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1   **Chemical differentiation, cold storage and remobilization of magma in the Earth's crust**
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7
8   **The formation, storage and chemical differentiation of magma in the Earth's crust is of**
9   **fundamental importance in igneous geology and volcanology. Recent data are challenging the**
10 **high melt fraction 'magma chamber' paradigm that has underpinned models of crustal magmatism**
11 **for over a century, suggesting instead that magma is normally stored in low melt fraction 'mush**
12 **reservoirs'.[1-9] A mush reservoir comprises a porous and permeable framework of closely packed**
13 **crystals with melt present in the pore space.[1,10] However, many common features of crustal**
14 **magmatism have not yet been explained by either the 'chamber' or 'mush' reservoir concepts.[1,11]**
15 **Here we show that reactive melt flow is a critical, but hitherto neglected, process in crustal mush**
16 **reservoirs, occurring because buoyant melt percolates upwards through, and reacts with, the**
17 **crystals.[10] Reactive melt flow in mush reservoirs produces the low crystallinity, chemically**
18 **differentiated (silicic) magmas which ascend to form shallower intrusions or erupt to the**
19 **surface.[11-13] The magmas can host much older crystals, stored at low and even sub-solidus**
20 **temperatures, consistent with crystal chemistry data.[6-9] Changes in local bulk composition caused**
21 **by reactive melt flow, rather than significant increases in temperature, produce the rapid increase**
22 **in melt fraction that remobilizes these cool- or cold-stored crystals. Reactive flow can also**
23 **produce bimodality in magma compositions sourced from mid- to lower-crustal reservoirs.[14,15]**
24 **Trace element profiles generated by reactive flow are similar to those observed in a well-studied**
25 **reservoir now exposed at the surface.[16] We propose that magma storage and differentiation**
26 **primarily occurs by reactive melt flow in long-lived mush reservoirs, rather than by the commonly**
27 **invoked process of fractional crystallisation in magma chambers.[14]**
28   Magma reservoirs occur at several depths within the crust and typically grow incrementally
29 through the intrusion of dykes or sills.[1,11,13,16,17] High melt fractions must sometimes be present in
30 these reservoirs to produce eruptible, low-crystallinity magmas.[1,7,8,9,13] However, geophysical data
31 suggest that reservoirs have low melt fraction even beneath active volcanoes[2-5] and crystal
32 chemistry data indicate that long-term magma storage occurs at low or even sub-solidus
33 temperature.[6-9] High melt fractions are therefore ephemeral, yet geochemical models typically
34 assume differentiation occurs by crystal fractionation from low-crystallinity magmas;[11,14] moreover,
35 geochronological data demonstrate that crustal magma reservoirs can be long-lived, spanning
36 hundreds of thousands to millions of years.[17-21] Existing models of crustal magma storage and
37 differentiation cannot reconcile these conflicting observations.
38   We use numerical modelling to investigate the storage and chemical differentiation of magma in
39 crustal reservoirs. The model describes repeated intrusion of mafic to intermediate sills into the
40 mid- to lower crust,[12,13,16,21-23] the associated transport of heat via conduction and advection and, in a
41 key advance, mass transport via reactive flow of buoyant melt through the compacting crystal
42 framework.[10] Transport of chemical components with the melt modifies the local bulk composition,
43 and melt fraction changes in response to the chemical reactions that maintain local thermodynamic
44 equilibrium. Phase behaviour is modelled using a two-component, eutectic phase diagram that,
45 although much simplified compared to natural systems, captures the critically important impact of
46 bulk composition on melting behaviour and the complex non-linear relationships between
47 composition, melt fraction and permeability (see Methods).[10] Melting relationships obtained from
48 the phase diagram approximate common crustal igneous systems (Extended Data Fig. 1). The
49 concentration of an incompatible trace element is also modelled assuming a constant partition
50 coefficient.
51   Typical results are shown in Figure 1 (see also Supplementary Video 1). In this example, 100m
52 thick basalt (mafic) sills are intruded randomly over a depth range of 600m, initially at 18km depth

53 and then around a depth that is controlled by the density contrast between intruding magma and
54 host mush, reflecting the evolving reservoir composition and melt fraction (see Methods). We
55 emplace 7.8km of basalt in total, at an average rate of 5mm·yr^{-1} typical of crustal magmatic
56 systems,[22-24] into solid crust with an initial geotherm of 20K·km^{-1}.[21-23] Our example was chosen to
57 facilitate comparison with data from a well-studied deep crustal section.[16,21] The key findings are
58 replicated over the depth range of 10-30km typical of many crustal magma reservoirs and following
59 intrusion of intermediate as well as mafic magma, using model parameters over a wide range
60 reasonable for such systems (see Methods and Extended Data Table 1).
61   Initially, following each sill intrusion, the melt fraction rapidly falls to zero so there is no
62 persistent magma reservoir (Supplementary Video 1 and Extended Data Figure 2). This is the
63 'incubation phase' of the incipient magma reservoir, observed also in models that neglect reactive
64 flow.[22,23] However, in our model, chemical differentiation occurs within each intrusion before it
65 solidifies, with more evolved melt (enriched in the incompatible trace element) accumulating at the
66 top of the intrusion, and more refractory and depleted crystals accumulating at the base. The rapid
67 increase in crystallinity traps the magma at the site of intrusion, but differentiation creates
68 compositional contrasts that cause the intrusion depth to progressively increase (Supplementary
69 Video 1 and Extended Data Figure 3a).
70   The incubation phase ends when the melt fraction is greater than zero between successive sill
71 intrusions, whereupon a magma reservoir has formed (Figure 1a; Supplementary Video 1). Melt is
72 now persistently present, but melt fraction remains low except for a brief period after each new
73 intrusion (Extended Data Figure 2b). The reservoir comprises a mush, rather than a high melt
74 fraction magma chamber. Reactive flow now significantly modifies the predicted reservoir behaviour
75 compared to previous models.[22,23]
76   Buoyant melt migrates upwards through the mush, accumulating in the upper part of the
77 reservoir because it cannot travel beyond the solidus isotherm where the melt fraction and
78 permeability fall to zero (Supplementary Video 1). Melt composition evolves as it flows into, and
79 reacts with, progressively cooler mush. Reactive flow reduces, or removes, early-formed
80 compositional contrasts, so the locally varying melt fraction controls the depth of later sill intrusions,
81 which decreases as melt migrates upwards (see Methods). This is the 'growing phase' of the
82 reservoir.
83   The growing phase ends when melt accumulates below the solidus isotherm to form a high melt
84 fraction (typically >0.7) layer overlying a thick (several km), low melt fraction (typically <0.2) mush
85 (Figure 1b and Supplementary Video 1). The melt-rich layer contains chemically differentiated felsic
86 magma and can grow to several 100's m in thickness. Although not captured by the model, buoyant
87 magma in the layer will be prone to leave the reservoir to produce shallower intrusions or volcanic
88 eruptions.[25,26] Once magma leaves, a new layer grows by the same mechanism (see Methods).
89   This is the 'active phase' during which the reservoir can deliver evolved, low crystallinity magma
90 (Extended Data Figure 2b). We suggest that, although geophysical surveys are probing active
91 reservoirs, they image only the low melt fraction mush;[2-5] the overlying high melt fraction layers are
92 not observed, because they are ephemeral and/or too thin to be resolved. Geophysical detection of
93 such a layer would suggest that magma mobilisation (and possible eruption) was imminent.[7]
94   When intrusion of new sills ends, reactive flow continues wherever the temperature is above the
95 solidus but, overall, the reservoir cools. This is the 'waning phase' (Supplementary Video 1;
96 Extended Data Figure 3b) that persists until the mush has completely solidified (Extended Data
97 Figure 2b). If exhumed, the resulting body of rock is termed a deep crustal section of which there
98 are several natural examples.[16,21]
99   During the active phase, the high melt fraction layer forms towards the top of the reservoir
100 where the temperature is low, rather than at the highest temperature (Figure 1b). This counter-
101 intuitive result is a consequence of reactive flow, whereby melt accumulation causes the local bulk
102 composition to evolve towards the eutectic. Melt composition in more chemically complex systems
103 will evolve towards other low-variance states such as cotectics, peritectics or multiple-saturation

104   points (see Methods), but the overall behaviour will be similar. A key finding here is that high melt
105   fraction layers in crustal mush reservoirs can form in response to changes in bulk composition
106   caused by reactive melt flow, rather than significant increases in temperature.
107   Magma in a high melt fraction layer contains c. 10% crystals (Figure 2a). These 'antecrysts' can
108   long pre-date magma formation, because they derive from crystallisation of early sills at the top of
109   the reservoir. Once formed, the antecrysts are stored at near- or sub-solidus temperature (i.e. 'cool'
110   or 'cold'; Figure 2b). The local temperature gradually increases in response to ongoing intrusion of
111   sills deeper in the reservoir and, eventually, exceeds the solidus. Soon afterwards, buoyant, evolved
112   melt, migrating upwards through the pore-space, accumulates around these older antecrysts,
113   causing the local melt fraction to increase rapidly and by far more than would be possible by heating
114   alone (Figure 2b).[6,7,18,27] Cold mush is remobilized here not by a significant increase in temperature,
115   but by buoyancy-driven reactive flow supplying evolved melt from deeper, more refractory parts of
116   the reservoir, where temperature can be high but the melt fraction remain low (Figure 2c).
117   Remobilization is primarily caused by changes in local bulk composition, rather than temperature.
118   In our example, melt accumulation forms a low crystallinity magma a few centuries after the local
119   temperature exceeds the solidus, yet the magma contains antecrysts formed up to c. 1–1.4Ma
120   earlier (Figure 2b). The range of antecryst ages reflects the timing of sill intrusions relative to the
121   timing of melt accumulation. Crystal chemistry data show cool or cold storage and remobilization of
122   older antecrysts hosted by younger felsic magma;[6-9] our results suggest that this could result from
123   reactive melt flow accumulating young, felsic melt within older mush. The antecrysts are not in
124   equilibrium with the younger melt, creating disequilibrium crystal textures such as partial resorption.
125   Flow of buoyant melt into the high melt fraction layer will drive convective overturn and
126   homogenisation before, or during, evacuation of magma, yielding a range of antecryst ages that may
127   span the entire reservoir history.[28]
128   Magmas in the high melt fraction layers have evolved composition. Conversely, magmas in the
129   sills shortly after intrusion have compositions close to the intruded basalt. Low crystallinity, mafic or
130   felsic magmas can therefore leave the reservoir, but not magmas with intermediate composition.
131   Many volcanic settings are characterised by bimodal volcanism (the 'Daly Gap'), especially in oceanic
132   settings (hotspots and island arc environments) and continental hotspots (Figure 3a).[14,15,29] Our
133   results suggest that compositional bimodality is another consequence of differentiation by reactive
134   melt flow in mush reservoirs. However, not all systems show bimodality.[30] Intermediate
135   compositions could result from magma mixing[15] or differentiation within multiple mush reservoirs
136   comprising a vertically extensive magmatic system.[1]
137   The modelled incompatible trace element concentration in the solidified reservoir shows a
138   characteristic pattern. Towards the base, the spiky signature produced by differentiation in each sill
139   during the incubation phase is preserved (Figure 3b). In the upper part, the profile is smoother and
140   shows depletion relative to the initial concentration, reflecting extraction of melt. The top shows
141   enrichment, reflecting accumulation of melt during the growing and active phases. Data from a
142   deep crustal section show a similar pattern (Figure 3c).[16] We suggest this pattern is another
143   characteristic product of reactive melt flow in crustal mush reservoirs. Reactive melt flow at low
144   melt fraction, rather than fractional crystallisation at high melt fraction, is the dominant mechanism
145   controlling magma storage, accumulation and chemical differentiation in the continental crust (Fig.
146   4).

147

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224
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231
232 **Author Contribution Statement**
233 MDJ wrote the code and produced the numerical results. JB prepared the phase equilibria model
234 and calibrated this to experimental data. RSJS provided information on context and background for
235 the study. All authors jointly designed the numerical experiments presented and drafted the
236 manuscript text. MDJ prepared the figures.
237
238 **Author Information**

239 Reprints and permissions information is available at www.nature.com/reprints. The authors have no
240 competing financial and/or non-financial interests to declare in relation to the work described.
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242 **Main Figure Legends**
243
244 **Figure 1 | Snapshots showing temperature, melt fraction, bulk composition and incompatible
245 trace element concentration as a function of depth during the growing and active phases of the
246 reservoir** after **a**, 0.97Ma following the onset of sill intrusions and **b**, 1.39Ma. Snapshots taken from
247 Supplementary Video 1. At early times (not shown; see Extended Data Fig. 3a), during the
248 incubation phase, individual sills cool rapidly. At later times **a**, during the growing phase, a
249 persistent mush reservoir forms but the melt fraction is low. Buoyant melt migrates upwards and
250 begins to accumulate at the top of the mush. During the active phase **b**, the accumulating melt
251 forms a high melt fraction layer containing mobile magma. The composition of the melt in the layer

252 is evolved and enriched in incompatible trace elements. Elsewhere in the mush, the melt fraction
253 remains low. At late times (not shown; see Extended Data Fig. 3b) during the waning phase, sill
254 intrusions cease and the mush cools and solidifies. To illustrate the key processes, intruding basalt
255 and crust are assumed in this example to have the same initial incompatible trace element
256 concentration. Shaded area in all plots denotes the vertical extent of basalt intrusions at that time.
257 Equivalent results for sill intrusions at 10km depth are shown in Extended Data Fig. 5.
258
259 **Figure 2 | Cold storage and rapid remobilization of magma.** Plot **a** shows melt fraction as a function
260 of depth at three different snapshots in time (1.346Ma and 4ka before and after). Reactive flow of
261 buoyant melt produces a high melt fraction layer that migrates upwards. Plot **b** shows temperature
262 and melt fraction as a function of time at a depth of 18.2km, close to the top of the reservoir. Similar
263 results are obtained over the depth range 18-18.5km. Early sill intrusions rapidly cool and crystallize.
264 The crystals are kept in 'cold storage' at sub-solidus temperature, but the temperature gradually
265 increases in response to sill intrusions deeper in the reservoir. Soon (<0.3ka) after the temperature
266 exceeds the solidus, the high melt fraction layer arrives at this depth (coloured arrow denotes the
267 corresponding snapshot in plot a) and the reservoir is remobilised: the melt fraction increases
268 rapidly to form a low crystallinity magma. The melt fraction increases much more rapidly and to a
269 higher value than would be possible by melting alone. Plot **c** shows temperature and melt fraction as
270 a function of time at a depth of 20km. Similar results are obtained over the depth range 18.5-
271 21.5km. Melt fraction remains low because reactive flow has left a refractory residue at this depth.
272 There is no remobilization, despite the increase in temperature. Data extracted from Supplementary
273 Video 1. Equivalent results for intrusion at 10km depth are shown in Extended Data Fig. 6.
274
275 **Figure 3 | Geochemical consequences of reactive melt flow in crustal magma reservoirs.** Plot **a**
276 shows SiO_2 content of low crystallinity (crystal fraction <30%) magmas. Solid curves show bulk
277 magma composition (melt+crystals); dashed curves show melt composition alone. The peak at low
278 SiO_2 corresponds to magma within the intruding sills; the peak at high SiO_2 corresponds to magma
279 within high melt fraction layers near the top of the reservoir. Also shown for comparison are data
280 from the Snake River Plain.[29] The bimodality is clear, although the basalt has a lower SiO_2 content
281 than modelled here. Results for different intruding sill compositions are shown in Extended Data
282 Figure 7. Plots **b** and **c** show modelled and observed neodymium concentration along a paleo-
283 vertical transect through the Upper Mafic Complex in the Ivrea-Verbano zone. LBZ denotes Lower
284 Basal Zone, IBZ denotes Intermediate Basal Zone, UBZ denotes Upper Basal Zone, MG denotes Main
285 Gabbro and DIO denotes Diorite.[16] Both modelled (b) and observed[16] (c) data show a spiky profile at
286 the base of the reservoir, depletion in the middle part of the reservoir, and enrichment at the top.
287
288 **Figure 4 | Reactive flow of buoyant melt at low melt fraction** is a critical mechanism controlling
289 magma storage, accumulation and differentiation in mid- to lower-crustal reservoirs. The middle
290 and lower parts of the reservoir comprise a thick (order km) mush layer, with low and relatively
291 uniform melt fraction, formed by early sill intrusions during the incubation and growing phases. This
292 layer is typically imaged in geophysical data. During the active phase, the upper part of the reservoir
293 comprises transient layers containing either intermediate/mafic, or felsic magma, that can feed
294 shallower intrusions or surface eruptions. The felsic magma layer is formed in response to changes
295 in local bulk composition caused by upwards reactive flow of buoyant melt through the mush. The
296 evolved melt accumulates around older antecrysts, which may have formed during the earliest sill
297 intrusions and hence long pre-date magma formation. In the schematic shown here, the felsic
298 magma hosts a mixture of old and young antecrysts. The old antecrysts were formed during early sill
299 intrusions; the young antecrysts formed during late sill intrusion at similar depth.
300

Methods

Model Formulation

To understand processes within crustal mush reservoirs, a quantitative model is required that includes three key features. First, the model must include the addition of hot magma or heat, to initially solid crust, in order to create and grow the reservoir.[22,23,31-34] Second, the model must include the relative motion of melt and crystals, to allow chemical differentiation.[10,34-37] Third, the model must operate primarily at low melt fraction, consistent with a wealth of evidence that crustal magma reservoirs are normally low melt fraction mushes rather than high melt fraction magma chambers.[1-9,38,39] At low melt fraction, a magma reservoir comprises a mush of crystals forming a solid framework with melt distributed along grain boundaries.[1,10,39-41] At higher melt fraction, the reservoir comprises a slurry of melt containing suspended crystals that can flow via fractures, faults or other pathways to be intruded at shallower depths or erupt at the surface.[1,26,39,40] The latter process is not modelled explicitly in this study.

The intrusion of magma to form sills can occur in numerous tectonic settings, providing both a source of heat and a source of magma that can differentiate to produce evolved melt.[1,13,16,17,21-23,31-33] Here, we follow earlier numerical approaches and model the repetitive intrusion of sills into the mid- to lower-crust (modelling magma reservoirs at depths over the range 10-30km), consistent with numerous contemporary magma reservoirs imaged in geophysical data, and magma reservoirs interpreted in deep crustal sections.[2-5,16,21-23,31-34,42-44] It is assumed that the magma in the sills is delivered from some deeper reservoir in the crust or upper mantle. In most of the example cases shown, the intruding magma is mantle-derived basalt, recognizing that crustal magmatism is largely driven by basalt[13] and consistent with numerous natural examples.[16,21,23,42-44] However, in a later section we also show results for a case when the intruding sills contain more evolved (intermediate) magma.

Most models of repetitive sill intrusion do not include relative motion of melt and crystals and, therefore, there is no chemical differentiation: the bulk composition of the mush reservoir remains constant.[22,23,31-33] Here, it is assumed that melt within the mush reservoir, produced by cooling and crystallisation of the intruded sills and also heating and melting of the surrounding crust, is present along grain boundaries and forms an interconnected network at low melt fraction.[10,39-41] The melt is buoyant because it is less dense than the surrounding crystals, so a pressure gradient is present which causes upwards flow of melt relative to the crystalline matrix.[10,34,39] The matrix can deform in response to melt flow.[45-47] This coupled process of melt migration and matrix deformation is termed compaction.[48] There is abundant evidence that compaction occurs in a wide variety of crustal igneous systems, and our assumptions are consistent with previous models of compaction.[10,26,34,48-52]

Melt flow along grain boundaries in a mush reservoir allows efficient exchange of heat and mass between melt and solid phases, so that in most of the mush and over most of its lifetime, the phases remain in local thermal and chemical equilibrium.[53] The local bulk composition of the mush therefore changes as the melt migrates upwards and the crystals compact downwards. To capture this, our model includes component transport and chemical reaction.[10,36,37,54] The results shown here demonstrate that reactive flow of melt is a critically important process controlling the storage, accumulation and chemical evolution of magma within the mush reservoir.

Governing equations and method of solution

The governing equations and method of solution are modified from Solano et al.[10] The enthalpy method is used to describe conservation of heat[55] and a binary eutectic phase diagram is used to describe solid and melt compositions assuming local thermodynamic equilibrium.[10,37]

In common with many previous studies, compaction is modelled using a modified version of the McKenzie formulation,[48] assuming that deformation of the matrix occurs by melt-enhanced diffusion creep.[45-47] This is reasonable in supra-solidus mush reservoirs deforming at low strain rates ($<<10^{-15}$ s^{-1}) and yields a Newtonian rheology for the mush.[56] The matrix shear viscosity is assumed constant, but the matrix bulk viscosity has a power-law relationship with melt fraction.[26,50,57] The melt is also

352 assumed to have a Newtonian rheology, which is reasonable for crystal-free melts containing a few
353 wt% water.[57,58]

354   Surface tension and interphase pressure are neglected. The compaction formulation is currently
355 being extended to include these potentially important effects, but a single, self-consistent model that
356 includes phase change has not yet been presented.[59-63] Differential stresses imposed by tectonic
357 forces, and magma chamber over-pressuring and loading[64] are also neglected, recognizing that at least
358 some grain boundary flow is essential to separate melt and crystals in a mush reservoir and buoyancy
359 is always available to drive this. Volatiles are assumed to remain in solution, so are not present as a
360 separate phase. In shallow crustal reservoirs, an exsolved volatile phase likely plays an important role
361 in controlling phase relations and melt flow, and driving magma mobilization.[65,66]

362   As outlined in Solano et al.,[10] the transport of heat, mass and components is modelled in 1-D,
363 using a continuum formulation of the governing conservation equations. Typical sill intrusions and
364 crustal mush reservoirs have high aspect ratio.[2-5,16,21,38,42-44,67,68] Given this, and the predominately
365 vertical flow of buoyant melt in the mush, a 1-D model is a reasonable starting point to determine the
366 effects of reactive melt flow on magma storage and differentiation. However, a 1-D model does not
367 admit the formation of high porosity, sub-vertical channels caused by reactive infiltration instability.[54]
368 Numerical modelling in 2-D has suggested that such channels are created during reactive melt flow in
369 the mantle,[54,69,70] but their formation and significance in crustal mush reservoirs is not yet clear.
370 Future work should investigate whether additional and important controls on flow in crustal magma
371 reservoirs are observed in 2- and 3-D models. Such models are likely to be computationally expensive.

372   The Boussinesq approximation is applied, so density differences between solid and liquid are
373 neglected except for terms involving gravity.[10,34,51] Melt fraction and porosity are synonymous in this
374 model. However, in contrast to previous models of crustal magma reservoirs, changes in local bulk
375 composition resulting from melt migration mean that the local melt fraction here cannot be simply
376 related via temperature to the melt fraction in the initial bulk composition (Extended Data Fig. 1c).
377 This is a very important aspect of our model and one that pertains in both simple chemical systems
378 (as employed here) and complex natural systems.

379   The governing equations can be expressed in dimensionless form as[10]

380
$$\frac{\partial h'}{\partial t'} = \kappa \frac{\partial^2 T'}{\partial z'^2} + Ste \frac{\partial}{\partial z'}\left((1-\varphi) w'_s\right) \tag{1}$$

381
$$\frac{\partial C}{\partial t'} = -\frac{\partial}{\partial z'}\left((1-\varphi) w'_s C_s\right) - \frac{\partial}{\partial z'}\left(\varphi w'_m C_m\right) \tag{2}$$

382
$$\frac{\partial I}{\partial t'} = -\frac{\partial}{\partial z'}\left((1-\varphi) w'_s I_s\right) - \frac{\partial}{\partial z'}\left(\varphi w'_m I_m\right) \tag{3}$$

383
$$\frac{\partial}{\partial z'}\left(\varphi^{-\beta} \frac{\partial w'_s}{\partial z'}\right) = \frac{\mu' w'_s}{\varphi^\alpha} + (1-\varphi)\Delta\rho' \tag{4}$$

384
$$\varphi w'_m = -(1-\varphi) w'_s \tag{5}$$

385 where h is enthalpy per unit mass; T is temperature; t is time; z is the vertical coordinate; w is velocity;
386 ϕ is melt fraction; C is composition, defined using the phase diagram described in the next section; I is
387 trace element concentration; $\Delta\rho$ is the density contrast between melt and crystals and μ is melt shear
388 viscosity, both discussed in the next section. Subscripts s and m denote solid and melt, respectively.
389 Primes denote the dimensionless equivalents of the dimensional variables.

390   The characteristic time- and length scales used to non-dimensionalize the equations are given
391 by[10]

$$\tau = \frac{1}{\Delta\rho_r g}\left(\frac{\mu_r \eta_0}{a^2 b}\right)^{\frac{1}{2}} \tag{6}$$

$$\delta = \left(\frac{\eta_0 a^2 b}{\mu_r}\right)^{\frac{1}{2}} \tag{7}$$

where $\Delta\rho_r$ is a reference density contrast and μ_r a reference melt shear viscosity discussed in the next section, g is the acceleration due to gravity, and the matrix viscosity is related to the melt fraction by[10,34,50]

$$\frac{4}{3}\eta + \xi = \eta_0 \varphi^{-\beta} \tag{8}$$

where η is the shear viscosity, ξ is the bulk viscosity, η_0 is a reference shear viscosity and β is an adjustable constant. The permeability of the mush k_ϕ is given by[10,34,48,51]

$$k_\varphi = a^2 b \varphi^\alpha \tag{9}$$

where a is the grain size, and b and α are adjustable constants.

Temperature and enthalpy are scaled using[10]

$$T' = \frac{T - T_S}{T_L - T_S} \tag{10}$$

$$h' = \frac{h - h_S}{h_L - h_S} \tag{11}$$

where the subscripts L and S denote liquidus and solidus respectively. The dimensionless scaling factor κ in equation (1) is given by[10]

$$\kappa = \frac{k_T \tau (T_L - T_S)}{\rho_r \delta^2 \left(c_p (T_L - T_S) + L_f\right)} \tag{12}$$

and the Stefan Number by[10]

$$Ste = \frac{L_f}{c_p (T_L - T_S) + L_f} \tag{13}$$

where k_T is the thermal diffusivity, c_p the sensible heat capacity, L_f the latent heat of fusion and ρ_r is a reference density discussed in the next section.

The initial condition is chemically homogenous crust with a constant linear geotherm T_{geo} and no melt present. Temperature, melt fraction and velocity are zero at the upper boundary (Earth surface); the lower boundary has constant T_{geo} and is set sufficiently deep that melt fraction and velocity remain zero. Equations (1)-(5) were solved numerically using finite difference methods and a code developed by the authors. Equation (1) was approximated using a forward-time-centered-space scheme; equations (2) and (3) using a modified Lax-Wendroff scheme; and equation (4) using a centered scheme. Node spacing and time-steps were chosen based on the results of convergence tests. Solutions reported here used 20-40 nodes per individual sill intrusion with time steps that were always less than the well-known CFL condition.[34,71]

421   The numerical methods and code have been validated extensively against analytical
422   solutions.[10,34,51,71] The energy conservation equation (1) is a special case of the general parabolic
423   diffusion-advection equation, while the mass conservation equations (2) and (3) are special cases of
424   the general hyperbolic flux conservative equation. Analytical solutions are available for simplified
425   forms of these general equations, and the numerical methods were tested against these. An analytical
426   solution is available for a simplified set of the compaction equations and the code was also tested
427   against this.
428
429   **Phase behaviour and composition-dependent material properties**
430   Phase change and compositions are described using a binary eutectic phase diagram that
431   approximates the behaviour of natural systems. Several previous studies of crustal igneous systems
432   have used a similar approach, which is preferable to more complex models involving, for example, the
433   thermodynamic software MELTS[72] for two reasons. First, reactive flow leads to local changes in bulk
434   composition, so the local phase equilibria must be recalculated at each location and time; this is trivial
435   using a simple phase diagram, but computationally intensive (albeit possible) using MELTS. Second, it
436   allows fundamental aspects of compositional evolution to be identified without the additional
437   complexity associated with modelling the phase behaviour of natural systems.[10,37]
438   Melt fraction is related to composition through

439
$$\varphi = \frac{C - C_s}{C_m - C_s} \tag{14}$$

440   where C is the local bulk composition. Assuming a linear release of enthalpy during melting, enthalpy
441   is related to temperature through

442
$$h = c_p T + L_f \varphi \tag{15}$$

443   Using equations (14) and (15), and the temperature-dependent liquid and solid compositions
444   determined from the phase diagram, the melt fraction is determined locally.
445   The binary eutectic phase diagram is described by a quadratic function given by[10]

446
$$C_m = \begin{cases} \dfrac{-a_2 - \sqrt{a_2^2 - 4a_1(a_3 - T)}}{2a_1} & C < e \\[2ex] \dfrac{-b_2 + \sqrt{b_2^2 - 4b_1(b_3 - T)}}{2b_1} & C > e \end{cases} \tag{16}$$

447
$$C_s = \begin{cases} 0 & C < e \\ 1 & C > e \end{cases} \tag{17}$$

448   Here, only compositions with $C < e$ were used. Values of the constants a_1 - a_3 were selected so that
449   the melting relations obtained for starting compositions chosen to represent crust and intruded basalt
450   match typical experimental data for the equilibrium melting/crystallization of metagreywackes and
451   basalt, respectively, over the pressure range 400 – 900MPa (Extended Data Figure 1a; Extended Data
452   Table 1).[12,73,74] The match is surprisingly good given the simple phase behaviour adopted.
453   It is important to recognize that the static melt fraction versus temperature relations shown in
454   Extended Data Figure 1a are specific to the chosen starting bulk compositions. They are not valid if
455   the bulk composition changes in response to reactive melt flow. The phase diagram provides a family
456   of melting curves for all bulk compositions encountered in the reservoir; we show just two in Extended
457   Data Fig. 1a. The effect of reactive melt flow in the reservoir is to decouple melt fraction and
458   temperature (Extended Data Figure 1c). High melt fraction can be found at low temperature where

459 reactive melt flow has caused the bulk composition of the mush to evolve towards the eutectic and
460 *vice-versa*. It is often assumed that high melt fraction necessitates high temperature.[6-8,22,23,31-33,75,]
461 Reactive melt flow means that this is not the case in crustal mush reservoirs.
462   We choose to relate composition C to a simple measure of differentiation, the SiO_2 content, by

463
$$S_{SiO_2} = a_5 + a_6 \tanh\left(a_7 + a_8 C\right) \tag{18}$$

464 Values of the constants a_5 - a_8 were selected to yield a variation in SiO_2 content with temperature that
465 matches melt SiO_2 content from the same experimental melting/crystallization data (Extended Data
466 Figure 1b; Extended Data Table 1). Again, the match is surprisingly good given the simple phase
467 behavior adopted.
468   Rearrangement of equations (14), (16) and (17), followed by substitution into (15), yields a cubic
469 polynomial in melt fraction, dependent on enthalpy h and bulk composition C, which can be solved
470 analytically[10]

471
$$\phi = \frac{h}{L_f} - \frac{c_p}{L_f}\left(\frac{a_3 + \left(\left(2a_1\left(\frac{C - C_s + C_s\phi}{\phi}\right) + a_2\right)^2 + a_2^2\right)}{4a_1} \right) \tag{19}$$

472   The model includes partitioning of a trace element between crystals and melt. The concentration
473 in the melt is given by

474
$$I_m = \frac{I}{K + \varphi\left(1 - K\right)} \tag{20}$$

475 and in the solid by

476
$$I_s = KI_m \tag{21}$$

477 In the cases modelled here, the intruding magma and crust have the same initial concentration of an
478 incompatible trace element. This is unlikely in nature, but allows the evolution of trace element
479 concentration in response to reactive melt flow in the mush to be more clearly observed and
480 understood. Trace element concentration does not affect the evolution of temperature or melt
481 fraction, so the other key model results remain unchanged.
482   The density of the melt and matrix, and the viscosity of the melt, both vary as a function of
483 composition. Solid and melt densities are given by

484
$$\rho_m = C\rho_{m\,min} + \left(1 - C\right)\rho_{m\,max} \tag{22a}$$

485
$$\rho_s = C\rho_{s\,min} + \left(1 - C\right)\rho_{s\,max} \tag{22b}$$

486 where the subscripts max and min denote, respectively, the most evolved and least evolved (most
487 refractory) compositions in the system. The average density of the crystals+melt mixture (mush or
488 magma) is given by

489
$$\overline{\rho} = \phi\rho_m + \left(1 - \phi\right)\rho_s \tag{23}$$

490 The dimensionless density is obtained by dividing by a reference density ρ_r chosen as the initial density
491 of the crust, and the dimensionless density contrast is obtained by dividing by a reference density

492   contrast $\Delta\rho_r$ chosen to be the difference in density between the most refractory crystals (ρ_{smax}) and
493   most evolved melt (ρ_{mmin}).
494   The logarithm of melt shear viscosity μ is linearly related to the dimensionless silica content of
495   the melt s_{SiO_2}

496
$$\mu = 10^{\left(\left(\mu_{max}-\mu_{min}\right)S_{SiO_2}+\mu_{min}\right)} \tag{24}$$

497   with

498
$$s_{SiO_2} = \frac{S_{SiO_2} - S_{SiO_2}^{min}}{S_{SiO_2}^{max} - S_{SiO_2}^{min}} \tag{25}$$

499   where S_{SiO_2} is the silica content of the melt (wt%).[58] The dimensionless melt shear viscosity is then
500   obtained by dividing by a reference viscosity μ_r chosen to be the maximum melt viscosity in the system
501   (corresponding to the most evolved composition), to yield

502
$$\mu' = 10^{\left(\log\left(\mu_{min}/\mu_{max}\right)\left(1-s_{SiO_2}\right)\right)} \tag{26}$$

503   In the illustrative models shown here, melt viscosity varies from a minimum of 1 Pa·s to a maximum
504   of 10^5 Pa·s (Extended Data Table 1) for the most mafic and felsic compositions respectively, which is
505   reasonable for melt containing a few wt% water.[58] A range of maximum melt viscosities is investigated
506   in the sensitivity analysis described below.
507
508 **Modelling of sill intrusion**
509   The governing equations do not include terms representing addition of heat and mass in response to
510   repetitive sill intrusion. Each sill intrusion is modelled numerically, using a simple approach in which
511   new nodes, populated with the properties (enthalpy, melt fraction, major element composition and
512   trace element concentration) of the magma in the sill, are added into the model at the target intrusion
513   depth.[22,32,34] The number of new nodes is chosen to yield the desired sill thickness. Pre-existing nodes
514   below the location of sill intrusion are shifted downwards to accommodate the new nodes
515   representing the sill; this approach represents, numerically, the case that intrusion of each new sill
516   causes downwards displacement of deeper crust and approximates isostatic equilibrium. Intrusion of
517   each sill is assumed to occur over a timescale that is small compared to the thermal and chemical
518   evolution of the magma reservoir and within a single time-step in the model. We note that injection
519   of magma may generate local over-pressure, fracturing and, during the growing and active phases of
520   the magma reservoir, locally disrupt the mush. Future refinements will focus on methods to better
521   couple thermal and mechanical models.
522
523 **Sill intrusion depth**
524   Previous numerical studies have modelled repetitive intrusion by over-accretion, in which each new
525   sill is intruded immediately above the previous sill; under-accretion, in which each new sill is intruded
526   immediately below the previous sill; and random intrusion of sills and dykes around a fixed depth.[21-
527   23,32-34] The approach used here to link sill intrusion depth to the state of the mush reservoir at the time
528   of intrusion yields variations in intrusion depth that are not captured by these previous models.
529   Controls on the depth of sill intrusions include rigidity contrasts and rheology anisotropy, resulting
530   from variations in lithology and (if present) melt fraction; rotation of deviatoric stress such that the
531   minimum deviatoric stress becomes vertical; and density contrasts between the surrounding country
532   rock and intruding magma.[67,68] The initial intrusion depth is chosen here to match the depth of an
533   observed magma reservoir. Understanding why a sill should be initially emplaced at a given depth is
534   beyond the scope of the paper. Once the first sill is emplaced, the depth of subsequent intrusions is

535   controlled by the density contrast between the intruding magma and the surrounding reservoir. The
536   next sill intrusion occurs at the deepest level of the mush that has a lower bulk density (crystals + melt)
537   than the intruding magma. The top of the resulting reservoir tends to be close to the initial intrusion
538   depth.
539   Density contrasts are controlled by the local composition and/or melt fraction of the mush
540   reservoir. We use density contrasts here as a proxy for rigidity contrasts resulting from changes in
541   rock composition or mush melt fraction.[21,56,57,67,68] Density is calculated using equations (22) and (23);
542   the chosen values of density for refractory crystals ($\rho_{s\max}$) and most evolved melt ($\rho_{m\min}$) (Extended
543   Data Table 1) yield densities of c.3000kg·m^{-3} and c.2600kg·m^{-3} for solid basalt and evolved (felsic) rock
544   compositions respectively, and densities of c.2800kg·m^{-3} and c.2350kg·m^{-3} for their corresponding
545   molten counterparts. These values are consistent with measured data.[25,76,77] The initial (reference)
546   density of the solid crust is c.2850kg·m^{-3}, consistent with data for intermediate rocks.[77]
547   During the incubation phase, melt fraction falls to zero between successive sill intrusions (Extended
548   Data Figure 2), but variations in density arise in response to differentiation within each intruded sill as
549   it cools. Differentiation yields a lower density, evolved top and a higher density, more refractory base
550   (Extended Data Figure 3a and Supplementary Video 1). Similar compositional trends are observed in
551   sills now exposed at the surface.[10,78,79] The density-controlled intrusion depth of each new sill is,
552   therefore, located below the deepest evolved top of a previous, now solidified, sill intrusion.
553   During the growing and active phases of the reservoir (Extended Data Figure 2b), melt is
554   persistently present and the compositional and density variations formed during the incubation phase
555   are reduced by reactive melt flow (Figure 1 and Supplementary Video 1). Variations in density are
556   then primarily controlled by melt fraction, so the density-controlled intrusion depth of each new sill is
557   located below the deepest high melt fraction layer.
558   Field observations from deep crustal sections suggest that intrusions progressively accumulate to
559   form a mush zone.[16,21,76] At early times, when the heat content of the reservoir is still low, intrusions
560   cool without causing significant melting of the surrounding crust, leaving septa of crust interleaved
561   with the intruded sills. We model this by intruding sills at random over a range of 300m above and
562   below the intrusion depth determined by density contrast.
563   Random intrusion preserves septa of crustal rock between sill intrusions, whereas strictly density-
564   controlled intrusion does not (see also Solano et al.[10]). Varying the depth range of random intrusion
565   affects the frequency and volume of preserved septa, but does not otherwise affect significantly the
566   results obtained. Although septa between early intrusions are preserved, septa between later
567   intrusions, when the heat content of the reservoir is higher, are partially assimilated into the melt
568   phase, causing crustal contamination of the melt.[16,76,80-81]
569
570   **Validity of the model at high melt fraction**
571   The reactive flow and compaction formulation is applied in the model regardless of local melt
572   fraction. However, it is strictly valid only when the crystals form a solid framework that will expel melt
573   if it undergoes mechanical disruption or viscous deformation.[82] Estimates of the melt fraction at which
574   this framework forms vary widely (over the range c. 0.4 - 0.7) and likely depend on local shear stresses
575   and strain rates, and the crystal morphology and size distribution.[40,45,82-84] Melt fractions higher than
576   this are present in each sill immediately after intrusion and in the melt layers that form in response to
577   reactive flow. However, we argue below that the formulation captures enough of the physics to yield
578   informative results.
579   High melt fractions are present in the intruding sills over very short timescales (of order 100's years)
580   because the sills cool very rapidly, losing heat to the surrounding reservoir and/or crust (e.g. Extended
581   Data Figure 2a). Over these short timescales following each intrusion, crystal-melt separation is
582   assumed in the model to occur only by reactive flow and compaction, omitting other mechanisms of
583   crystal-melt separation;[82] moreover, it is assumed that there is no bulk flow of melt+crystals driven by
584   convection.[28,85,86] However, the modelled cooling timescale is correct, because the rate of heat loss
585   from each sill is dominated by conduction and this is described by equation (1).[71,86] Furthermore, in

586 each sill, the model captures enough crystal/melt separation to yield a more evolved top, relatively
587 enriched in incompatible trace elements, and a more refractory base, relatively depleted in
588 incompatible trace elements, consistent with observations (e.g. Figure 3).[10,78,79]
589   High melt fractions are also present in the layers that form in response to reactive melt flow (e.g.
590 Figure 1). These layers are persistently present once formed and the model again assumes crystal-
591 melt separation in each layer occurs only by reactive flow and compaction and that there is no bulk
592 flow of melt+crystals driven by convection. However, the rate of delivery of new melt into the layer
593 is controlled by reactive flow and compaction of the underlying mush where the formulation is valid.
594 Moreover, the rate of upwards movement of the layer, which affects cold storage, is controlled by the
595 rate of upwards movement of the solidus isotherm; this depends on conductive heat transfer in the
596 overlying mush and solid rock, and is captured by the formulation. Thus we argue that the model
597 captures the overall growth and upwards migration of the layers.
598   Within each high melt fraction layer, the formulation likely does not correctly capture the
599 variation in melt fraction. However, the modelled temperature in each layer is constant at the solidus;
600 melt fraction is also high and approximately constant, controlled primarily by the local bulk
601 composition (e.g. Extended Data Video 1; Figure 1). Thus, the modelled temperature and melt fraction
602 assuming reactive flow with no bulk flow of melt+crystals are similar to what would be observed for
603 the opposite end-member model of vigorous convection in which crystals are suspended and mixed
604 in the magma.[28] We argue that vigorous convection may be more likely given the results of earlier
605 studies of single sill intrusions.[28,85,86]
606
607 **Magma mobilisation**
608 Accumulation of melt creates a high melt fraction layer which, as it migrates upwards, can remobilize
609 old mush by causing a rapid increase in melt fraction. The short timescale of this process may not
610 allow for local chemical equilibrium to be maintained, so older crystals can be preserved in the
611 younger magma. Disequilibrium between melt and crystals may also give rise to resorption and
612 zonation of crystals which is not described here.
613   The model does not attempt to capture migration out of the reservoir of the high melt fraction
614 (low crystallinity) magmas in the layer. Felsic magma that accumulates at the top of the reservoir is
615 buoyant relative to the surrounding mush reservoir, so there is a pressure gradient to drive ascent to
616 higher crustal levels or eruption at the surface.[25,26] The magma in each sill also evolves during cooling
617 to become more buoyant relative to the more refractory mush, which may drive ascent of less evolved
618 magmas. Preliminary work, not reported here, suggests that removal of felsic magma accumulating
619 in a high melt fraction layer at the top of the reservoir does not affect the formation of subsequent
620 layers, so long as ongoing sill intrusions continue to supply new magma to the reservoir.
621   This preliminary work is not reported because the model does not yet include clearly defined
622 criteria for magma removal and ascent. Moreover, we note that the presence of volatile species, such
623 as H_2O, whose solubility is pressure-dependent, complicates phase relations and physical properties
624 during magma ascent, and consequently is not considered here. Further work should determine the
625 controls on mobilization and eruption of the low crystallinity magmas present in crustal mush
626 reservoirs. What is clear from the results obtained here is that the compositions of low crystallinity
627 magmas that can leave the reservoir, regardless of how or why that happens, are bimodal. In our
628 model, the melt composition evolves to the eutectic; in more chemically complex systems, melt
629 composition will evolve to other low-variance states, such as cotectics or peritectics (reaction
630 boundaries). In all cases, the effect is to buffer chemically the composition of accumulated melts, as
631 recently suggested on the basis of phase equilibrium experiments.[87]
632
633 **Magmatic systems at shallower depth**
634 The results shown in Figure 1 (and also Supplementary Video 1 and Extended Data Figure 3)
635 illustrate the key processes occurring within a crustal mush reservoir and were obtained using values

636 of the model parameters that are typical of crustal systems (Extended Data Table 1 and associated
637 references). The initial intrusion depth was chosen to allow model results to be compared against a
638 deep crustal section now exposed at the surface: the Upper Mafic Complex of the Ivrea-Verbano
639 zone, Italy.[16,21,76] The complex is interpreted to represent c.8km of basalt intruded into the crust
640 over a few Ma (i.e. at intrusion rates of order a few mm·a⁻¹). The top of the complex is interpreted
641 to have been located at a depth of c.18 km at the time of formation.

642 The model results can explain a wide range of magmatic phenomena. However, we recognize that
643 many of the magmatic systems that provide compelling evidence for these phenomena cannot be
644 approximated by a model tuned specifically to match data from the Upper Mafic Complex.   In
645 particular, systems providing evidence for cold storage and/or compositional bimodality are often
646 located at shallower levels in the crust.[6-9] Moreover, major and trace element and isotopic data for
647 these systems suggest they may be supplied by magmas of more evolved composition than basalt.[6-
648 9,88] In transcrustal magmatic systems[1] there are likely multiple zones of intrusion: primitive basalt
649 magmas may form intrusions deep in the crust that generate more evolved magmas; these magmas
650 ascend through the crust to form intrusion zones at shallower depths.

651 We test here whether similar results are obtained if the first sill is intruded at a depth of 10km,
652 rather than 18km.   Numerous magmatic systems are observed in geophysical data at similar
653 depth.[2,4,5,38] All model parameters are the same as used previously (Extended Data Table 1), except
654 that we assume the initial geotherm is appropriate for thermally mature crust where, for example, a
655 deeper magmatic zone has thermally primed the upper crust prior to the onset of shallower
656 magmatism. Previous studies have shown that this is necessary to allow upper crustal magmatic
657 systems to form without a prohibitively long incubation period or unreasonably high rate of magma
658 intrusion.[23]

659 The results obtained are qualitatively similar to those observed at 18km depth. There is an
660 incubation period, during which the melt fraction rapidly falls to zero, with compositional contrasts
661 formed by chemical differentiation within each sill intrusion prior to solidification causing the intrusion
662 depth to increase progressively (Supplementary Video 2; Extended Data Figure 4a). During the
663 growing phase, buoyant melt again migrates upwards through the mush and reactive melt flow
664 reduces, or removes, early formed compositional contrasts, so that the intrusion depth becomes
665 controlled by the locally varying melt fraction (Supplementary Video 2; Extended Data Figure 5a).

666 During the active phase, the reservoir can again produce evolved, low crystallinity magmas from
667 the high melt fraction layer that forms beneath the solidus isotherm, close to the top of the reservoir
668 (Supplementary Video 2; Extended Data Figure 5b). When intrusion of new sills ends, the reservoir
669 enters the waning phase (Supplementary Video 2; Extended Data Figure 4b) until the mush has
670 completely solidified.

671 Cold storage is again observed where upwards migrating, evolved melt rapidly accumulates around
672 older antecrysts derived from crystallisation of early sills (Extended Data Figure 6). In this shallower
673 example, melt accumulation forms a low crystallinity magma a few 100's years after the local
674 temperature exceeds the solidus, but the magma contains antecrysts formed c. 1.3Ma earlier
675 (Extended Data Figure 6). Compositional bimodality is again observed, as magmas in the high melt
676 fraction layers have evolved composition, but magmas in the sills shortly after intrusion have
677 compositions close to that of the intruded basalt (Extended Data Figure 7a). Thus, the key results are
678 consistently observed in models of shallower magmatic systems created and sustained by basaltic
679 magmatism.
680

681 **Intrusion of more evolved magma**
682 We now test whether similar results are obtained at 10km if the intruding sills contain magma of
683 intermediate (andesitic) rather than basaltic composition. All other model parameters are the same
684 as used in the previous 10km model (Extended Data Table 1). We do not model intrusion of rhyolite
685 magma because our density controlled intrusion depth model does not apply for rhyolite magma:

686 density controlled intrusion alone would suggest that rhyolite should mostly erupt. That evolved, low
687 density magmas often intrude rather than erupt has been a challenge to density driven models of
688 magma ascent and intrusion for many years.[67,68]

689   Intrusion of intermediate composition (c. 61% SiO_2) magma yields qualitatively similar behavior to
690 that observed in response to intrusion of basaltic magma. The incubation, growing, active and waning
691 phases of reservoir life are all observed and, during the active phase, a high melt fraction layer
692 containing evolved (felsic) magma overlies a thicker, low melt fraction mush (e.g. Extended Data
693 Figure 8). Older antecrysts are again rapidly remobilized by the arriving melt layer although, in this
694 case, storage is 'cool' rather than 'cold': the temperature remains above the solidus, but the melt
695 fraction remains low until the melt layer arrives. Whether crystals are kept in cold (sub-solidus) or
696 cool (supra-solidus) conditions may be difficult to determine from crystal chemistry data, requiring
697 accurate estimates of reservoir and solidus temperatures;[6-9] the key point is that the crystals are
698 stored at low (non-eruptible) melt fraction, as opposed to 'warm storage' where the magma remains
699 eruptible.[18]

700   Compositional bimodality is again observed, but here the magma compositions are either evolved
701 (felsic), reflecting melt accumulation in the upwards migrating layer, or intermediate, reflecting the
702 composition of the intruding magma (Extended Data Figure 7b). In general, we suggest that crustal
703 mush reservoirs deliver magmas with compositions that reflect either (i) low-variance states, such as
704 eutectics, cotectics or peritectics (reaction boundaries)[87] or (ii) the intruding magma that creates the
705 reservoir.
706

707 **Intrusion depth model**
708 Numerical tests show that compositionally bimodal, low crystallinity magmas are obtained regardless
709 of whether the intrusion depth is modelled using our sill intrusion depth model or simple under- or
710 over-accretion. 'Cold' (or at least 'cool') storage of crystals, in a non-eruptible state, is also observed
711 (e.g. Extended Data Figure 9a,b), except when intrusion depth is modelled using simple over-accretion.
712 Over-accretion cannot yield cold or cool storage of antecrysts formed as part of the same magmatic
713 event, as persistent sill intrusion at the top of the magma reservoir causes the melt layer to migrate
714 upwards and form in the overlying crust (e.g. Extended Data Figure 9c,d). The crystals here are rapidly
715 mobilized by the arrival of the melt layer, but the history of the crystals and their genetic relationship
716 with the magmatic event may be much more complex. However, simple over-accretion requires the
717 magma supplying each sill to pass through the mush reservoir regardless of local melt fraction,
718 rheology or density, which is inconsistent with available evidence and models.[67,68] We argue that our
719 sill intrusion model better captures the effect of the local mush state on intrusion depth.
720

721 **Sensitivity analysis**
722 Extended Data Table 1 shows that crustal magma reservoirs are described by a broad range of material
723 properties. Values of many of these are poorly constrained. A simple sensitivity analysis was used to
724 confirm that the results obtained are typical.

725   Previous work has shown that solutions to equations (1) – (5) are largely dictated by the value of
726 the dimensionless scaling factor κ.[51,71] The effect of varying the other dimensionless parameter Ste is
727 much smaller. Other studies, confirmed by additional numerical experiments conducted here, have
728 shown that, for a given depth of intrusion and initial geothermal gradient, the thermal impact of
729 intruding sills is controlled by the intrusion rate, irrespective of the model used to choose the sill
730 intrusion depth.[22,23,32,33] The chosen intrusion rate of 5mm·yr^{-1} for the example results shown here
731 corresponds to the time-averaged magma productivity in arc settings simplified to a 1D geometry.[24]
732 We now explore a range of intrusion rates around this value, consistent with estimates for different
733 crustal magmatic systems and previous studies.[21-24,32-34]

734   A simple Monte-Carlo analysis[89] shows that 90% of the calculated values of κ for typical crustal
735 parameters lie within the range $0.028 < \kappa < 2160$ ($-1.55 < \log \kappa < 3.335$; see Extended Data Figure 10a).
736 Numerical solutions were obtained for ten values of $\log \kappa$ sampled evenly over this range in log space,

737   for a range of values of the sill intrusion rate, three different intrusion depths and basalt that is
738   intruded to a maximum thickness of 20km (Extended Data Table 1). The results are summarized in
739   Extended Data Fig. 10b,c by plotting the incubation time (the time required to reach the end of the
740   incubation phase and produce a persistent mush reservoir; see Extended Data Figure 2b), the
741   activation time (the time required to produce an active reservoir with a low crystallinity felsic magma
742   layer; see Extended Data Figure 2b), the bulk composition of the mobile magmas (i.e. magmas with
743   melt fraction >0.7), and the 'cold storage time' of antecrysts at the top of the reservoir, as a function
744   of sill intrusion rate for the different intrusion depths. The 'cold storage time' is the time elapsed
745   between the last intrusion and the local melt fraction exceeding 0.7 at locations close to the top of
746   the reservoir (see Fig. 2 and Extended Data Figures 6, 8). The cold storage time reflects the likely range
747   of crystal ages in magmas that have achieved melt fractions exceeding 0.7.

748   The incubation time scales with the reciprocal of the intrusion rate squared q^{-2} (Extended Data Fig.
749   10b). The same scaling has been obtained in previous, purely thermal, models of repetitive sill
750   intrusion using a variety of intrusion depth schemes, showing that the incubation time is relatively
751   insensitive to the details of sill intrusion.[22,23,90] Varying the value of κ over the range specified has a
752   negligible impact on the incubation time, regardless of intrusion rate or depth, reflecting the relatively
753   small range of uncertainty in thermal parameters such as thermal conductivity, specific heat capacity
754   and latent heat (Extended Data Table 1).

755   The impact of varying κ on the accumulation time is more significant, especially at lower intrusion
756   rates when the accumulation time may be several million years longer than the incubation time
757   (Extended Data Fig. 10b). Longer incubation times are observed for large values of κ that correspond
758   to larger values of the melt shear viscosity and smaller values of matrix grain size, and for small values
759   of κ that correspond to large values of the matrix bulk viscosity (Extended Data Table 1). Nevertheless,
760   the accumulation time is finite so long as the incubation time is reached within the maximum intruded
761   thickness of basalt; in other words, the formation of a high melt fraction layer is inevitable, so long as
762   a persistent mush reservoir is present.

763   The composition of the high melt fraction (eruptible) magma in the reservoir is always bimodal,
764   irrespective of the value of κ, the intrusion rate or intrusion depth (Extended Data Fig. 10c). The
765   magma in the intruded sills has a composition close to that of the intruding basalt, while the magma
766   in the layer that accumulates at the top of the reservoir has an evolved (approximately eutectic)
767   composition, consistent with the results shown earlier for specific cases (Figure 3; Extended Data
768   Figure 7a).

769   The impact of varying κ on the cold storage time is more significant, as is the effect of intrusion
770   rate (Extended Data Fig. 10c). Smaller cold storage times are observed for larger values of κ that
771   correspond to larger values of the melt shear viscosity and smaller values of matrix grain size
772   (Extended Data Table 1). Smaller cold storage times are also observed for higher intrusion rates,
773   because evolution of the system as a whole occurs more rapidly. The cold storage time reflects the
774   relative timing of sill intrusion relative to remobilization. Nevertheless, the cold storage time is always
775   non-zero; in other words, some antecrysts are stored in a cold (or cool), non-eruptible state, prior to
776   remobilization by reactive flow.

777

778   **Code availability.**
779   The code (MUSHREACT) used to solve equations (1)-(5) and produce the results reported here is
780   available from the corresponding author on request. The code is platform dependent and is not
781   optimized or tested for broad distribution, but the methodology is described within the article and
782   preceding studies.[10,34]

783

784   **Data Availability Statement**
785   No original data are reported that were not created using the software code (MUSHREACT). Data
786   can be recreated using the code.

787

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929

930   **Extended Data Figure and Table Captions**

931

932   **Extended Data Figure 1 | Phase behaviour and compositions of the modelled system. a**, Static melt
933   fraction versus temperature for the modelled basalt and crust, extracted from the binary phase
934   diagram for the chosen initial bulk compositions.   Also shown are experimental equilibrium
935   melting/crystallization data for metagreywackes and basalt over the pressure range 400-
936   900MPa.[12,73,74] Triangles denote data from ref. 73; circles data from ref. 74; squares data from ref. 12.
937   Static melt fraction denotes the melt fraction obtained if there is no relative motion of melt and
938   crystals, so the bulk composition remains constant. **b**, SiO_2 content versus temperature modelled

939   here. Also shown are experimental data corresponding to those shown in (a). **c**, Melt fraction versus
940   temperature obtained from the numerical model (data extracted from Supplementary Video 1 at
941   three snapshots in time (0.97Ma, 1.39Ma and 1.66Ma after the onset of sill intrusions) corresponding
942   to Figure 1 and Extended Data Figure 3b. Reactive flow in the mush decouples temperature and melt
943   fraction: high melt fraction can be found at low temperature where reactive melt flow has caused the
944   bulk composition of the mush to evolve and *vice-versa*.

945   **Extended Data Figure 2 | Maximum melt fraction as a function of time. a,** following a single sill
946   intrusion during the incubation period, and **b**, over the life of the reservoir. In a, the sill cools rapidly,
947   with the melt fraction falling below 0.7 (i.e. the crystallinity exceeding 30%) within 63a after intrusion,
948   and the sill solidifying within 225a. The sharp decrease in melt fraction prior to full solidification is
949   physical and represents the arrival of the solidification front during crystallisation at the eutectic. In
950   b, during the *incubation phase*, maximum melt fraction spikes after each sill intrusion, but decreases
951   rapidly and falls to zero between sill intrusions. The incubation phase ends when the melt fraction
952   remains greater than zero between sill intrusions. During the *growing phase*, the maximum melt
953   fraction at the top of the mush reservoir increases in response to reactive flow of buoyant melt. Spikes
954   in melt fraction correspond to ongoing sill intrusions deeper in the reservoir. Melt fraction at the top
955   of the mush increases until, during the *active phase*, evolved, low crystallinity (<30%) magma is
956   present which is likely to rapidly leave and ascend to shallower crustal levels. New sill intrusions cease
957   and, sometime later, the melt fraction at the top of the mush also starts to decrease. Overall, the
958   reservoir is cooling. This is the *waning phase*, at the end of which the reservoir has completely
959   solidified. Data in both plots extracted from Supplementary Video 1.

960   **Extended Data Figure 3 | Snapshots showing temperature, melt fraction, bulk composition and**
961   **incompatible trace element concentration as a function of depth through a crustal section at 18km**
962   **during the incubation and waning phases of the reservoir** after **a**, 0.82Ma following the onset of sill
963   intrusions and **b**, 1.66Ma. Snapshots are taken from Supplementary Video 1. At early times **a**, during
964   the incubation phase, individual sills cool rapidly. During the growing phase (not shown here; see
965   Figure 1a), a persistent magma reservoir forms but the melt fraction is low and relatively uniform.
966   However, buoyant melt migrates upwards and begins to accumulate at the top of the reservoir. During
967   the active phase (not shown here; see Figure 1b), a high melt fraction layer forms. At late times **b**,
968   during the waning phase, sill intrusions cease and the mush cools and solidifies. Shaded area in all
969   plots denotes the vertical extent of basalt intrusion at that time. Equivalent results for intrusion at
970   10km depth are shown in Extended Data Fig. 4.

971   **Extended Data Figure 4 | Snapshots showing temperature, melt fraction, bulk composition and**
972   **incompatible trace element concentration as a function of depth through a crustal section at 10km**
973   **depth during the incubation and waning phases** after **a**, 0.82Ma following the onset of sill intrusions
974   and **b**, 1.66Ma. Snapshots are taken from Supplementary Video 2. The results are qualitatively very
975   similar to those obtained at 18km depth (Extended Data Figure 3). During the incubation phase **a**,
976   individual sills cool rapidly. During the waning phase **b**, sill intrusions cease and the mush cools and
977   solidifies. Shaded area in all plots denotes the vertical extent of basalt intrusion at that time.

978   **Extended Data Figure 5 | Snapshots showing temperature, melt fraction, bulk composition and**
979   **incompatible trace element concentration as a function of depth through a crustal section at 10km**
980   **depth during the growing and active phases** after **a**, 0.99Ma following the onset of sill intrusions and
981   **b**, 1.39Ma. Snapshots are taken from Supplementary Video 2. The results are qualitatively very similar
982   to those obtained at 18km depth (Figure 1). During the growing phase **a**, a persistent mush reservoir
983   forms but the melt fraction is low. Buoyant melt migrates upwards and begins to accumulate at the
984   top of the reservoir. During the active phase **b**, the accumulating melt forms a high melt fraction layer

985   containing mobile magma. The composition of the melt in the layer is evolved and enriched in
986   incompatible trace elements. Elsewhere in the mush, the melt fraction remains low. Shaded area in
987   all plots denotes the vertical extent of basalt intrusions at that time.

988   **Extended Data Figure 6 | Cold storage and rapid remobilization of magma in a reservoir at 10km**
989   **depth.** Results are qualitatively very similar to those obtained at 18km depth (Figure 2). Plot **a** shows
990   melt fraction as a function of depth at the first snapshot after remobilization at 10km (1.441Ma).
991   Shaded area denotes intruded basalt. Reactive flow of buoyant melt produces a high melt fraction
992   layer that migrates upwards. Plot **b** shows temperature and melt fraction as a function of time at a
993   depth of 10km. Similar results are obtained over the depth range 10-10.5km. Early sill intrusions
994   rapidly cool and crystallize. The crystals are kept in 'cold storage' at sub-solidus temperature, but the
995   temperature gradually increases in response to sill intrusions deeper in the reservoir. Soon (<0.3ka)
996   after the temperature exceeds the solidus, the high melt fraction layer arrives at this depth and the
997   reservoir is remobilized: the melt fraction increases rapidly to form a low crystallinity magma. The
998   melt fraction increases much more rapidly and to a higher value than would be possible by melting
999   alone. Plot **c** shows temperature and melt fraction as a function of time at a depth of 12km. Similar
1000 results are obtained over the depth range 10.5-15km. Melt fraction remains low because reactive flow
1001 has left a refractory residue at this depth. There is no remobilization, despite the increase in
1002 temperature. Data extracted from Supplementary Video 2.

1003 **Extended Data Figure 7 | Geochemical consequences of reactive melt flow in crustal magma**
1004 **reservoirs at 10km depth** created by intrusion of (**a**) mafic sills and (**b**) intermediate sills. Both plots
1005 show SiO_2 content of low crystallinity (crystal fraction <30%) magmas. Solid curves show bulk magma
1006 composition (melt plus crystals); dashed curves show melt composition alone. The peak at low SiO_2
1007 corresponds to magma within the intruding sills; the peak at high SiO_2 corresponds to magma within
1008 high melt fraction layers near the top of the reservoir. In plot (a), measured data from the Snake River
1009 Plain are shown for comparison;[29] the bimodality is clear although the basalt has a lower SiO_2 content
1010 than modelled here. Bimodal compositions correspond to (1) the magma intruded into the reservoir,
1011 and (2) the most evolved composition obtained by differentiation.

1012 **Extended Data Figure 8 | Cool storage and rapid remobilization of magma in a reservoir created by**
1013 **intrusion of intermediate magma at 10km depth.** Results are qualitatively similar to those obtained
1014 by intruding basalt magma. Plot **a** shows melt fraction as a function of depth at the first snapshot
1015 after remobilization at a depth of 11.4km (1.28Ma). Reactive flow of evolved, buoyant melt produces
1016 a high melt fraction layer that migrates upwards. Plot **b** shows temperature and melt fraction as a
1017 function of time at a depth of 11.4km. Early sill intrusions rapidly cool and crystallize. The crystals are
1018 kept in 'cool storage' at near-solidus temperature. At 1.28Ma, the high melt fraction layer arrives at
1019 this depth and the reservoir is remobilized: the melt fraction increases rapidly to form a low
1020 crystallinity magma. The melt fraction increases much more rapidly and to a higher value than would
1021 be possible by melting alone. Melt fraction deeper in the reservoir remains low because reactive flow
1022 has left a refractory residue at this depth.

1023 **Extended Data Figure 9 | Consequences of emplacement during (a, b) under-accretion and (c, d)**
1024 **over-accretion.** During under-accretion, plot **a** shows melt fraction as a function of depth at the first
1025 snapshot after remobilization at a depth of 22km (1.02Ma). Reactive flow of evolved, buoyant melt
1026 produces a high melt fraction layer that migrates upwards. Plot **b** shows temperature and melt
1027 fraction as a function of time at a depth of 22km. Similar results are obtained over the depth range
1028 22-22.5km. Under-accretion causes the sill intrusion depth to progressively increase from 18km; in
1029 this case, an intrusion at 22km occurs at 0.75Ma that rapidly cools and crystallises. The crystals are
1030 kept in 'cool storage' at close-to-solidus temperature. At 1.02Ma the high melt fraction layer arrives

1031 at this depth and the reservoir is remobilized. During over-accretion, plot **c** shows melt fraction as a
1032 function of depth at a snapshot in time (1.53Ma). In this case, the high melt fraction layer has migrated
1033 into the overlying country rock. Plot **d** shows temperature and melt fraction as a function of time at a
1034 depth of 17.5km, close to the top of the active magma reservoir. Similar results are obtained over the
1035 depth range 17.5-18km. Crystals in the magma are sourced from the country rock and may be
1036 genetically unrelated to the melt. There is no cold storage of crystals brought into the reservoir by
1037 basaltic sill intrusions, as intrusion occurs deeper in the reservoir. In plots a and c, the shaded area
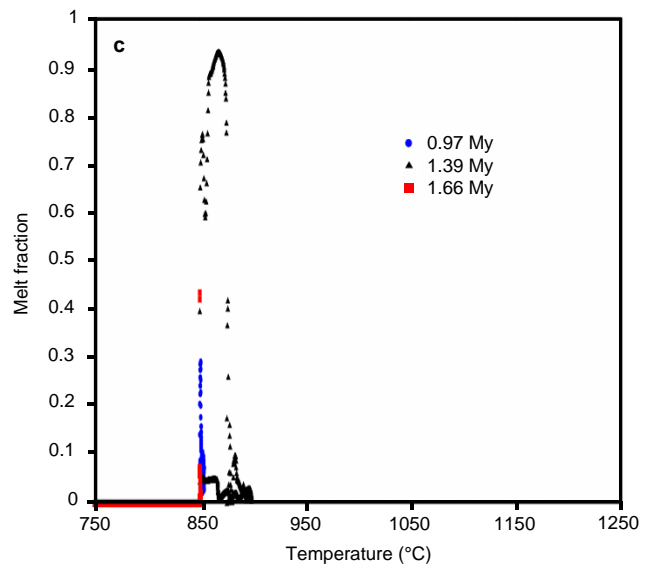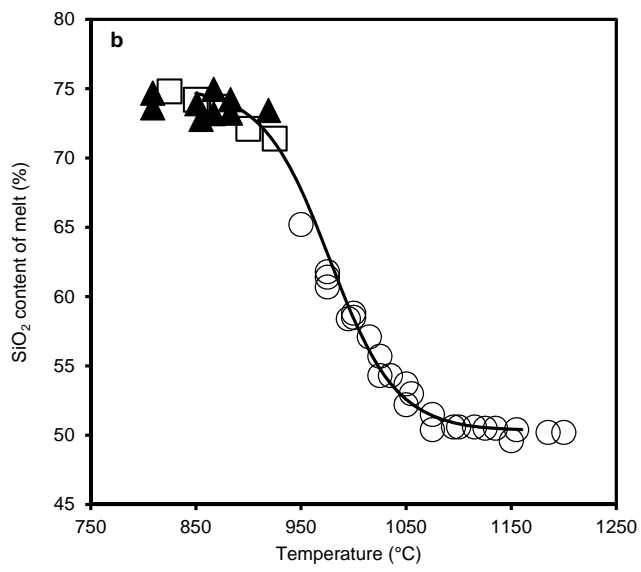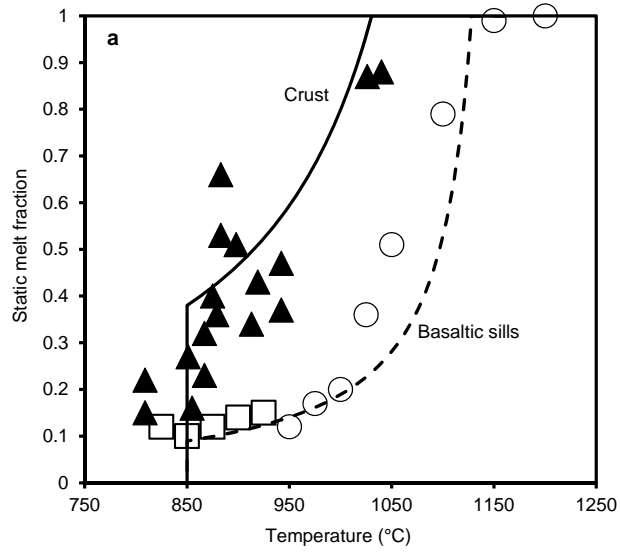1038 denotes intruded basalt.

1039 **Extended Data Figure 10 | Sensitivity analysis.** Plot (a) is a frequency plot showing values of the
1040 dimensionless scaling factor κ calculated using equation (12). Values of the input values were varied
1041 uniformly over the range given in Extended Data Table 1 in a simple Monte-Carlo analysis.[89] Plot (b)
1042 shows incubation and activation time; plot (c) shows cold storage time and eruptible magma
1043 composition. Error bars and shaded regions in (b) and (c) denote the effect of varying the
1044 dimensionless scaling factor κ over the range 0.028 < κ < 2160. Error bars on the incubation time are
1045 within the symbol size. Dashed lines denote fit to the incubation time of the form q^{-2} where q is the
1046 intrusion rate. Colours in (b) and (c) denote different initial emplacement depths of 10km, 18km and
1047 30km. Models were run for a maximum 20km of intruded basalt.

1048 **Extended Data Table 1 | Parameters used in the numerical experiments.** Values used to produce the
1049 results shown in all figures except Extended Data Figure 10. A steeper geotherm suitable for thermally
1050 mature crust[23] was assumed for the results shown in Extended Data Figures 4-8 which have intrusion
1051 at 10km depth. The range of values for the sensitivity analysis was used to calculate the range of
1052 values of the dimensionless scaling factor κ shown in Extended Data Figure 10a and produce the
1053 associated numerical modelling results shown in Extended Data Figure 10b,c. Data sources are
1054 indicated.

1055

1056

a

Crust

Basaltic sills

Static melt fraction

Temperature (°C)

b

SiO$_2$ content of melt (%)

Temperature (°C)

c

Melt fraction

- 0.97 My
- 1.39 My
- 1.66 My

Temperature (°C)

a

Melt fraction (y-axis, 0 to 1)
Time (yr) (x-axis, 0 to 250)

Arrival of solidification front at eutectic composition

b

Spikes in melt fraction correspond to sill intrusions

Persistent high melt fractions between sill intrusions correspond to the high melt fraction layer at top of mush reservoir

Crystallinity of felsic magma during active phase is c. 10%

Magma

Mush

Final sill intrusion

Melt fraction (y-axis, 0 to 1)
Time (Myr) (x-axis, 0 to 1.8)

Incubation phase

Growing phase

Active phase

Waning phase

Incubation time

Activation time

a

Location of most recent sill intrusion →

No melt present except for a few ka after each sill intrusion

Spiky trace element signature reflecting segregation in each sill intrusion

Intruding basalt composition

Initial crust composition

Initial trace element concentration

Depth (km)

Temperature (°C)

Melt fraction

SiO$_2$ (%)

Incompatible trace-element concentration (p.p.m.)

b

Temperature buffered at the solidus

Solidus

Low and approximately uniform melt fraction

Enrichment

Depletion

Spiky trace element signature preserved in deeper reservoir

← Septa of preserved crust

Depth (km)

Temperature (°C)

Melt fraction

SiO$_2$ (%)

Incompatible trace-element concentration (p.p.m.)

a

Location of most recent sill intrusion →

No melt present except for a few ka after each sill intrusion

Intruding basalt composition

Initial crust composition

Spiky trace element signature reflecting segregation in each sill intrusion

Initial trace element concentration

Depth (km)

Temperature (°C)

Melt fraction

SiO$_2$ (%)

Incompatible trace-element concentration (p.p.m.)

b

Temperature buffered at the solidus

Solidus

Low and approximately uniform melt fraction

Enrichment

Depletion

Spiky trace element signature preserved in deeper reservoir

Septa of preserved crust ←

Depth (km)

Temperature (°C)

Melt fraction

SiO$_2$ (%)

Incompatible trace-element concentration (p.p.m.)

a

Buoyant melt migrates upwards and reacts with compacting crystals

Initial crust composition

Intruding basalt composition

Initial trace element concentration

Solidus

b

Maximum temperature

High melt fraction layers of evolved magma

Low melt fraction 'mush'

Evolved bulk composition in response to upwards reactive melt flow

Spiky major/trace element signature in deeper reservoir

Temperature (°C)

Melt fraction

SiO$_2$ (%)

Incompatible trace-element concentration (p.p.m.)

Depth (km)

b

Sill intrusion at this depth
Temperature and melt fraction
rapidly decrease
after intrusion

Remobilization by reactive melt
flow from deeper reservoir

Melt fraction

Cold (sub-solidus) crystal storage

Solidus

Temperature
(oscillations show
thermal response to sill
intrusions deeper in
reservoir)

Temperature (°C)

Melt fraction

Time (Myr)

1.441 Myr

a

Depth (km)

Buoyant melt
migrates upwards

Melt fraction

c

Solidus

Temperature
(oscillations show
thermal response to sill
intrusions shallower in
reservoir)

Melt fraction

Temperature (°C)

Melt fraction

Time (Myr)

b

Sill intrusion at this depth

Cool (near-solidus) crystal storage

Remobilization by reactive melt flow from deeper reservoir

Melt fraction

Temperature (oscillations show thermal response to sill intrusions elsewhere in reservoir)

Solidus

Temperature (°C)

Time (Myr)

Melt fraction

a

Buoyant melt migrates upwards

Depth (km)

Melt fraction

b Sill intrusion at this depth →

Remobilization by reactive melt flow from deeper reservoir

Cool crystal storage

Melt fraction

Solidus

Temperature

a Buoyant melt migrates upwards

c Buoyant melt migrates upwards

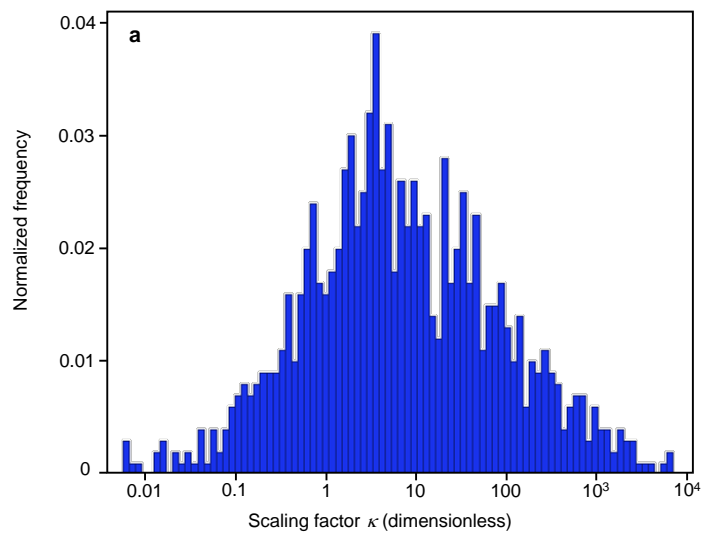**d** Melt layer created by reactive melt flow from deeper reservoir

Melt fraction

Solidus

Temperature

Symbol	Description and sources	Example case	Sensitivity analysis	Units
k_T	thermal conductivity[22,23,33,51]	3	1 - 3	$W \cdot {}^{\circ}C^{-1} \cdot m^{-1}$
c_p	specific heat capacity[51]	1100	1,020 - 1,220	$J \cdot kg^{-1} \cdot {}^{\circ}C^{-1}$
L_f	latent heat[51]	550000	400,000 - 600,000	$J \cdot kg^{-1}$
$T_L - T_S$	liquidus-solidus interval[12,73,74]	310	310	${}^{\circ}C$
T_S	solidus[12,73,74]	850	850	${}^{\circ}C$
T_{geo}	initial geotherm[21-23,51]	20	20, 40	${}^{\circ}C \cdot km^{-1}$
a	matrix grain radius[51]	2.75×10^{-3}	5×10^{-4} - 5×10^{-3}	m
α	permeability exponent[51]	3	3	None
β	bulk viscosity exponent[51]	0.5	0.5	None
b	permeability constant[51]	1/125	1/2500 - 1/50	None
μ_{max}	shear viscosity of most evolved melt[58]	10^5	$10^4 - 10^6$	Pa·s
μ_{min}	shear viscosity of least evolved melt[58]	1	1	Pa·s
η_r	reference matrix shear viscosity[26,50,51,57]	10^{15}	$10^{14} - 10^{17}$	Pa·s
q	sill intrusion rate[21-24,32-34]	5	1 - 20	$mm \cdot yr^{-1}$
z_s	sill thickness[21-23,42]	100	50-200	m
a_1, a_2, a_3	phase behavior parameters	50, -360, 1433.15	50, -360, 1433.15	${}^{\circ}C$
a_4, a_5, a_6, a_7, a_8	silica content modelling parameters	62.7, 12.38,-0.0158, 15.44	62.7, 12.38, -0.0158, 15.44	-
ρ_r	reference density[77]	2850	2850	$kg \cdot m^{-3}$
ρ_{smin}	density of most evolved solid composition[25,76,77]	3000	3000	$kg \cdot m^{-3}$
ρ_{smax}	density of least evolved solid composition[25,76,77]	2600	2600	$kg \cdot m^{-3}$
ρ_{mmin}	density of most evolved melt composition[26,76,77]	2880	2880	$kg \cdot m^{-3}$
ρ_{mmax}	density of least evolved melt composition[25,76,77]	2350	2350	$kg \cdot m^{-3}$
$S_{SiO_2}^{max}$	SiO_2 of most evolved composition[12,73,74]	75	75	%
$S_{SiO_2}^{min}$	SiO_2 of least evolved composition[12,73,74]	50	50	%
K	Trace element Nernst partition coefficient[16]	0.08	0.08	-