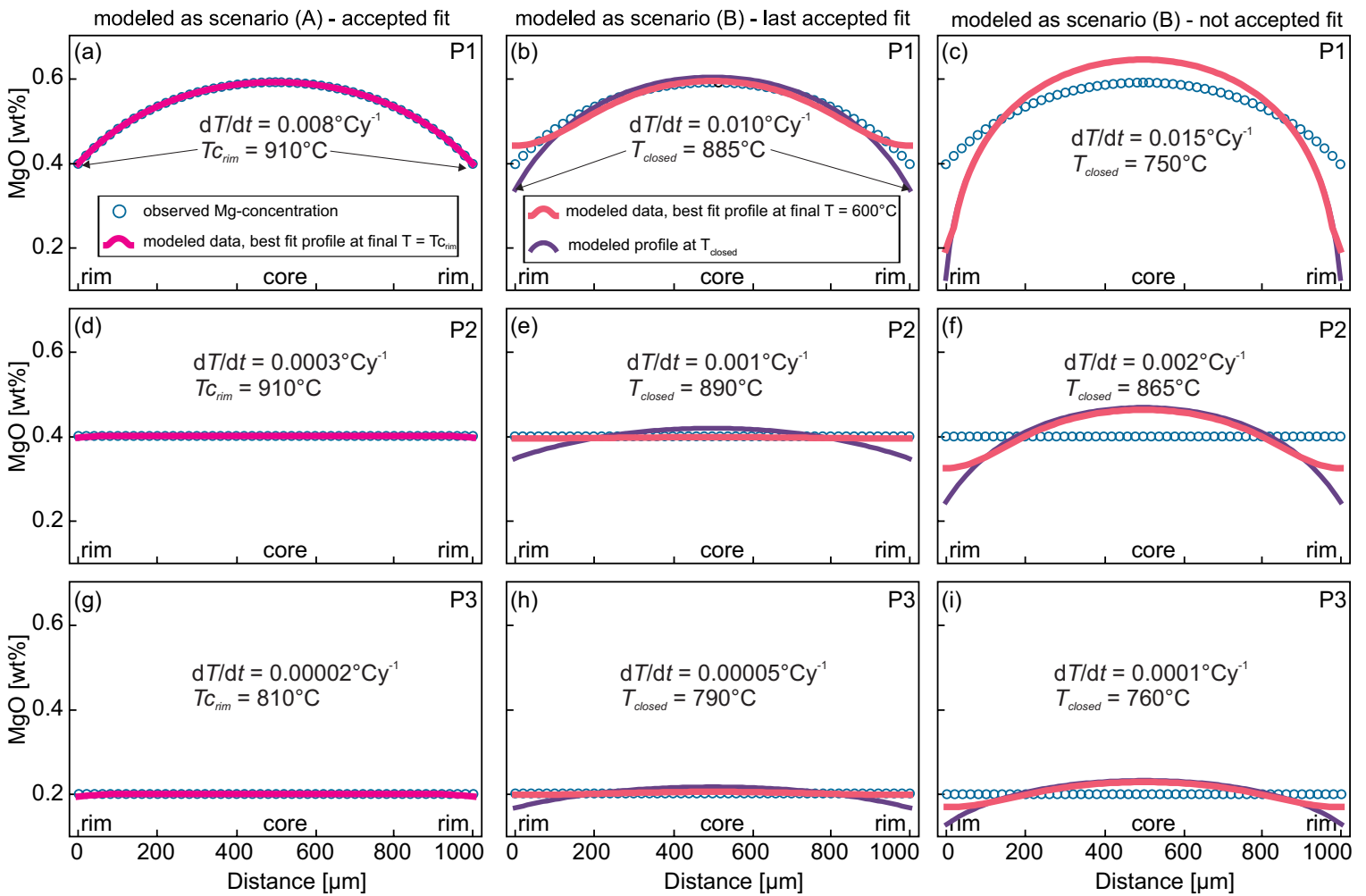


Table 1
[Click here to download Table: Table 1.docx](#)

Table 1. Summary of the cooling rates that were obtained from each used Mg-profile in the individual plagioclase crystals from the three sample suites. The sample depth is given in m below the DGB of the individual sample suite (mbDGB). The given temperature range refers to the *T*-interval over which the diffusion process was modeled (see Section 4.2 and Appendix I for details). *italics* = the observed Mg-profile is very homogeneous, so that only a maximum estimate of the cooling rate can be obtained.

Sample	Depth [mbDGB]	Plag	Profile	Length [μm]	dT/dt [$^{\circ}\text{C}/\text{y}$]	log dT/dt	T-interval [$^{\circ}\text{C}$]
Hess Deep							
2212-1358	0	Pl4	2	610	0.35	-0.46	1155-960
2212-1338	17	Pl1	1	500	0.35	-0.46	1133-940
2212-1338	17	Pl3	1	1590	0.3	-0.52	1223-950
2212-1338	17	Pl3	2	1550	0.32	-0.49	1221-800
3369-1355	82	Pl1	2	860	0.05	-1.30	1179-980
3369-1355	82	Pl3	1	530	0.033	-1.48	1145-870
3369-1355	82	Pl3	2	380	0.04	-1.40	1121-900
3369-1349	90	Pl1	1	590	0.03	-1.52	1152-900
3369-1349	90	Pl3	2	350	0.027	-1.57	1115-890
3374-1031	127	Pl1	1	590	0.028	-1.55	1152-920
3374-1031	127	Pl1	2	260	0.03	-1.52	1094-910
3374-1031	127	Pl2	2	250	0.025	-1.60	1092-920
3369-1250	144	Pl1	1	265	0.017	-1.77	1096-840
3369-1250	144	Pl2	1	190	0.018	-1.74	1072-820
3369-1221	208	Pl1	1	207	0.018	-1.74	1078-830
3369-1110	211	Pl1	1	460	0.015	-1.82	1135-920
3369-1050	282	Pl2	2	660	0.02	-1.70	1160-900
3369-1050	282	Pl3	1	960	0.01	-2.00	1187-950
3369-1050	282	Pl3	2	450	0.012	-1.92	1133-930
3369-1042	282	Pl1	1	1410	0.011	-1.96	1211-920
3369-1042	282	Pl1	2	1590	0.009	-2.05	1223-890
2213-1110	380	Pl2	1	640	0.005	-2.30	1158-940
2213-1110	380	Pl3	1	470	0.007	-2.15	1136-915
2213-1110	380	Pl4	2	510	0.009	-2.05	1142-920
2213-1110	380	Pl6	2	560	0.008	-2.10	1149-920
2218-1111	470	Pl1	1	350	0.0045	-2.35	1115-910
2218-1111	470	Pl1	2	690	0.0038	-2.42	1164-910
2218-1111	470	Pl2	2	510	0.003	-2.52	1142-900
2218-1132	520	Pl2	2	105	0.002	-2.70	1030-850
2218-1132	520	Pl3	1	600	0.0008	-3.10	1154-865
2218-1132	520	Pl3	2	300	0.0009	-3.05	1104-870
2218-1132	520	Pl5	1	170	0.0025	-2.60	1064-840
Pito Deep							
022205-0259	41	Pl2	1	480	1.3	0.11	1188-930
022205-0259	41	Pl2	2	702	0.8	-0.10	1215-900
022205-0248	45	Pl2	2	888	0.15	-0.82	1182-640
022205-0248	45	Pl4	1	200	0.11	-0.96	1076-720
022205-0230	72	Pl1	1	370	0.014	-1.85	1119-780

022005-1052	355	PI2	2	420	0.00005	-4.30	1128-730
022005-1052	355	PI3	2	500	0.00007	-4.15	1141-720
022005-0910	386	PI1	2	240	0.009	-2.05	1089-835
022005-0910	386	PI2	2	180	0.01	-2.00	1068-810
022005-0910	386	PI4	2	460	0.005	-2.30	1135-800
022005-0534	569	PI2	1	560	0.0025	-2.60	1149-760
022005-0155	780	PI1	2	830	0.0009	-3.05	1177-710
022005-0155	780	PI2	1	850	0.0005	-3.30	1178-720
022005-0056	836	PI2	2	830	0.0004	-3.40	1177-680
022005-0056	836	PI3	1	470	0.0003	-3.52	1136-690
022005-0056	836	PI5	1	840	0.00035	-3.46	1178-670
IODP Site 1256D							
216R1 15-20	12.1	PI3	1	460	0.1	-1.00	1135-860
216R1 49-57	12.4	PI1	1a	313	0.33	-0.48	1107-810
216R1 49-57	12.4	PI1	1b	313	0.35	-0.46	1122-820