# Hot climate inhibits volcanism on Venus: Constraints from rock deformation experiments and argon isotope geochemistry

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## 11 Abstract

12 The disparate evolution of sibling planets Earth and Venus has left them markedly different. Venus' hot (460 °C) surface is dry and has a hypsometry with a very low standard deviation, whereas 13 Earth's average temperature is 4 °C and the surface is wet and has a pronounced bimodal 14 15 hypsometry. Counterintuitively, despite the hot Venusian climate, the rate of intraplate volcano formation is an order of magnitude lower than that of Earth. Here we compile and analyse rock 16 17 deformation and atmospheric argon isotope data to offer an explanation for the relative contrast in 18 volcanic flux between Earth and Venus. By collating high-temperature, high-pressure rock 19 deformation data for basalt, we provide a failure mechanism map to assess the depth of the brittle-20 ductile transition (BDT). These data suggest that the Venusian BDT likely exists between 2–12 km 21 depth (for a range of thermal gradients), in stark contrast to the BDT for Earth, which we find to be at a depth of ~25-27 km using the same method. The implications for planetary evolution are 22 23 twofold. First, downflexing and sagging will result in the sinking of high-elevation structures, due 24 to the low flexural rigidity of the predominantly ductile Venusian crust, offering an explanation for

25 the curious coronae features on the Venusian surface. Second, magma delivery to the surface-the most efficient mechanism for which is flow along fractures (dykes; i.e., brittle deformation)-will 26 be inhibited on Venus. Instead, we infer that magmas must stall and pond in the ductile Venusian 27 crust. If true, a greater proportion of magmatism on Venus should result in intrusion rather than 28 29 extrusion, relative to Earth. This predicted lower volcanic flux on Venus, relative to Earth, is supported by atmospheric argon isotope data: we argue here that the anomalously unradiogenic 30 present-day atmospheric <sup>40</sup>Ar/<sup>36</sup>Ar ratio for Venus (compared with Earth) must reflect major 31 differences in <sup>40</sup>Ar degassing, primarily driven by volcanism. Indeed, these argon data suggest that 32 the volcanic flux on Venus has been three times lower than that on Earth over its 4.56 billion year 33 34 history. We conclude that Venus' hot climate inhibits volcanism.

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#### 36 **1 Introduction**

37 The present-day differences in the expression and intensity of volcanism on the telluric planets serves as a testament to the dynamic nature of planetary evolution (Wilson, 2009). For example, 38 Earth and Venus are colloquially referred to as sibling planets because of their similar mass and 39 40 bulk composition (i.e., bulk petrology). However, their contrasting atmospheric mass and chemistry (e.g., Gaillard and Scaillet, 2014; Mikhail and Sverjensky, 2014), climate (e.g., Pollack et al., 41 1980), and geomorphology (e.g., Head and Solomon, 1981; Donahue and Russell, 1997; Basilevsky 42 43 and Head, 2003; Ghail, 2015) and volcanic character (e.g., Fegley and Prinn, 1989; Head et al., 1992; Wilson, 2009) is striking: Earth is a crucible of life, whereas Venus is a barren wasteland. 44 45 Suffice to say, then, Earth and Venus are not identical siblings. The major differences between 46 Venus and Earth are discussed in detail below.

First, the average surface temperatures are 460 and 4 °C on Venus and Earth, respectively. The Earth also has an excess in surface water of about  $1.2 \times 10^{21}$  kg compared to Venus, a difference between five and six orders of magnitude (Donahue, 1999; Lécuyer et al., 2000). The high temperature and low water content of the Venusian surface are a combined consequence of the absence of a magnetic field (Donahue and Russell, 1997), the presence of a dense atmosphere dominated by CO<sub>2</sub> (at a pressure of 9 MPa), and its proximity to the Sun (with a solar irradiance of 2611 W/m<sup>2</sup>, compared with 1366 W/m<sup>2</sup> on Earth).

54 Second, hypsometric data show that >80% of the surface elevation of Venus ranges from -1.0 to 55 +2.5 km; only  $\sim$ 2% of the surface lies >2 km above the median radius (Fig. 1) (Head and Solomon, 1981; Basilevsky and Head, 2003; Taylor and McLennan, 2009). The surface of Earth, by contrast, 56 57 has a pronounced bimodal hypsometry (i.e., it has continental rises and ocean basins; Fig. 1). The 58 fact that Venus has a hypsometry with a very low standard deviation is not easily attributable to the absence of plate tectonics on Venus, because Mars-a planet that, like Venus, operates a stagnant-59 60 lid tectonic regime (Head and Solomon, 1981; Head et al., 1992; Donahue and Russell, 1997; Basilevsky and Head, 2003)—has a surface hypsometry with a very large standard deviation (Fig. 61

62 1).

63 Third, the way in which volcanism is manifest on Earth and Venus differs substantially (e.g., Wilson and Head, 1983; Wilson, 2009). For example, while the majority (ca. 90%) of Earth's 64 65 volcanism occurs along curvilinear belts and rift-margins, which collectively define tectonic plate boundaries (Cottrell, 2015), Venus operates a stagnant-lid tectonic regime and is dominated by 66 67 features interpreted to be related to mantle plumes (e.g., Head et al., 1992). Although Venus is host 68 to volcanic features commonly observed on Earth, such as lava plains, discrete lava flows, shield 69 volcanoes, and shield fields, it is also home to enigmatic, flat landforms such as coronae (Head et al., 1992; Stofan et al., 1992; Squyres et al., 1992; McKenzie et al., 1992; Grosfils and Head, 1994; 70 71 Addington, 2001; Krassilnikov and Head, 2003; Grindrod and Hoogenboom, 2006; Robin et al., 2007; Wilson, 2009; Krassilnikov et al., 2012; Ivanov and Head, 2013). 72

An important difference between volcanism on Earth and Venus is that, by comparing intraplate 73 74 volcanic fluxes on both Earth and Venus, it is clear that Earth is the most volcanically active of the 75 two planets, possibly by an order of magnitude (Ivanov and Head, 2013). Indeed, while volcanic 76 activity on Earth is evidently abundant, evidence for ongoing, present-day volcanism on Venus is 77 comparatively sparse, although it is thought that the vast majority of the Venusian surface is 78 volcanic in origin (Head et al., 1992; Basilevsky and Head, 2003; Wilson, 2009). However, a 79 number of recent findings suggest that volcanic activity on Venus persists to the present: [1] 80 infrared radiation from three volcanic regions showed some flows to be warmer than their 81 surrounding rocks, implying that these lavas are younger than 2.5 Ma (Smrekar et al., 2010); [2] 82 sporadic atmospheric SO<sub>2</sub> fluctuations have been observed at Venus (Esposito, 1984; Marcq et al., 2011); and [3] thermal spikes have been reported at Ganiki Chasma, a rift valley proximal to Ozza 83 84 and Maat Montes (Shalygin et al., 2015). In addition, the sulfuric clouds that envelop the entire 85 planet would not persist beyond 1–50 Ma without the replenishment of SO<sub>2</sub>, the source of which is 86 presumed to be magmatic (Fegley and Prinn, 1989; Bullock and Grinspoon, 2001).

87 To emphasise the difference between volcanic activity on Earth and Venus: while Earth's oceanic

88 crust (that covers 60% of Earth's surface) has created >100,000 individual volcanoes (including 89 seamounts) in <100 Ma (e.g., Wessel et al., 2001 and references therein), Venus' entire surface has 90 produced roughly 70,000 individual volcanoes in <700 Ma (Head et al. 1992). The difference in the 91 rate of volcano production is therefore about an order of magnitude greater on Earth than on Venus. 92 We further note that, because >70% of all extrusive volcanism on Earth occurs beneath ocean 93 depths >1000 m under pressures >9 MPa, the presence coronae, a landform unique to the surface of 94 Venus, cannot simply be explained by the high Venusian atmospheric pressure (Smith, 1996). To 95 wit, Earth's ocean basins are not littered with coronae, but with well-formed stratovolcanoes (i.e., 96 seamounts).

97 The principal goal of this contribution is to explore the reasons as to why Earth hosts vastly more 98 intraplate volcanoes than Venus. To do so, we formulate a conceptual model that combines data 99 from rock deformation experiments on basalts, which inform on the mechanical behaviour of the 100 crust and therefore the depth of the brittle-ductile transition (BDT) on both planets, with 101 atmospheric noble gas isotope data from Earth and Venus, which inform on planetary volcanic flux. 102 Additionally, our model also offers an explanation as to why volcanoes on Venus are 103 morphologically distinct from those on Earth.

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#### 105 2 The deformation mode of the Terran and Venusian crusts

106 The depth of the BDT on Venus has been estimated numerous times. For example, first-order 107 morphological differences between fold and thrust belts on Earth and Venus can be explained by a 108 shallow BDT on Venus relative to Earth (Williams et al., 1994). Spacing between adjacent 109 extensional structures may match the spacing between linear bands seen in the mountains of Ishtar 110 Terra on Venus if the surficial brittle layer is no more than a few km in thickness (Solomon and 111 Head, 1984). Preservation of substantial crater topographic relief on Venus is likely the result of a thin (<10 km) brittle crust (Grimm and Solomon, 1988). Further, surface features within tesserae 112 (e.g., ribbons, long-wavelength folds, and grabens) offer a wealth of information as to the depth and 113

114 evolution of the BDT on Venus (Phillips and Hanson, 1998). For example, ribbons within tesserae 115 (Hansen and Willis, 1996) suggest a BDT as shallow as ~1 km during their formation (Hansen and 116 Willis, 1998; Ghent and Hansen, 1999; Ruiz, 2007). Of interest, long-wavelength folds and graben 117 are thought to reflect a deepening of the BDT over time (Phillips and Hanson, 1998)—but the depth 118 of the BDT during the formation of long-wavelength folds is estimated at only ~6 km depth (Brown 119 and Grimm, 1997; Ghent and Hansen, 1999). The pervasive deformation of the plateau highland 120 tesserae, the oldest preserved terrain, requires a weak, thin lithosphere (Brown and Grimm, 1999). 121 However, the presence of highland regions and large shield volcanoes (e.g., Crumpler et al., 1986; 122 Smrekar and Soloman, 1992; McGovern et al., 2014) implies localised crustal domains where the 123 BDT is deep enough to provide support for these structures. Nonetheless, these studies suggest that, 124 on average, the BDT on Venus is shallower than that on Earth.

125 We use here an experimental rock deformation approach to provide an alternate assessment for the 126 depth of the present-day BDT on Venus and Earth (see also Heap et al., 2017), which we interpret 127 here as a purely mechanical boundary between brittle and ductile behaviour. To do so, we compiled 128 experimental rock deformation data on basaltic (and diabase) samples deformed over a range of confining pressures (analogous to depth) and temperatures (Table 1). We used these data to 129 130 construct a failure mode map that highlights the pressures and temperatures at which basaltic (and 131 diabase) rocks behave either in a brittle or a ductile manner in response to applied stress. We then 132 used this map to assess the position (depth) of the BDT on Earth and Venus. We first review some 133 important considerations for our experimental approach.

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## 135 2.1 Considerations for our experimental approach

## 136 2.1.1 Composition of the Venusian crust

137 There is a dearth of *in-situ* quantitative geochemical data for Venusian surface rocks, and the 138 planet's thick CO<sub>2</sub>-dominated atmosphere makes optical geological observations from orbit or

139 Earth-based telescopes impossible. The only available *in-situ* geochemical data from Venus are the major element composition of surface rocks, measured using gamma-ray and X-ray fluorescence 140 141 spectroscopy. The three localities measured show basaltic compositions with SiO<sub>2</sub>, FeO, MnO, and 142 MgO abundances similar to mid-ocean ridge basalts on Earth (summarised in Bougher et al., 1997). 143 Furthermore, the data from Venera 13 and 14 (Fe/Mg, Mg/Mn, K/U, and U/Th ratios) suggest 144 Venus and Earth are made of the same chondritic material and have a similar internal structure, and 145 that Venusian basalts are the product of similar degrees of partial (peridotite) mantle melting as 146 those on Earth (Treiman, 2007; Hess and Head, 1990). Combined with the geomorphological data 147 of Venus from radar imagery (i.e., reflectance spectra), it appears that most of the Venusian surface 148 is volcanic in origin. This means the vast majority of the Venusian and Terran crusts are basaltic in 149 their bulk composition (Basilevsky and Head, 2003). Therefore, we consider the deformation mode 150 (i.e., brittle or ductile) of basaltic rocks collected on Earth to be analogous to the deformation mode 151 of those on Venus.

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#### 153 2.1.2 Hydration of the Venusian crust

The Venusian atmosphere is extremely arid, with 150 times less H<sub>2</sub>O compared with Earth's 154 155 exosphere (Donahue and Russell, 1997). However, the lack of water in Venus' atmosphere and on its surface does not necessarily imply a desiccated crust. We suggest that the degree of hydration for 156 157 Venusian crust and mantle (e.g., Kaula, 1990; Nimmo and McKenzie, 1996; Mackwell et al., 2008) 158 requires re-examination. Note, the degassing of water is extremely inefficient for one-plate planets 159 such as Venus or Mars. For example, it has been modelled that 90–95% of Mars' primordial water 160 reserves should be retained in the mantle following accretion (Hunten, 1993), and recent data show 161 the Martian mantle to be as 'wet' as the Terran mantle (McCubbin et al., 2012).

162 Combined, these studies conclude that substantial aqueous fluids can remain within planetary 163 interiors, irrespective of the plate tectonic regime and without correlation to the degree of surface 164 desiccation. For instance, if one were to distribute all of the water in the Earth's oceans into the

165 Venusian mantle, the water abundance (distributed in nominally anhydrous minerals) would not exceed the storage capacity of a peridotitic mantle (Bell and Rossman, 1992; Kohlstedt et al., 1996; 166 Bolfan-Casanova et al., 2000; Lécuyer et al., 2000; Hirschmann, 2006; Smyth et al., 2006). 167 168 Furthermore, the Martian surface and atmosphere are both very water poor, but we know that the 169 crust on Mars is hydrated (Carr and Head, 2010; 2015). A volatile-rich interior on Venus (or at least 170 a hydrated mantle) could result in explosive volcanism (Thornhill, 1993; Fagents and Wilson, 1995; 171 Glaze et al., 2011; Airey et al., 2015), and some workers have proposed that some morphological 172 units on the Venusian surface are pyroclastic deposits (Campbell and Rogers, 1994; McGill, 2000; 173 Grosfils et al., 2011; Ghail and Wilson, 2013). Therefore, it is difficult to definitively conclude 174 whether the crust and upper mantle on Venus is desiccated or hydrous, and only future missions to 175 Venus can resolve this question. Because of this ambiguity, we contend that the consideration of all 176 of the available experimental rock deformation data for basalt and diabase (including the ultra-dry 177 diabase data from Mackwell et al., 1998) is an effective approach to investigate the failure mode of rock within the Venusian crust. We also note that the majority of the basalts deformed in these 178 studies only contain a subordinate glass phase, if any. As a result, the impact of a glass phase, 179 180 hydrated or otherwise, should only play a very minor role in dictating the rheological behaviour of a 181 given sample (Smith et al., 2011; Violay et al., 2012; 2015).

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#### 183 2.2 Determining the depth of the brittle–ductile transitions for Earth and Venus

184 2.2.1 Essential nomenclature: brittle and ductile

Before interpreting the collated experimental rock deformation data it is important to outline some essential nomenclature. The terms 'brittle' and 'ductile' are sometimes interpreted differently across disciplines, which can cause confusion. To avoid such confusion, we define how we use these terms.

189 Here, we use 'brittle' and 'ductile' to describe the failure mode of a rock sample on the lengthscale

190 of that sample (typically between 10 and 100 mm). Brittle behaviour is characterized by localised 191 deformation, typically manifest as axial splits or shear fractures. During a deformation experiment, 192 it is typical to observe an increase in porosity of a sample as the peak stress is approached. This 193 increase in porosity is the result of the growth and formation of dilatant microcracks. Following a 194 peak stress, a brittle experiment involves a stress drop (i.e., strain softening). This stress drop marks the point at which a macroscopic (i.e., across the lengthscale of the sample) fracture is forming or 195 196 has formed-the hallmark of a brittle failure mode (see Hoek and Bieniawski, 1965; Brace et al., 197 1966; Scholz, 1968). We note that, in the case of highly porous samples, brittle deformation can be 198 associated with a net decrease in porosity. In these cases, inspection of the post-deformation sample 199 is required to verify the presence of axial splits or shear fractures.

200 We use the term ductility as per the definition of Rutter (1986), who described it as simply the 201 capacity of a material to accommodate qualitatively substantial strain without the tendency to 202 localise the flow into faults—localisation does not occur on the sample lengthscale. The concept of 203 ductility is not dependent on the mechanism of deformation (Rutter, 1986). Although brittle 204 behaviour is always the result of cracking on the microscale, ductile behaviour can be the product of 205 a number of micromechanisms. For example, the micromechanism behind low-temperature, high-206 pressure cataclastic flow (i.e., ductile behaviour) is microcracking (Menéndez et al., 1996; Wong et al., 1997). Ductile behaviour typically involves the loss of porosity. We note that ductile behaviour 207 208 can be associated with strain localization in certain circumstances: ductile behaviour in porous 209 rocks can involve the formation of compaction bands (e.g., Baud et al., 2004) or bands of collapsed pores (e.g., Heap et al., 2015). The formation of such features is also associated with small stress 210 211 drops in the mechanical data. In ambiguous cases, inspection of the post-deformation sample is required to verify the absence of axial splits or shear fractures, features synonymous with a brittle 212 213 failure mode. Mechanical behaviour for two experiments is shown in Fig. 2, a typical brittle test and 214 a typical ductile test (Violay et al., 2012; Heap et al., 2017).

217 In the context of our study, we are interested in the transition between brittle behaviour and ductile 218 behaviour as a result of viscous flow (i.e., the change in micromechanism from microcracking to viscous flow). We interpret viscous flow as non-recoverable viscoelastic deformation; this type of 219 220 deformation is referred to as 'plastic' by some authors, but this term is sometimes also used to describe non-recoverable deformation in the brittle field. Since we are interested in the change in 221 222 deformation micromechanism, we are not concerned here with low-temperature ductility driven by 223 microcracking or cataclastic pore collapse, although very few studies exist on this topic for basaltic rocks (e.g., Shimada, 1986; Shimada et al., 1989; Adelinet et al., 2013; Zhu et al., 2016). We 224 225 included all available experimental rock deformation data for basalt and diabase into our analysis 226 (Table 1), with the exception of room-temperature experiments under uniaxial conditions (e.g., Al-227 Harthi et al., 1999; Heap et al., 2009; Schaefer et al., 2015), because they are of little use for determining the BDT, and those triaxial experiments that yielded non-viscous ductile behaviour 228 229 such as cataclastic pore collapse (e.g., Shimada, 1986; Shimada et al., 1989; Adelinet et al., 2013; 230 Zhu et al., 2016).

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## 232 2.2.3 Limitations to our approach

One obvious limitation of our collation approach is that typical laboratory strain rates ( $\sim 10^{-5} \text{ s}^{-1}$ ) are 233 234 much faster than tectonic strain rates (e.g., Grimm, 1994). However, we recognise that [1] 235 experiments already classified as ductile at laboratory strain rates will remain ductile at lower (i.e., 236 natural) strain rates, and [2] lowering the strain rate at low experimental pressures and temperatures will reduce rock strength—because of the increased time available for subcritical crack growth (see 237 238 Brantut et al., 2013 for a review)—but may not promote ductile deformation per se. For example, 239 the experiments of Heap et al. (2011) showed that basalt can still fail in a brittle manner at a low laboratory strain rate of 10<sup>-9</sup> s<sup>-1</sup>. Although our approach utilises experiments conducted at high 240

strain rates, and so should be considered with this caveat in mind, our method does not assume arepresentative basalt for the Venusian crust (see section 2.1 above).

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#### 244 2.2.4 Calculating depth

245 Each published experimental datum was assigned a failure mode: brittle or ductile, defined above. Where necessary, and when possible, our definitions supersede those outlined in the studies from 246 247 which these data were collated. The effective pressure under which each experiment was performed were converted to a depth with the relation  $P = \rho \cdot g \cdot h$ , where P is lithospheric or hydrostatic 248 pressure and g is surface gravitational acceleration, taken as 9.807 and 8.87  $m/s^2$  for Earth and 249 Venus, respectively. This approach allowed us to determine the lithostatic pressure gradients for 250 Earth and Venus. The bulk rock density,  $\rho$ , was determined with the following relation (Wilson and 251 Head, 1994): 252

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$$\rho(h) = \frac{\rho_{\infty}}{\left[1 + \{V_0 - (1 - V_0)\}\exp(-\lambda\rho_{\infty}gz)\right]}$$
(1)

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where  $\rho_{\infty}$  (the density of porosity-free rock) was taken as 2900 kg/m<sup>3</sup>,  $V_0$  is the void space fraction (i.e., total porosity) at the surface (assumed here to be 0.25; see Wilson and Head, 1994), and the constant  $\lambda$  was assumed to be  $1.18 \times 10^{-8}$  Pa<sup>-1</sup> (Head and Wilson, 1992). Because of the very high atmospheric pressure of Venus, the lithostatic pressure at the surface was taken as 9 MPa. The hydrostatic pressure gradient for Earth was calculated using a constant water density of 1,000 kg/m<sup>3</sup> (yielding a pore pressure gradient of ~9.8 MPa/km). We note that the density of water does not vary considerably at the pressures and temperatures relevant for the Earth's crust.

However, the nature of the pore fluid, and therefore the pore pressure gradient, for Venus is enigmatic. The behavior of  $CO_2$  at the atmospheric pressure and temperature of Venus is that of a 265 supercritical fluid and, if one assumes that supercritical CO<sub>2</sub> is a plausible pore fluid for Venus, the density will vary with pressure and temperature. For example, the density of CO<sub>2</sub> at the surface of 266 Venus (at a pressure of 9 MPa and a temperature of 460 °C) is 65 kg/m<sup>3</sup>. CO<sub>2</sub> density increases to 267 457 kg/m<sup>3</sup> at a pressure of 100 MPa and a temperature of 600 °C. Because of the relatively broad 268 parameter space for pore fluid behaviour (and composition) on Venus, we considered three 269 270 scenarios that likely capture the range of possible pore fluid densities within the Venusian crust. In one, the pore fluid had a constant density of  $1,000 \text{ kg/m}^3$  (i.e., the same as on Earth, yielding a pore 271 pressure gradient of ~8.9 MPa/km); in the second scenario, pore fluid had a density of 500 kg/m<sup>3</sup> 272 273 and so a pore pressure gradient of ~4.4 MPa/km. In the third scenario, pore fluid density was 100 274  $kg/m^3$ , giving a pore pressure gradient of ~0.89 MPa/km. In all cases, the pore pressure at the surface was taken as 9 MPa. In our analysis, we interpreted the pressure within the crust as the 275 276 lithostatic pressure minus the pore fluid pressure.

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#### 278 2.2.5 Thermal gradients

279 Because we are discussing planetary-scale processes, we have opted to constrain the BDT on Earth 280 using an average Terran geothermal gradient of 25 °C/km and an average surface temperature of 4 281 °C. Due to the lack of heat-flux measurements on Venus, all of the published thermal gradients are inferred. Importantly, as a result of the greenhouse effect imposed by an average atmospheric 282 283 pressure of 9 MPa and a permanent cloud cover on Venus (Pollack et al., 1980), there is no 284 meaningful difference in average surface temperature across the Venusian day-night cycle (where 285 one Venusian day is equal to 116 Earth days) or with changes in latitude from the equator. In 286 addition, since Venus has a hypsometry with a very low standard deviation (Fig. 1) there is an 287 insignificant effect of altitude on the surface temperature when one considers a global average. 288 Therefore, a representative surface temperature for Venus should have a small standard deviation from the assumed average value of 460 °C. To account for the uncertainty in the Venusian thermal 289 290 gradient, we have used a selection of values from 5-40 °C/km (e.g., Sclater et al., 1980; Solomon and Head, 1982; 1984; Grimm and Solomon, 1988; Burt and Head, 1992; Turcotte, 1993; 1995;

Solomatov and Moresi, 1996; Turcotte et al., 1999; Leitner and Firneis, 2006).

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294 2.2.6 BDT estimates for Venus and Earth using experimental data

295 Once the effective pressure of each experiment was converted to a depth, these data were plotted 296 against the experimental temperature to examine the predicted depth of the present-day brittle-297 ductile transition on Earth (Fig. 3) and Venus (Fig. 4). The majority of experiments performed with 298 basaltic rock samples were conducted under pressures equating to depths from 0 km (i.e., the 299 surface) to 7 km (Shimada and Yukutake, 1982; Caristan, 1982; Bauer and Handin, 1983; Shimada, 300 1986; Duclos and Paquet, 1991; Schultz, 1993; Rocchi et al., 2004; Apuani et al., 2005; Benson et al., 2007; Ougier-Simonin et al., 2010; Heap et al., 2011, Violay et al., 2012; Adelinet et al., 2013; 301 302 Violay et al., 2015; Schaefer et al., 2015; Zhu et al., 2016); few studies were performed under 303 pressures corresponding to depths of up to 40 km (Griggs et al., 1960; Caristan, 1982; Hacker and 304 Christie, 1991; Mackwell et al., 1998; Violay et al., 2012; 2015). In all cases, ductile behaviour was not observed below temperatures of 500 °C, even under an effective pressure of 500 MPa (e.g., 305 306 Griggs et al., 1960). As expected, ductile behaviour is more commonly observed under combined 307 high pressure and high temperature. Surprisingly, ductile behaviour was observed under room 308 pressure (i.e., 0.1 MPa) at 800 °C (Figs. 3 and 4), which was likely the result of the presence of a 309 melt phase; therefore, although these samples were of a basaltic bulk composition, they may not 310 typify basaltic rocks. However, we prefer to retain all data for rocks of a basaltic composition in our 311 analysis, for two reasons: first, not all of the experimental studies offer a detailed microstructural 312 and compositional breakdown of their basaltic samples; second, we do not want to remove data based on our interpretation of what constitutes a basaltic rock typical of Venus or Earth. 313

Our analysis predicts that the BDT for the oceanic crust of Earth occurs at a depth of ~25-27 km (Fig. 3), consistent with the broad temperature-dependent (i.e., ~10–40 km depth) brittle–ductile transition predicted for a predominantly basaltic oceanic crust on Earth (Kohlstedt et al., 1995). 317 Assuming a pore fluid pressure gradient on Venus of ~8.7 MPa/km (Fig. 4a), we find that most of 318 the thermal gradients for Venus (i.e.,  $5-40^{\circ}$ C/km) pass through a zone (from ~5 to ~18 km depth) 319 characterised by both brittle and ductile deformation. The difference in failure mode over this depth 320 interval arises from differences in rock properties such as composition, crystal size and content, and porosity, as well as in factors such as strain rate (although we note that typically laboratory strain 321 rates rarely deviate from  $10^{-5}$  s<sup>-1</sup>). We interpret this depth interval on Venus as a failure mode 322 'transitional' zone. Below a depth of ~20 km, our collated experimental data predict exclusively 323 324 ductile behaviour when the thermal gradient is 15 °C/km or above for a pore pressure gradient of 325 ~8.7 MPa/km (Fig. 4a). However, this failure mode transitional zone is much shallower in the 326 (arguably more plausible) scenarios under which the Venusian pore pressure gradient is lower (Figs. 4b and 4c). The failure mode transition zone on Venus exists at a depth of ~4-14 km (Fig. 327 4b) or ~2–12 km (Fig. 4c) for pore pressure gradients of ~4.4 or ~0.89 MPa/km, respectively. 328

Based on these collated experimental data, we conclude that the BDT occurs at a substantially shallower depth on Venus than on Earth (when one considers global averages) (Figs. 3 and 4). Therefore, these data show that although much of the crust on Earth behaves in a brittle manner, the majority by volume of the Venusian crust should respond to stress in a ductile manner.

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#### **334 3 Implications of a dominantly ductile crust on Venus**

## 335 *3.1 Implications for volcano morphology*

The tallest volcanoes on Earth, Venus, and Mars are intraplate volcanoes fed by deep-seated mantle plumes (Head and Solomon, 1981; Donahue and Russell, 1997; Herrick et al., 2005; Wilson, 2009): Mauna Loa on Earth (17.2 km of relief), Maat Mons on Venus (9 km of relief; Mouginis-Mark, 2016), and Olympus Mons on Mars (21.9 km of relief; Plescia, 2004), respectively). However, shield volcanoes on Earth and Venus are dramatically different in terms of morphology: those on Venus are, on average, wider and of lower elevation than those on Earth (700 km wide and 1.5 km 342 relief vs. 120 km wide and 8 km relief, respectively) (Head and Solomon, 1981; Head et al., 1992; Herrick et al., 2005; Wilson, 2009). Because the loci of intraplate volcanism on Earth vary as 343 344 tectonic plates move across fixed mantle plumes, the maximum elevation of a volcano is therefore 345 not only supply limited, but is also constrained by the velocity of the plate (Morgan, 1971). By contrast. Venus' stagnant-lid tectonic regime enables a volcano to grow for as long as the magma 346 347 source persists. Note, although it is debateable if plumes on Earth and Venus are geometrically 348 similar (Schubert et al., 1990; Stofan et al., 1995; Smrekar and Stofan, 1997; Jellinek et al., 2002; 349 Johnson and Richards, 2003; Ernst et al., 2007; Robin et al., 2007), the large shield volcanoes 350 observed on the Venusian surface are taken as evidence for long-lived mantle plumes in the 351 Venusian interior (Head and Solomon, 1981; Head et al., 1992; Herrick et al., 2005; Wilson, 2009). 352 With all else being equal, the average relief of shield volcanoes on Venus should therefore be 353 greater than their Terran counterparts (Wilson, 2009). But, other than some rare if notable 354 exceptions (e.g., Maat and Skadi Montes), Venusian volcanoes are not higher in relief than their Terran counterparts. To explore this discrepancy we assess here three first-order variables that we 355 356 consider important in controlling the elevation of a volcanic construct: [1] surface gravity, [2] the 357 viscosity of extruded lavas, and [3] the flexural response of the lithosphere to geological loads.

358

#### 359 3.1.1 Surface gravity

360 Mars is host to the largest volcanoes in the Solar System. This is, in part, because high-relief structures are easier to build and retain on Mars because of its relatively low surface gravity (i.e., 361 3.71 m/s<sup>2</sup>, compared with 9.81 m/s<sup>2</sup> and 8.81 m/s<sup>2</sup> on Earth and Venus, respectively) (Heap et al., 362 363 2017). However, the surface gravitational acceleration on Venus is very similar to that of Earth meaning that, if all else were equal, both planets should extrude lava flows of a similar thickness 364 365 and build shield volcanoes of a similar size over a given timescale. Large basaltic flows on Earth are typically <30 m thick, and—again, because of the comparable surface gravitational 366 accelerations of Earth and Venus-the same should be true for Venus. This inference is consistent 367

with radar imaging of Venus that shows that flows rarely exceed the vertical resolution of the
Magellan topographic data (which has a height resolution of 5-50 m; e.g., Pettengill et al., 1991;
Roberts et al., 1992; Wilson, 2009). We conclude therefore that the minor difference in surface
gravity between Earth and Venus cannot explain the considerable contrast in volcano relief.

372

## 373 3.1.2 Viscosity of extruded lava flows

374 On Earth, high-viscosity lavas are better able to construct a tall volcanic edifice than low-viscosity lavas, which tend to travel much greater distances from the vent (e.g., Harris and Rowland, 2009). 375 Although the bulk composition of Earth and Venus are similar (Bougher et al., 1997), the 376 377 substantial influence of water content on the viscosity of melts (e.g., Dingwell et al., 1996) means that if Venusian melts are anhydrous (dry) then the lavas erupted onto its surface should have a 378 379 higher viscosity than their Terran counterparts. It is possible that the Venusian mantle is about an 380 order of magnitude more viscous than that of Earth, based on the assumption that Venusian melts 381 are anhydrous and derived from an anhydrous mantle (Kaula, 1990; Nimmo and McKenzie, 1996; Mackwell et al., 1998). However, and as outlined above, recent data that suggest a hydrated Martian 382 383 mantle (McCubbin et al., 2012) demand a reappraisal of the assumption that the Venusian mantle is 384 anhydrous. Indeed, the vast majority of basaltic lava flows on the Venusian surface are of a similar 385 spatial magnitude and thickness to the flows observed in basaltic large igneous provinces (LIPs) on 386 Earth (e.g., Columbia River and Deccan Traps; Wilson, 2009); this similarity, together with the similar surface gravity of Earth and Venus, implies a similar basaltic flow viscosity. We also note 387 388 that an increase in temperature results in a decrease in melt viscosity (Hess and Dingwell, 1996; 389 Giordano et al., 2008), even for anhydrous melts (Hess et al., 2001). Therefore, if Venusian lavas 390 are indeed anhydrous, the high temperature of the Venusian surface may decrease their nominal eruptive viscosity to a value closer to lavas extruded on Earth. We conclude, therefore, that the 391 392 difference in viscosity of erupted lavas cannot explain the difference in morphology between the 393 volcanoes on Earth and Venus.

394

#### 395 3.1.3 Response of the lithosphere to geological loads

396 An additional parameter that controls the height of a volcanic structure is the mechanical rigidity of 397 the basement upon which the volcano is situated (Watts, 2001). The flexural rigidity of the 398 lithosphere depends on its rheology such that a strong and brittle lithosphere is better adapted to 399 support high-elevation structures than a weak and ductile lithosphere (Watts, 2001). Indeed, a thick 400 and predominantly brittle crust has been used to explain the presence of the ultra-high-elevation 401 volcanoes on Mars (Turcotte et al., 1981; Heap et al., 2017), with the mechanical response of the 402 Martian crust even influencing the eruptive behaviour of these shield volcanoes (Byrne et al., 2013). 403 We contend that the experimental rock deformation data collated in Figs. 3 and 4 provide a simple 404 explanation as to why Venus hosts volcanoes that, although perhaps as voluminous, are wider and 405 of lower elevation than those on Earth. On a global scale, high-elevation structures cannot be 406 supported on Venus to the same extent as they are on Earth due to the dominantly ductile Venusian 407 crust. Recent analogue modelling by Byrne et al. (2013) aligns with this conclusion. This prediction 408 further suggests that the volcanic topographic highs on Venus (e.g., Maat Skadi Montes) may be 409 relatively young, because our model predicts that high-elevation structures on Venus will force the 410 underlying lithosphere to yield over geological timescales (according to the models presented by 411 Byrne et al. (2013); see also Smrekar and Solomon (1992) and Herrick et al. (2005)). Large 412 volcanoes may even evolve into corona-like structures over time, evidenced by the number of volcano-corona 'hybrids' on the Venusian surface (e.g., Atai Mons; Grindrod et al., 2006). We also 413 414 note that the downflexing of the lithosphere beneath a volcano imposes a constrictional strain upon 415 the edifice, manifest as imbricate shortening structures arrayed around its flanks (Byrne et al., 2009; 416 2013). Unfortunately, the flanks of Venusian volcanoes are not sufficiently resolved with currently 417 available data to test this hypothesis (full resolution Magellan topographic imagery has a resolution 418 of about 100 m; Herrick et al., 2005).

419

421 The dominant mode of magma migration through Earth's crust (in terms of volume) is via fractures (e.g., Gudmundsson, 2006). The experimental data collated here suggest that, on Venus, faulting 422 423 could be restricted to shallow depths (i.e., 2-12 km) (Fig. 4). Similar to Earth (Burov and Gerya, 424 2014), a rising mantle-derived melt on Venus will pond and spread laterally, underplating the crust 425 at depths greater than that of the BDT (as shown in Fig. 5 a-c for Venus). However, based on our 426 depth estimates for the BDT on Venus (Fig. 4), the mechanism by which magmas on Venus can 427 continue to migrate towards the BDT is through buoyancy-driven diapirism. Importantly, however, the lengthscale for magma migration by diapirism is considerably shorter than migration through 428 429 dykes (Rubin, 1995; Gudmundsson, 2002; Petford, 2003; Gudmundsson, 2006; 2011) and diapirs 430 will inevitably pond and create sills due to a stress-related equilibrium when the forces driving 431 ascent are equal (or less than) the forces acting against ascent (i.e., crystallisation increasing 432 viscosity) – unless more magma is added to further drive ascent via buoyancy-driven diapirism. 433 Therefore, if magma transport through the lower to middle Venusian crust is dominated by diapirism, then a lower fraction of crust-situated melt can reach the surface and erupt, relative to 434 435 Earth. Occasionally however, a sill may form that is large enough to generate enough uplift to 436 initiate faulting in the brittle crust, creating a set of concentric vertical faults (see Galgana et al., 437 2013). If the magma reaches these faults (or vice versa) then melts can propagate upwards, forming ring-dikes or arachnoids (Head et al., 1992; Donahue and Russell, 1997; Basilevsky and Head, 438 439 2003; Wilson, 2009). Should it reach the surface, this melt will erupt as lava, and we conceptually show how this can result in the formation of the curious coronae features on Venus in Fig. 5. 440

A combination of lateral flow and dyke-facilitated volcanism will cause the sill (magma chamber) to contract vertically, and this can cut off the magma supply to the surface as the collapsing brittle crust closes the fracture network. This may result in subsidence beneath the forming or formed coronae with either negative or positive relief (both of which are commonly observed on Venus: Head et al., 1992; Donahue and Russell, 1997; Basilevsky and Head, 2003; Herrick et al., 2005; 446 Wilson, 2009), which we argue is the result of variable ratios of the erupted lava to the amount of subsidence. If the magma supply from the plume to the crust is large enough and is active over 447 448 sufficient timescales, then a shallow-flanked shield volcano could form (e.g., Maat Mons), the 449 vertical growth of which is likely tempered by the inability of the predominantly ductile Venusian 450 crust to support high-elevation structures (due to its low flexural rigidity). However, if the magma 451 chamber (sill) cannot connect with the faults, because the sill has stalled below the BDT, then 452 surface eruption will not ensue. In this eventuality, grabens (fossae and lineae), fractures, scarps 453 (rupes), or troughs will form, tectonic landforms common to the Venusian surface (Head et al., 454 1992; Donahue and Russell, 1997; Basilevsky and Head, 2003; Wilson, 2009).

455 Most volcanic systems on Earth show complex magmatic plumbing with several reservoirs situated 456 at different depths. However, most primary mantle melts that reach the Earth's crust form sill-like 457 magma chambers at the base of the crust (defined as primary magma chambers) and are typically found at depths considerably greater than 10 km (Kelley and Barton, 2008; Stroncik et al., 2009; 458 459 Becerril et al., 2013). Therefore, on Earth, most shallow magma chambers are connected to a deeper 460 primary magma chamber at depths of >10 km (Hill et al., 2009; Michon et al., 2015). This magma 461 system architecture suggests that magma ponds at the crust-mantle boundary on Earth. Therefore, if 462 Venusian melts form magma chambers at similar depth, or at a similar depth with respect to the stratigraphy of the crust (i.e., the crust-mantle boundary), as predicted for shallow magma 463 464 chambers (Wilson and Head, 1994), then those chambers will be hosted below the BDT (predicted 465 to occur between 2 and 12 km on Venus: see Fig. 4a-c), restricting magma mobility to the short lengthscales typical of diapirism (Rubin, 1995; Gudmundsson, 2002; Petford, 2003; Gudmundsson, 466 2006; 2011). 467

We can therefore predict, albeit qualitatively, that a greater proportion of magmatism on Venus does not result in volcanism, but instead results in plutonism, than on Earth. Indeed, lava flow unit thickness estimates from Magellan topographic data suggest that coronae are probably underlain by large magma bodies that are not emptied during eruption (Grindrod et al., 2010). Any crustal 472 thickening in areas of high magmatic activity should thus be compensated by delamination back 473 into the mantle with or without crustal uplift (Smrekar and Stofan, 1997; Ghail, 2015). To test the 474 hypothesis that plutonism is favoured over volcanism on Venus (relative to Earth), we will now 475 compare differences in volcanic flux on Earth and Venus with the available geochemical data.

476

## 477 **4 Measuring the volcanic eruptive flux of Venus and Earth**

478 Finding a suitable metric to compare the eruptive fluxes of Venus and Earth is challenging. For 479 example, there is a large uncertainty for both the longevity and frequency of Venusian volcanism 480 due to the lack of reliable chronostratigraphic or radiogenic isotopic data for the Venusian surface 481 (Head et al., 1992; Basilevsky et al., 2003; Kreslavsky et al, 2015). However, there is evidence that 482 Venus has experienced some voluminous volcanism in the past, coined 'global resurfacing events'. 483 The model for catastrophic volcanic resurfacing is based on the relatively few (ca. 1,000) impact 484 craters, and is thought to have occurred between 300 Ma and 1 Ga (e.g., McKinnon et al., 1997). 485 Assuming a frequency of resurfacing episodes that declined with the rate of heat generation (based on K-Th-U systematics of the mantle), Kaula (1991) proposed that there could have been eight 486 487 resurfacing events over Venus' 4.56 Ga history. Volcanism on Venus appears to be mostly 488 quiescent between these resurfacing events, which are either random or occurring roughly once every 0.5 Ga (Kaula, 1991). If in fact magmatism during these largely passive periods does not 489 490 result in extrusive volcanism, then by our inference it may instead be manifest as massive magmatic 491 underplating of basaltic melts at the base of the crust and subsequent plutonism in the crust, 492 possibly followed by delamination back into the mantle (Smrekar and Stofan, 1997).

An important and poorly constrained parameter is the thermal structure of the Venusian interior. Nimmo and McKenzie (1997; 1998) cite the composition (specifically the FeO abundance) of the basaltic rocks analysed by the Venera and Vega landers to argue that the potential temperature of the Venusian mantle was similar to the Earth's during the emplacement of these rocks. Note, the FeO data used by Nimmo and McKenzie (1997; 1998) are by no means absolute or accurate (they 498 have large uncertainties), but this is the only data presently available and future missions are 499 required to provide an improved insight. Nevertheless, they do provide a quantitative model with 500 which to demonstrate the point. Since these basalts are between 300–800 Ma one must calculate the 501 mantle temperature for the present day; this is because resurfacing events would have cooled the 502 Venusian upper mantle, which would have been followed by an increase in temperature due to U-Th-K decay and thermal insulation by the crust. Nimmo and McKenzie (1997, 1998) concluded 503 that it is unlikely that the Venusian mantle increased in temperature by more than 200 °C over 800 504 505 Ma. Hence, these workers proposed an upper limit of 1500 °C for the potential temperature of the 506 present-day Venusian mantle (Nimmo and McKenzie, 1998). This temperature is below the solidus 507 for water-undersaturated peridotite (Kohlstedt et al., 1996; Hirschmann, 2006), and so melt 508 production would be restricted to adiabatic melting of thermochemical plumes rising through the 509 mantle (e.g., such as the Hawaiian plume on Earth; Morgan, 1971).

510 A key feature of the conceptual model presented here is that, all else being equal, the volcanic 511 eruptive flux of Venus should be lower than that of Earth. Since we cannot rely on estimates of 512 volcanic flux from chronostratigraphic methods, we must look elsewhere. For example, the chemistry of a planet's atmosphere is a passive recorder of surface and subsurface processes -513 514 including volcanism (e.g., Mather, 2008; Gaillard and Scaillet, 2014; Mikhail and Sverjensky, 2014). Therefore, if Venus has experienced a relatively retarded volcanic eruptive flux (relative to 515 Earth) over its geological history then this will have left a geochemical fingerprint in the chemistry 516 517 of the Venusian atmosphere. Herein therefore we focus on the stable isotopes of argon, principally  $^{36}$ Ar,  $^{38}$ Ar, and  $^{40}$ Ar, as useful tools for investigating the origin of volatiles (with  $^{38}$ Ar/ $^{36}$ Ar) and the 518 degassing history (with  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$ ) of Venus. This is because [1] there are data for the  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$  and 519 <sup>38</sup>Ar/<sup>36</sup>Ar ratios for the atmospheres of Earth, Mars, Venus, and solar wind (Porcelli and Pepin, 520 2002), and [2] <sup>36</sup>Ar and <sup>38</sup>Ar are primordial isotopes whereas <sup>40</sup>Ar is produced from the decay of 521 <sup>40</sup>K, with a half-life of 1.25 Ga, meaning that <sup>40</sup>Ar in planetary atmospheres can be used to derive 522 523 information regarding the degassing of planetary interiors (e.g., Halliday, 2013).

524

#### 525 **5** Validating the model: argon isotope data

Despite the dearth of missions into and below the Venusian atmosphere over the past 40 years, there are valuable data for the major and minor element geochemistry of the Venusian atmosphere, including argon isotope ratios. Indeed, argon isotopes have been previously used to inform on the evolution of Venus (e.g., Istomin et al., 1980; Hoffman et al., 1980a, b; Turcotte and Schubert, 1988; Kaula, 1990; 1991; Namiki and Solomon, 1998; Porcelli and Pepin, 2002; Mikhail and Sverjensky, 2014; Halliday, 2013; O'Rourke and Korenaga, 2015).

532 In December 1978, seven gas analysers (four mass spectrometers and two gas chromatographers) 533 provided in-situ measurements of the Venus atmospheric chemical and isotopic composition 534 (summarised by Hoffman et al., 1980a). The Soviet Union's Venera 11 and 12 landers (Istomin et 535 al., 1979) and the United States Pioneer Venus entry probe (Hoffman et al., 1980a) determined the 536 argon isotope composition of the lower Venusian atmosphere (below the altitude limit of isotopic homogenisation). Importantly, these two independent measurements provided a <sup>38</sup>Ar/<sup>36</sup>Ar ratio 537 within error of one another (summarised by Hoffman et al., 1980b). The similarity for the  ${}^{38}\text{Ar}/{}^{36}\text{Ar}$ 538 539 ratios for Earth and Venus is indicative of a shared source of volatile elements (Fig. 6). We consider 540 that the most surprising result of these measurements was that the ratio of radiogenic to primordial argon in the Venusian atmosphere was shown to be highly unradiogenic, with a  ${}^{40}$ Ar/ ${}^{36}$ Ar ratio of 541 542 only  $1.03 \pm 0.04$ . For comparison, most argon in the atmospheres of Earth and Mars is strongly radiogenic, with  ${}^{40}$ Ar/ ${}^{36}$ Ar ratios of 298.56 and 1900 ± 300, respectively (Fig. 6). Below, we outline 543 why atmospheric loss, Venus being a K-deficient planet, and diffusive degassing cannot explain the 544 difference between the <sup>40</sup>Ar/<sup>36</sup>Ar ratios of Earth and Venus. We then finish by proposing a solution 545 546 (that leans on the notion of a shallow BDT for Venus), where we conclude that this discrepancy can 547 be explained by a relatively low volcanic eruptive flux for Venus (compared to Earth).

548

One of the principle mechanisms leading to stable isotope fractionation of atmosphere-forming 550 elements is low-temperature atmospheric loss (i.e., hydrodynamic escape). This process induces 551 mass dependant stable isotope fractionation, and therefore preferentially removes the lighter 552 isotopes over the heavy isotopes (e.g., <sup>36</sup>Ar over <sup>38</sup>Ar, and <sup>38</sup>Ar over <sup>40</sup>Ar). This, in turn, means that 553 the <sup>38</sup>Ar/<sup>36</sup>Ar ratio would reflect substantial stable isotope fractionation if atmosphere loss to space 554 were the sole reason for the unradiogenic <sup>40</sup>Ar/<sup>36</sup>Ar ratio in the Venusian atmosphere. Note, this is 555 not the case for the Venusian and Terran datasets (Fig. 6). Hydrodynamic escape of <sup>36</sup>Ar cannot 556 explain the low  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$  ratio of the Venusian atmosphere, because the  ${}^{36}\text{Ar}/{}^{38}\text{Ar}$  data for the 557 Venusian and Terran datasets are almost identical (i.e., 5.5 vs. 5.3; see Fig. 6), and Earth and Venus 558 559 have very similar escape velocities for argon (ca. 12 and 13 km/s, respectively). Because Earth and Venus both show primordial  ${}^{36}$ Ar/ ${}^{38}$ Ar ratios, both planets appear to share identical (isotopic) source 560 materials (i.e., both are similar to their initial value recorded by solar wind: Porcelli and Pepin, 561 2002; Halliday, 2013). This in turn implies that both Earth and Venus had the same initial 562 atmospheric  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$  ratio. A conundrum thus ensues: where is the missing  ${}^{40}\text{Ar}$  in the Venusian 563 atmosphere? 564

565

#### 566 5.2 The case against Venus being a K-deficient planet

The unradiogenic  ${}^{40}$ Ar/ ${}^{36}$ Ar ratio for the Venusian atmosphere also cannot be explained by proposing Venus to be a K-deficient planet, because the average observed K/U ratio in rocks on the Venusian surface is 7,220 (akin to mid-ocean ridge basalts on Earth). Therefore, assuming an initial K/U and  ${}^{38}$ Ar/ ${}^{36}$ Ar ratio for Earth and Venus, the present-day  ${}^{40}$ Ar/ ${}^{36}$ Ar ratio of the Venusian atmosphere is not a reflection of the overall K abundance, but would therefore reflect either the flux of  ${}^{40}$ Ar diffused or degassed out of the mantle and/or crust.

The efficient transport of <sup>40</sup>Ar from the interior of a planet into its atmosphere can be, conceptually, 575 576 achieved by diffusion. The entire Venusian crust is at a temperature above the closure temperature for argon in most silicate systems (Kelley and Wartho, 2000). However, efficient (or total) diffusion 577 of <sup>40</sup>Ar through the crust cannot be proposed, because the Venusian atmosphere is strongly 578 unradiogenic (for argon). This indicates that the Venusian crust has retained considerable <sup>40</sup>Ar 579 580 produced continually over the age of the planet (4.56 Ga). The BSV must therefore be saturated in <sup>40</sup>Ar. The lack of <sup>40</sup>Ar-diffusion at high Venusian surface temperatures can be explained by the lack 581 of a chemical gradient. A buildup of <sup>40</sup>Ar in the crust above the closure temperature does not 582 necessarily mobilize the <sup>40</sup>Ar into the atmosphere. Buoyancy drives ascent, but pathways and 583 584 mobilising agents are also required (note, gravity and physical inhibition are also acting as opposing forces). The lack of <sup>40</sup>Ar transport can be explained by a system that rapidly reaches equilibrium 585 with the intergranular medium, despite diffusion coefficients great enough to model efficient 586 587 mobilisation, conceptually (Cassata et al., 2011), Furthermore, mass-transfer along the grain 588 boundary of silicates and oxides is limited to a very thin layer (ca. 1 nm; Joesten, 1991), so the bulk diffusivity should be reduced by the ratio of the thickness of the grain boundary to the diameter of 589 590 the grain (Faver and Yund, 1992). For a grain diameter of 0.1 to 1 mm, the diffusive lengthscales of 591 argon is <1.2 km in 1 Ga. Since the lengthscale is less than the likely thickness of the Venusian crust (which is most certainly >1.2 km), the nominal diffusive flux of  $^{40}$ Ar to the atmosphere is 592 effectively zero over 1 Ga (Namiki and Solomon, 1998). Therefore, the nominal diffusive flux of 593  $^{40}$ Ar to the atmosphere will be negligible over 4.5 Ga. 594

595

## 596 5.4 The case for a low volcanic eruptive flux on Venus relative to Earth

597 We propose volcanism is the main liberating agent for transporting <sup>40</sup>Ar to the Venusian 598 atmosphere. During mantle melting on Earth and Venus, <sup>40</sup>K and <sup>40</sup>Ar are mobilised in melts, 599 because they are both incompatible elements in primary mantle silicates, e.g., olivine (Chamorro et

al., 2002; Brooker et al., 2003). This degassing implies that the strongly unradiogenic low  ${}^{40}$ Ar/ ${}^{36}$ Ar 600 ratio in the Venusian atmosphere is mirrored by a higher crustal excess of <sup>40</sup>Ar than is observed for 601 the crust on Earth (which is known to contain excess <sup>40</sup>Ar: Allègre et al., 1996; Kelley, 2002). We 602 argue that most of the <sup>40</sup>Ar transported in melts from the Venusian mantle is locked in plutons and 603 stored within the Venusian crust, implying that there is a large excess of <sup>40</sup>Ar in the BSV. Our 604 contention that a dominantly ductile Venusian crust (Fig. 4a-c) inhibits volcanism but results in 605 606 abundant plutonism (relative to Earth; Fig. 7) forms a testable hypothesis: Venus should have degassed less <sup>40</sup>Ar, relative to Earth. Mars, for example, has a highly fractionated <sup>36</sup>Ar/<sup>38</sup>Ar ratio of 607 608 4.1 (Porcelli and Pepin, 2003; Halliday, 2013), which reflects a substantial low-temperature loss of 609 its atmosphere (Porcelli and Pepin, 2003; Halliday, 2013) (Fig. 6). Consequently, their present-day atmospheric <sup>40</sup>Ar/<sup>36</sup>Ar ratios will reflect their relative efficiencies in <sup>40</sup>Ar degassing. The present-610 day Venusian atmosphere has a strongly unradiogenic  ${}^{40}$ Ar/ ${}^{36}$ Ar ratio of  $1.03 \pm 0.04$ , compared with 611 612 298.56 for Earth (Kaula, 1991; Porcelli and Pepin, 2003; Halliday, 2013). However, the Venusian atmosphere also contains roughly two orders of magnitude more <sup>36</sup>Ar relative to Earth's atmosphere 613 (Porcelli and Pepin, 2003). If we correct the  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$  ratio for Venus then the  ${}^{40}\text{Ar}/{}^{36}\text{Ar}$  ratio of the 614 Venusian atmosphere would be approximately 103, meaning Earth has degassed three times more 615 <sup>40</sup>Ar than Venus. We view this implication here as a consequence of a higher rate of volcanism on 616 617 Earth than on Venus. This is because the majority of Earth's volcanism is directly related to Earth's 618 mobile-lid plate tectonic regime (Cottrell, 2015), but we argue that the high surface temperature and 619 dearth of deep crustal faults on Venus also plays an important role. Therefore plutonism, rather than 620 volcanism, is the dominant mode of magmatic activity on Venus (Fig. 7) and this is reflected in the unusually unradiogenic  ${}^{40}$ Ar/ ${}^{36}$ Ar ratio observed in the Venusian atmosphere (Fig. 6). 621

622

## 623 6 Concluding remarks

We present here an experimentally-constrained and isotopically-supported conceptual model that predicts Venus to have been less volcanically active relative to Earth by a factor of three, in terms 626 of eruptive flux. Since the volume of magma erupted cannot be directly discussed, we focus here on the degassing flux constrained by argon isotopes, which show that Earth has degassed three times 627 more <sup>40</sup>Ar than Venus. We conclude that the reduced eruptive flux on Venus, compared to Earth, is 628 629 the result of the hot Venusian climate, a factor that greatly impacts the dominant failure mode of, and therefore the method by which magma can travel up through, the Venusian crust. The higher 630 631 rate of intraplate volcanic activity on Earth is exemplified by the observation that Earth's relatively 632 young oceanic crust has seen the development of <100,000 individual volcanoes (i.e., seamounts) in 633 <100 Ma, whereas Venus has only produced ca. 70,000 individual volcanoes over a much longer time period (700-1000 Ma) – a difference of an order of magnitude. 634

635 An interrogation of high pressure, high temperature experimental rock deformation data suggests 636 that the unrelenting high temperature (460 °C) of the Venusian surface modifies the rheology of the Venusian crust such that the dominant failure mode within the Venusian crust is ductile. These data 637 highlight that the BDT on Venus could be as shallow as 2–12 km (Fig. 4), while the same method 638 639 yields a realistic estimate for the BDT on Earth of ~25-27 km (Fig. 3). The implications of a 640 dominantly ductile Venusian crust are twofold. First, the flexural rigidity of the Venusian lithosphere will be low, inhibiting the formation of high-relief volcanoes (via lithospheric flexure). 641 We further note that the low flexural rigidity of the Venusian lithosphere may not just impact 642 volcano morphology, but also the global hypsometric profile of Venus (Fig. 1). We therefore 643 644 speculate that the low standard deviation of the Venusian surface is also the consequence of its hot 645 climate. Second, magma delivery to the surface through fractures (i.e., dykes)-the dominant transport mechanism of magma to shallow crustal levels on the telluric planets (e.g., Wilson and 646 647 Head, 1994; Gudmundsson, 2006)-will be impeded on Venus. Our conceptual model therefore predicts that most magma on Venus will stall in the crust as sills, rather than be erupted at the 648 649 surface: plutonism, rather than volcanism, is the dominant mode of magmatic activity on Venus 650 (Fig. 7). Importantly, these implications are supported by the atmospheric argon isotope ratios for 651 Earth and Venus, which indicate that volcanic degassing, and therefore volcanic flux, has been three times lower on Venus than on Earth over the past 4.5 Ga (Fig. 6).

653 Our conceptual model falls short in describing, for example, the formation histories for the Venusian continents, Aphrodite Terra and Ishtar Terra (which, speculatively, could be the result of 654 isostatic rebound before the global resurfacing event, or crustal delamination of the lower 655 656 lithosphere back into the mantle, or a presently unknown mechanism; Smrekar and Stofan, 1997; Ghail, 2015). We also highlight that our conceptual model assumes various similarities between 657 658 Earth and Venus, such as similar mantle convective regimes, which may not be strictly true (e.g., Johnson and Richards, 2003; Robin et al., 2007). Nevertheless, our model offers a viable 659 explanation for the difference in volcano morphology between Earth and Venus (i.e., the presence 660 661 of coronae) and the relative quiescence of volcanism on Venus compared to Earth (i.e., the order of 662 magnitude difference in the rate of intraplate volcano formation between Earth and Venus). Furthermore, a Venusian BDT as shallow as predicted here also implies that faulting through the 663 vertical lengthscale of the crust is hindered. Therefore, the hot climate of Venus may also inhibit the 664 formation of the plate tectonic boundaries that sub-divide the crust (Foley et al., 2012; Bercovici 665 and Ricard, 2014). Our study highlights another example of the complex interplay between climate 666 and geodynamics. 667

668

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Fig. 1: Hypsography of Venus, Earth, and Mars (Head and Solomon, 1981; Basilevsky and Head,
2003; Taylor and McLennan, 2009). Dashed lines mark the mean surface elevation for each planet.



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**Fig. 2**: The mechanical behaviour of rock in compression (from Heap et al., 2017). Examples of brittle and ductile stress-strain curves for basalt deformed at a confining pressure of 300 MPa and a temperature of 650 °C (brittle test) and 850 °C (ductile test) (data from Violay et al., 2012). Inset shows cartoons depicting post-failure samples typical of brittle (throughgoing shear fracture) and ductile (distributed deformation) deformation.



Fig. 3: Failure mode map (brittle or ductile) for Earth assuming a pore pressure gradient of ~9.8
MPa/km, a surface gravity of 9.807 m/s<sup>2</sup>, an average thermal gradient of 25 °C/km, and an average
surface temperature of 4 °C. See text for details.

688



692

**Fig. 4**: Failure mode maps (brittle or ductile) for Venus assuming a surface gravity of 8.87 m/s<sup>2</sup> and an average surface temperature of 460 °C. Due to the uncertainty in the pore pressure gradient we provide three scenarios. (a) That the pore fluid has a constant density of 1000 kg/m<sup>3</sup> (i.e. the same as Earth; yielding a pore pressure gradient of ~8.9 MPa/km). (b) That the pore fluid has a constant density of 500 kg/m<sup>3</sup> (yielding a pore pressure gradient of ~4.4 MPa/km). (c) That the pore fluid has a constant density of 100 kg/m<sup>3</sup> (yielding a pore pressure gradient of ~0.89 MPa/km). Due to the

## 699 uncertainty in the thermal gradient we provide a range from 5 to 40 °C/km. See text for details.



Fig. 5: The formation of coronae on Venus. This cartoon depicts sill emplacement and growth,
followed by uplift and faulting of the crust above the brittle-ductile transition (BDT). The schematic
also shows how this only leads to volcanism after the magma chamber makes physical contact with
faults (see text for more details; not to scale). Arrows indicate directions of main differential
stresses.





Fig. 6: The atmospheric argon isotope composition of Earth, Mars, and Venus (data from Istomin et al., 1979; Hoffman et al., 1980b; Porcelli and Pepin, 2002; Mahaffy et al., 2013).



Fig. 7: Schematic illustration showing the relative differences for magma transport within the lithosphere on Earth and Venus. The cartoon shows that primary magma chambers on Venus rely on diapirism to move towards the surface, leading to stagnation and crystallisation (on average). Conversely for Earth, primary magma chambers can force dyking in the overlying (brittle) lithosphere and initiate volcanism.

715

# 716 **Table caption**

Reference	Pc (MPa)	Pp (MPa)	Peff (MPa)	<i>T</i> (°C)	$\sigma_p$ (MPa)	Failure mode	Notes
Griggs et al. 1960	500	0	500	25	1668	Brittle	Basalt
Griggs et al. 1960	500	0	500	300	1390	Brittle	Basalt
Griggs et al. 1960	500	0	500	500	1080	Brittle	Basalt
Griggs et al. 1960	500	0	500	700	-	Ductile	Basalt
Griggs et al. 1960	500	0	500	800	-	Ductile	Basalt
Caristan 1982	0	0	0	950	199	Brittle	Maryland diabase; strain rate = $10^{-3}$ s <sup>-1</sup>
Caristan 1982	0	0	0	970	223	Brittle	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Caristan 1982	0	0	0	995	193	Brittle	Maryland diabase; strain rate = $10^{-3}$ s <sup>-1</sup>
Caristan 1982	30	0	30	1000	370	Brittle	Maryland diabase; strain rate = $10^{-3}$ s <sup>-1</sup>
Caristan 1982	50	0	50	1000	440	Brittle	Maryland diabase; strain rate = $10^{-3}$ s <sup>-1</sup>
Caristan 1982	150	0	150	810	780	Brittle	Maryland diabase; strain rate = $10^{-6}$ s <sup>-1</sup>
Caristan 1982	150	0	150	970	385	Brittle	Maryland diabase; strain rate = $10^{-6}$ s <sup>-1</sup>
Caristan 1982	150	0	150	994	535	Brittle	Maryland diabase; strain rate = $10^{-3}$ s <sup>-1</sup>
Caristan 1982	150	0	150	1000	566	Brittle	Maryland diabase; strain rate = $10^{-4}$ s <sup>-1</sup>
Caristan 1982	150	0	150	1000	561	Brittle	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Caristan 1982	150	0	150	1000	573	Brittle	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Caristan 1982	350	0	350	1000	-	Ductile	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Caristan 1982	400	0	400	1000	-	Ductile	Maryland diabase; strain rate = $10^{-4}$ s <sup>-1</sup>
Caristan 1982	425	0	425	1000	-	Ductile	Maryland diabase; strain rate = $10^{-4}$ s <sup>-1</sup>
Caristan 1982	425	0	425	1000	-	Ductile	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Caristan 1982	425	0	425	1000	-	Ductile	Maryland diabase; strain rate = $10^{-6}$ s <sup>-1</sup>
Caristan 1982	450	0	450	1000	-	Ductile	Maryland diabase; strain rate = $10^{-5}$ s <sup>-1</sup>
Shimada and Yukutake 1982	57	0	57	25	400	Brittle	Yakuno basalt; Porosity = $0.07$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Shimada and Yukutake 1982	107	0	107	25	415	Brittle	Yakuno basalt; Porosity = $0.07$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Bauer et al. 1981	50	0	50	25	540	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>

Bauer et al. 1981	50	0	50	25	400	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	600	300	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08: strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	600	340	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	700	300	Brittle	Cuerbio basalt; Porosity = $0.05$ - $0.08$ ; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	940	125	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	940	200	Brittle	Cuerbio basalt; Porosity = $0.05$ - $0.08$ ; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	50	0	50	1000	100	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	100	0	100	700	465	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al.	100	0	100	900	240	Brittle	Cuerbio basalt; Porosity = $0.05$ - $0.08$ ; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al. 1981	100	0	100	950	110	Brittle	Cuerbio basalt; Porosity = $0.05$ - $0.08$ ; strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al. 1981	100	0	100	1000	180	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08: strain rate = $10^{-4}$ s <sup>-1</sup>
Bauer et al. 1981	100	50	50	820	180	Brittle	Cuerbio basalt; Porosity = $0.05$ - 0.08: strain rate = $10^{-4}$ s <sup>-1</sup>
Shimada 1986	57	0	57	25	410	Brittle	Yakuno basalt; Porosity = $0.07$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	300	399	Brittle	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	600	430	Brittle	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	700	445	Brittle	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	750	430	Brittle	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	800	-	Ductile	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	900	-	Ductile	Alkaline basalt; partially glassy; strain rate = $10^{-6}$ s <sup>-1</sup>
Duclos and Paquet 1991	0	0	0	1000	-	Ductile	Alkaline basalt; partially glassy; strain rate = $10^{-6} \text{ s}^{-1}$
Hacker and Christie 1991	1000	0	1000	675	-	Ductile	Tholeiitic basalt; partially glassy; 0.5 wt.% water added; strain rate = $10^{-4} - 10^{-7} \text{ s}^{-1}$
Hacker and Christie 1991	1000	0	1000	725	-	Ductile	Tholeiitic basalt; partially glassy; 0.5 wt.% water added; strain rate = $10^{-4} - 10^{-7} \text{ s}^{-1}$
Hacker and Christie 1991	1000	0	1000	775	-	Ductile	Tholeiitic basalt; partially glassy; 0.5 wt.% water added; strain rate = $10^{-4} - 10^{-7} \text{ s}^{-1}$
Hacker and Christie 1991	1000	0	1000	825	-	Ductile	Tholeiitic basalt; partially glassy; 0.5 wt.% water added; strain rate = $10^{-4} - 10^{-7} \text{ s}^{-1}$
Hacker and Christie 1991	1000	0	1000	875	-	Ductile	Tholeiitic basalt; partially glassy; 0.5 wt.% water added; strain rate = $10^{-4} - 10^{-7} \text{ s}^{-1}$
Schultz 1993	0	0	0	450	210	Brittle	Estimated strength value taken as 80% of the average uniaxial compressive strength for basalt; see Schultz (1993) for details
Mackwell et al. 1998	400	0	400	1000	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Mackwell et al. 1998	400	0	400	1050	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Mackwell et al. 1998	400	0	400	1050	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Mackwell et al. 1998	450	0	450	970	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Mackwell et al. 1998	450	0	450	1000	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Mackwell et al. 1998	450	0	450	1050	-	Ductile	Dehydrated Maryland and Columbia diabase; creep test; strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>

Mackwell et	500	0	500	1000	-	Ductile	Dehydrated Maryland and
al. 1998							strain rate = $10^{-5} - 10^{-7}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	300	89	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	300	104	Brittle	Etna "core" basalt; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	300	35	Brittle	Etna "crust" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	600	96	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	600	105	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	600	103	Brittle	Etna "core" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	600	181	Brittle	Etna "core" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	600	40.5	Brittle	Etna "crust" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	700	33	Brittle	Etna "crust" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	800	42	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	800	43	Brittle	Etna "core" basalt; strain rate = $10^{-4}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	800	25	Brittle	Etna "core" basalt; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	800	17	Brittle	Etna "core" basalt; strain rate = $10^{-6} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	800	20	Brittle	Etna "crust" basalt; strain rate = $10^{-4}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	900	50	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-4}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	900	38	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	900	29	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	0	0	0	900	31	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-6}$ s <sup>-1</sup>
Rocchi et al. 2004	5	0	5	25	108	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	10	0	10	25	104	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	10	0	10	300	101	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	10	0	10	300	88	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	10	0	10	600	116	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	10	0	10	916	62	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	12	0	12	25	93	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	15	0	15	25	101	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	17	0	17	25	100	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	20	0	20	25	109	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	20	0	20	300	95	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	20	0	20	300	91	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	20	0	20	600	118	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	25	112	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	25	103	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	300	105	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	300	87	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	600	104	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>
Rocchi et al. 2004	30	0	30	604	79	Brittle	Vesuvius basalt; Porosity = $0.08$ - 0.10; strain rate = $10^{-5}$ s <sup>-1</sup>

2004 2004	0	0	0	900	-	Ductile	Etna "crust" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	912	-	Ductile	Etna "core" basalt; strain rate = $10^{-5} \text{ s}^{-1}$
Rocchi et al. 2004	0	0	0	1001	-	Ductile	Vesuvius basalt; Porosity = $0.08$ - 0 10: strain rate = $10^{-5}$ s <sup>-1</sup>
Apuani et al.	4	0	4	25	98	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al.	4	0	4	25	72	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al.	4	0	4	25	67	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al.	8	0	8	25	88	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al.	8	0	8	25	99	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	12	0	12	25	104	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	12	0	12	25	109	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	16	0	16	25	54	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	16	0	16	25	62	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	16	0	16	25	87	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	16	0	16	25	94	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	20	0	20	25	56	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	20	0	20	25	109	Brittle	Vigna Vecchia basalt (Stromboli)
Apuani et al. 2005	20	0	20	25	178	Brittle	Vigna Vecchia basalt (Stromboli)
Benson et al. 2007	60	20	40	25	475	Brittle	Etna basalt; porosity = 0.04; strain rate = $10^{-6}$ s <sup>-1</sup>
Ougier- Simonin et al. 2010	15	0	15	25	370	Brittle	Seljadur basalt; porosity = 0.05; strain rate = $10^{-6}$ s <sup>-1</sup>
Heap et al. 2011	30	20	10	25	291	Brittle	Etna basalt; porosity = 0.4; strain rate = $10^{-5}$ s <sup>-1</sup>
Heap et al. 2011	50	20	30	25	287	Brittle	Etna basalt; porosity = 0.4; strain rate = $10^{-5}$ s <sup>-1</sup>
Heap et al. 2011	70	20	50	25	504	Brittle	Etna basalt; porosity = 0.4; strain rate = $10^{-5}$ s <sup>-1</sup>
Heap et al. 2011	50	20	30	25	375	Brittle	Etna basalt; porosity = 0.4; creep test: strain rate = $10^{-6}$ s <sup>-1</sup>
Heap et al. 2011	50	20	30	25	357	Brittle	Etna basalt; porosity = 0.4; creep test: strain rate = $10^{-7}$ s <sup>-1</sup>
Heap et al. 2011	50	20	30	25	329	Brittle	Etna basalt; porosity = 0.4; creep test: strain rate = $10^{-8}$ s <sup>-1</sup>
Heap et al. 2011	50	20	30	25	304	Brittle	Etna basalt; porosity = 0.4; creep test: strain rate = $10^{-9}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	400	1002	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	400	902	Brittle	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$
Violay et al. 2012	100	0	100	600	854	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	700	508	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	800	462	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	800	446	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	900	355	Brittle	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	600	749	Brittle	Aphanitic basalt; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	700	755	Brittle	Aphanitic basalt; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	800	518	Brittle	Aphanitic basalt; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	50	0	50	600	-	Ductile	Porphyritic basalt; partially glassy; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>

Violay et al. 2012	70	0	70	600	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	500	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	600	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	600	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	700	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	800	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	800	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	800	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	900	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	900	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	100	0	100	900	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	250	0	250	650	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	600	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	700	-	Ductile	Porphyritic basal; partially glassy; porosity = 0.02; strain rate = $10^{-5}$
Violay et al. 2012	300	0	300	750	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	800	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	800	-	Ductile	Aphanitic basalt; porosity = 0.02; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	850	-	Ductile	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	900	-	Ductile	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	900	-	Ductile	Porphyritic basalt; partially glassy; porosity = 0.02; strain rate = $10^{-3}$ s <sup>-1</sup>
Violay et al. 2012	300	0	300	950	-	Ductile	Aphanitic basalt; porosity = $0.02$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Adelinet et al. 2013	10	5	5	25	120	Brittle	Reykjanes basalt; porosity = $0.08$ ; strain rate = $10^{-6}$ s <sup>-1</sup>
Adelinet et al.	80	76	4	25	118	Brittle	Reykjanes basalt; porosity = $0.08$ ; strain rate = $10^{-6}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	600	877	Brittle	Aphanitic basalt; porosity = $0.03$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al.	130	30	100	650	834	Brittle	Aphanitic basalt; porosity = $0.03$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	700	792	Brittle	Aphanitic basalt; porosity = $0.03$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	750	699	Brittle	Aphanitic basalt; porosity = $0.03$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	800	717	Brittle	Aphanitic basalt; porosity = $0.03$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	900	382	Brittle	Aphanitic basalt; porosity = 0.03; strain rate = $10^{-5}$ s <sup>-1</sup>
Violay et al. 2015	130	30	100	1050	-	Ductile	Aphanitic basalt; porosity = 0.03; strain rate = $10^{-5}$ s <sup>-1</sup>

Schaefer et al. 2015	0	0	0	935	167	Brittle	Pacaya (Guatemala) basalt; porosity = 0.02; strain rate = $10^{-1}$
Schaefer et al. 2015	0	0	0	935	162	Brittle	Pacaya (Guatemala) basalt; porosity = 0.05; strain rate = $10^{-1}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	126	Brittle	Pacaya (Guatemala) basalt; porosity = 0.06; strain rate = $10^{-5}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	59	Brittle	Pacaya (Guatemala) basalt; porosity = 0.19; strain rate = $10^{-1}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	49	Brittle	Pacaya (Guatemala) basalt; porosity = 0.16; strain rate = $10^{-5}$ s <sup>-1</sup>
Schaeffer et al. 2015	0	0	0	935	93	Brittle	Pacaya (Guatemala) basalt; porosity = 0.19; strain rate = $10^{-1}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	44	Brittle	Pacaya (Guatemala) basalt; porosity = 0.19; strain rate = $10^{-5}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	75	Brittle	Pacaya (Guatemala) basalt; porosity = 0.23; strain rate = $10^{-1}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	64	Brittle	Pacaya (Guatemala) basalt; porosity = 0.21; strain rate = $10^{-5}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	28	Brittle	Pacaya (Guatemala) basalt; porosity = 0.32; strain rate = $10^{-1}$ s <sup>-1</sup>
Schaefer et al. 2015	0	0	0	935	16	Brittle	Pacaya (Guatemala) basalt; porosity = 0.31; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	20	10	10	25	281	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	20	10	10	25	240	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	20	10	10	25	221	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	20	10	10	25	327	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	30	10	20	25	329	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	30	10	20	25	361	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	40	10	30	25	399	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	50	10	40	25	403	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	60	10	50	25	500	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	60	10	50	25	493	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	60	10	50	25	561	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	80	10	70	25	563	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	90	10	80	25	560	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	90	10	80	25	574	Brittle	Etna basalt (EB_I); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	90	10	80	25	655	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	110	10	100	25	658	Brittle	Etna basalt (EB_I); porosity = $0.04$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	160	10	150	25	753	Brittle	Etna basalt (EB_I); porosity = $0.05$ : strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	60	10	50	25	365	Brittle	Etna basalt (EB_II); porosity = $0.08$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	90	10	80	25	349	Brittle	Etna basalt (EB_II); porosity = $0.08$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	20	10	10	25	224	Brittle	Etna basalt (EB_III); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	60	10	50	25	434	Brittle	Etna basalt (EB_III); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al. 2016	90	10	80	25	543	Brittle	Etna basalt (EB_III); porosity = $0.05$ ; strain rate = $10^{-5}$ s <sup>-1</sup>

Zhu et al.	110	10	100	25	640	Brittle	Etna basalt (EB_III); porosity =
2016							0.05; strain rate = $10^{-5}$ s <sup>-1</sup>
Zhu et al.	160	10	150	25	798	Brittle	Etna basalt (EB_III); porosity =
2016							0.05; strain rate = $10^{-5}$ s <sup>-1</sup>

718	Table 1: Summary of the experimental conditions for the rock deformation experiments used in this
719	study (for the construction of Figs. 3, 4, and 5) (see also Heap et al., 2017). $Pc =$ confining pressure;
720	$Pp$ = pore fluid pressure; $Peff$ = effective pressure; $T$ = experimental temperature; $\sigma_p$ = peak
721	differential stress (see Figure 1). In some cases, failure mode classification differs from that stated
722	in the original publication. Data not included in this compilation are uniaxial experiments
723	conducted at room temperature and instances of non-viscous ductile deformation (see text for
724	details).

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