1	Emplacement of shergottites in the martian crust inferred from 3D petrofabric and crystal
2	size distribution analyses
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18	Abstract - Shergottites are mafic to ultramafic igneous rocks that represent a majority of known
19	martian meteorites. They are subdivided into gabbroic, poikilitic, basaltic, and olivine-phyric
20	categories based on differences in mineralogy and textures. Their geologic contexts are unknown
21	so analyses of crystal sizes and preferred orientations have commonly been used to infer where
22	shergottites solidified. Such environments range from subsurface cumulates to shallow intrusives
23	to extrusive lava flows, which all have contrasting implications for interactions with crustal
24	material, cooling histories, and potential in situ exposure at the surface. In this study, we present
25	a novel three-dimensional (3D) approach to better understand the solidification environments of
26	these samples and improve our knowledge of shergottites' geologic contexts. Shape preferred
27	orientations of most phases and crystal size distributions of late-forming minerals were measured
28	in 3D using X-ray computed tomography (CT) on eight shergottites representing the gabbroic,
29	poikilitic, basaltic, and olivine-phyric categories. Our analyses show that highly anisotropic, rod-
30	like pyroxene crystals are strongly foliated in the gabbroic samples but have a weaker foliation
31	and a mild lineation in the basaltic sample, indicating a directional flow component in the latter.
32	Star volume distribution analyses revealed that most phases (maskelynite, pyroxene, olivine, and
33	oxides/sulfides) preserve a foliated texture with variable strengths, and that the phases within
34	individual samples are strongly to moderately aligned with respect to one another. In
35	combination with relative cooling rates during the final stages of crystallization determined from
36	interstitial oxide/sulfide crystal size distribution analyses, these results indicate that the olivine-
37	phyric samples were emplaced as shallow intrusives (e.g., dikes/sills) and that the gabbroic,
38	poikilitic, and basaltic samples were emplaced in deeper subsurface environments.
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40	INTRODUCTION
41	Martian meteorites are currently the only physical materials from Mars that can be
42	studied in the laboratory and have provided a wealth of information on the planet's geologic
45	divided into three main entrancing sharesettites, malthlites, and sharesignites, with
44	alvideu mito unee main categories — shergottites, nakinites, and chassignites — with shergottites comprising a vest majority with poorly 200 individual named stores to date
4J 16	Sucreace at al. 2023 McSween 2015 Lidry at al. 2020). Sharaottitos are subdivided into four
+0	(Ganacecca et al. 2025, Micsween 2015, Oury et al. 2020). Shergounes are suburvided into four

47 groups based on textural and mineralogical characteristics: basaltic, gabbroic, olivine-phyric, and

- 48 poikilitic (e.g., Udry et al. 2020). In addition to textural differences, the shergottites are
- 49 subdivided into three geochemical groups based on their bulk trace element and isotope
- geochemistry: geochemically enriched, intermediate, and depleted (Barnes et al. 2020,
 Sarbadhikari et al. 2011, Shearer et al. 2013, Tait and Day 2018). Regardless of lithology or
- 52 geochemistry, most, if not all, shergottites experienced complex crystallization histories
- 52 geochemistry, most, in not all, shergotites experienced complex crystalization instones 53 characterized by episodes of cooling and crystal growth at various depths throughout the martian
- 54 crust with a final period of cooling and solidification during emplacement in the shallow
- 55 subsurface or eruption onto the surface (Balta et al. 2015, Filiberto et al. 2018, First and Hammer
- 2016, Gross et al. 2013b, Gross et al. 2011, Howarth et al. 2014, Lentz and McSween 2000, Liu
 et al. 2013, Liu et al. 2016, McCoy et al. 1992, Rahib et al. 2019, Udry et al. 2017, Usui et al.
 2008, Usui et al. 2010). Shergottites represent the largest and most diverse sampling of martian
- secondary crustal material on Earth and provide a window into volcanic processes on Mars.
 Terrestrial volcanic systems can have a complex topology featuring multiple magma
- chambers, conduits, dikes, cone sheets, and/or sills emplaced in the subsurface (Tibaldi 2015). If 61 the magma erupts onto the surface, lava flows, tubes, and ponds of various morphologies can 62 63 form (Crisp 1984). Both subsurface and surface volcanic features can have a wide range of 64 thicknesses and lateral extents and subsurface features can be emplaced at a variety of depths. 65 Because magma can solidify in any of these environments, igneous rocks with the same bulk 66 composition can display considerable textural diversity. This holds true for shergottites, but their 67 various geological contexts are obscured by the fact that they were ejected from the martian near-surface during impact events. Textural and geochemical analyses of shergottites can provide 68 69 an opportunity to investigate the final stages of their parent melts' crystallization histories, which 70 are then used to infer modes of magma emplacement or eruption and the structure of volcanic 71 systems on Mars.
- 72 Basaltic and gabbroic shergottites are mineralogically similar, with pyroxene and 73 plagioclase (now maskelynite) as the dominant phases. The coarser texture of the gabbroic 74 shergottites (average pyroxene length of 1 mm, up to 5 mm) (e.g., Udry et al. 2020) is interpreted 75 to reflect prolonged and slower crystallization and solidification at greater depths (Filiberto et al. 76 2018, Wenzel et al. 2021) compared to the finer-grained basaltic shergottites (average pyroxene 77 length of 0.3 mm, up to 1 mm) (e.g., Udry et al. 2020), which are variably interpreted to have solidified in environments ranging from hypabyssal (shallow intrusives) to thin or thick extrusive 78 79 flows (Lentz and McSween 2000, McCoy et al. 1992, Mcsween et al. 1996, Smith and Hervig 80 1979, Stolper and McSween 1979). Olivine-phyric shergottites have porphyritic textures with 81 early-formed megacryst (0.5 - 3.0 mm) olivine that crystallized in the lower crust. These are set 82 in fine-grained to glassy groundmasses that are interpreted to have solidified in a large range of hypabyssal to extrusive environments (Ennis and McSween 2014, First and Hammer 2016, Gross 83 84 et al. 2011, Liu et al. 2013, Liu et al. 2016, Sarbadhikari et al. 2009, Usui et al. 2008). Poikilitic 85 shergottites are characterized by having olivine chadacrysts enclosed by large (cm-sized) 86 pyroxene oikocrysts, set in a coarse-grained groundmass consisting of olivine, pyroxene, and 87 plagioclase (now maskelynite). The poikilitic olivine-pyroxene assemblages are interpreted to 88 have formed in the lower crust and the groundmass crystallized during ascent and final 89 emplacement as a coarse-grained intrusive rock (Combs et al. 2019, Howarth et al. 2014, 90 McSween and Treiman 1998, Rahib et al. 2019).
- 91 Most interpretations of how shergottite parent melts were emplaced or erupted cited 92 above are based on analyses of crystal sizes and measurements of mineral shape preferred

orientations (SPO). There is a significant variation in crystal sizes, even within the same
 shergottite subgroups, indicating that a single solidification environment may not be applicable

- 95 for all samples belonging to a specific lithologic category. Further complications arise from the
- 96 fact that crystal size distribution (CSD) analyses have, thus far, been only extrapolated from two-
- 97 dimensional (2D) data, which assumes a mineral's common habits and hence can hinder accurate
- 98 interpretations when the mineral grows out of equilibrium. Similarly, all SPO measurements
- have been made on traditional thin/thick sections, limiting their analyses to 2D (Duke 1968,
- Filiberto et al. 2018, Greshake et al. 2004, Smith and Hervig 1979, Stolper and McSween 1979).
- 101 Without three orthogonal sections, which is uncommon for these rare and typically small 102 samples, interpretations of 2D SPO analyses are limited. Three-dimensional crystallographic
- samples, interpretations of 2D SPO analyses are limited. Three-dimensional crystallographic
 preferred orientations (CPO) have been made on shergottites (Orr et al. 2022) and nakhlites
- (Griffin et al. 2022), a suite of martian clinopyroxene cumulate meteorites unrelated to the
 shergottites, using electron backscatter diffraction (EBSD) analysis.

106 In this study, we examine 3D textures in eight shergottites using X-ray computed 107 tomography (CT) (Fig. 1). The samples were selected to represent all lithologic groupings and 108 provide the largest range of igneous textures. We investigate fabrics of several phases and crystal 109 size distributions of combined oxides and sulfides to derive stress/deformation conditions and 110 relative cooling rates during the final stages of crystallization, respectively. We use this 111 information to infer where in the martian crust these samples solidified, which is important for 112 characterizing the structure of volcanic systems on Mars, determining possible interactions 113 between shergottites and other crustal rocks or subsurface fluids at magmatic and subsolidus 114 temperatures, and identifying rock types that may be present at or near the surface. 115

SAMPLES AND ANALYTICAL METHODS

116117 Samples

118 Olivine-phyric

119 Small (<5 g) chips embedded in a 1-inch epoxy disc were obtained from the Meteorite 120 Working Group for samples Larkman Nunatak (LAR) 12095(40A), LAR 12011(54A), and 121 Elephant Moraine (EETA) 79001(536) lithology A. Their exact masses are unknown because 122 thin and thick sections of unknown masses were previously prepared from these embedded 123 samples. A 2.24 g chip of Tissint was given to M. Fries from the Dupont meteorite collection at 124 the Chicago Field Museum, who in turn made it available for this study. Of these samples, LAR 125 12095, EETA 79001 lithology A, and Tissint are all geochemically depleted shergottites, and 126 LAR 12011 is a geochemically enriched shergottite (Tait and Day 2018). A summary of 127 previously published descriptions for all analyzed samples is found in Supplementary Material 128 (Section A) (DOI: 10.17632/mzgp37947y.1).

129 130 *Basaltic*

An ~4 g chip of Zagami was obtained from the University of New Mexico Institute of
 Meteoritics. Zagami is an enriched shergottite (Borg and Draper 2003).

- 133
- 134 Gabbroic

An ~9 g chip of Northwest Africa (NWA) 6963 was purchased by J. Gross from a dealer in Morocco in 2016, and a 13.5 g piece of NWA 13134 was purchased by Y. Liu from a dealer in

- 137 2013 after being found in Morocco in 2012. Both NWA 6963 and NWA 13134 are enriched
- 138 shergottites (Burney et al. 2022, Filiberto et al. 2018).

139 Poikilitic

The entire mass (35.72 g) of Roberts Massif (RBT) 04261 was scanned as part of the
Astromaterials 3D project (Blumenfeld et al. 2019), and those data were used in this study. RBT
04261 is an enriched shergottite (Usui et al. 2010).

143

144 X-ray Computed Tomography

145 X-ray CT scanning was performed at the Astromaterials X-ray Fluorescence and 146 Computed Tomography (X-FaCT) Lab at NASA Johnson Space Center, the University of Texas 147 High-Resolution X-ray Computed Tomography Facility (UTCT), and the Analysis and Test Lab 148 at NASA Jet Propulsion Laboratory. Scanning conditions and instrumentation are summarized in 149 Table 1 and Supplementary Material (Section B). Software corrections were used to reduce beam 150 hardening and ring artifacts during reconstruction. The final reconstructed data are output as a 151 contiguous series of 2D (X and Y) 16-bit grayscale images (slices) oriented orthogonally to the 152 scan rotation axis (Z), which together comprise a 3D grid of cubic voxels (pixels with a third dimension). Each voxel value (CT number) reflects the effective X-ray attenuation of the 153 154 material comprising that voxel, which is a function of the mean atomic number (z) and mean 155 density of the material, as well as the X-ray energy. Air is the least X-ray attenuating phase and 156 sets the baseline CT #. Phases with a higher relative X-ray attenuation (i.e. high-z and high 157 density) are assigned a higher CT #, and are generally visualized with brighter gray values

- 158 (Ketcham and Carlson 2001).
- 159

160 X-ray CT data analysis

161 Interpreting CT #'s

162 There is no direct correlation of CT #'s to specific phases because the X-ray beam is polychromatic and beam energy varies with location within a sample due to beam hardening. As 163 164 a result, prior knowledge or complementary petrographic analysis of the sample is required to 165 map mineralogy. As shergottites have a primarily mafic/ultramafic mineralogy and have been 166 extensively studied (see Supplementary Material (Section A)), the dominant phases are known to 167 be plagioclase (now maskelynite) + pyroxene (mostly augite and pigeonite) + oxides/sulfides 168 (mostly titanomagnetite and ilmenite and pyrrhotite, respectively) \pm olivine (forsterite content 169 usually ranges from ~Fo₅₀ to Fo₇₅). Phosphates (apatite and merrillite) and impact-melt glass 170 commonly comprise <3 vol.% (Shearer et al. 2015). Figure S1 shows the linear X-ray 171 attenuation coefficient (μ) for common shergottite phases at 80 keV obtained using MuCalc 172 (Hanna and Ketcham 2017), which calculates X-ray attenuation at different X-ray energies using 173 the NIST XCOM database.

All samples feature small, high-attenuation phases, such as oxides and sulfides and some geochemically enriched samples have other trace minerals with high-µ components such as baddeleyite (Moser et al. 2013, Staddon et al. 2021). Distinguishing between these phases in the CT data is problematic because their CT #'s largely overlap (Fig. S1) and, due to their small sizes relative to the size of the whole rock, their grayscales will be affected by blurring and

179 partial-volume effects with surrounding material. Accordingly, we group them together as "high-

180 μ phases." 181

182 Machine learning segmentation

183 Digitally segmenting each CT dataset, which requires assigning each voxel to a specific 184 phase category, is required for object-based quantitative analysis. This is commonly done using a

185 global CT # threshold where a range of CT #'s corresponds to a single phase. However, this is 186 not possible for these samples because there is overlap in the CT #'s for pyroxenes, olivine, 187 phosphates, and basaltic glass (Fig. S1), To overcome CT # overlap, as well as other issues such 188 as mineral zoning and partial-volume effects inherent to CT scanning, we used machine learning 189 to segment the CT datasets. See Supplementary Material (Section C) for further details about 190 interpreting CT #'s and data segmentation. We used Dragonfly[™] software's (Object Research 191 Systems) "Segmentation Trainer" tool to train a classifier on a limited number of manually 192 segmented slices (usually 5 to 30 slices) to enable automatic segmentation of the entire dataset. 193 During training, all voxels were manually assigned to one of five phase categories based on CT 194 #'s and observed petrologic textures: air (both internal voids and external air), maskelynite, 195 pyroxene, olivine, and high- μ phases. Even though pyroxene and olivine have overlapping CT 196 #'s, olivine are easy to distinguish because of their large size, morphology, and/or distinct zoning 197 patterns. Additionally, we could not identify phosphates or impact melt pockets or veins because 198 their ranges of CT #'s overlap with the primary phases and the phosphate grain sizes are near or 199 below the resolution limits. The training slices were used as inputs to the Extra-Trees classifying 200 engine, utilizing the smoothing, neighbors, morphological, median, and moments mathematical 201 filters. Once the classifier was sufficiently trained (i.e., automatically segmented >98% of the 202 manually segmented voxels correctly), it was used to automatically segment the entire CT 203 dataset. A new classifier was trained for each sample. However, even after extensive iterative 204 training, some of the Mg-rich olivine cores in the olivine-phyric samples were improperly 205 segmented as pyroxene because of their overlapping CT #'s and similar textures. As a final 206 processing step after machine learning classification, olivines with incorrectly segmented cores 207 were manually filled to add the mislabeled voxels to the olivine category. The final segmented 208 dataset is a binarized 3D phase map of air, maskelynite, pyroxene, olivine (if present), and high-209 μ phases.

210

211 Measuring modal abundances

212 Normalized modal abundances were measured using the segmented data in *ImageJ* 213 software. Whole-rock modal abundances were determined by counting the number of voxels 214 assigned to each phase and normalizing to the total number of non-air voxels in the entire CT 215 dataset. To enable comparison with other studies that have utilized traditional modal analysis 216 based on 2D thin sections, we also calculated the modal abundance on a slice-by-slice basis 217 along the three orthogonal scan axes (XY, XZ, and YZ) (see Supplementary Material (Section 218 D)). Samples were not specially oriented during CT scanning, so the orthogonal axes are 219 random.

220

221 *CT fabric measurements*

222 Petrofabrics can be measured for each phase in a sample, and different quantification 223 methods are available based on whether the particular phase being examined is treated as a 224 continuum feature, or can be separated into individual mineral grains. We refer to the former as 225 continuum fabrics, and the latter as discrete fabrics, and both as CT fabrics in this paper. Both 226 approaches can provide the type, strength, and direction of preferred orientations, with the 227 principal difference that the discrete fabrics will honor grain boundaries and the continuum 228 fabrics will not. However, as phases constituting high modal percentages can be laborious, 229 problematic, or impossible to separate reliably into grains in CT data, the continuum approach 230 can provide information on more phases, more repeatably and with less effort. As such, we

231 measured continuum fabrics for maskelynite, pyroxene, olivine (if present), and high-µ phases in

- 232 all samples and discrete fabrics for high-µ phases (all samples) and pyroxenes (basaltic and
- 233 gabbroic samples). To facilitate comparing different measurements, all fabrics were
- 234 parameterized using K and C (Woodcock and Naylor 1983), which quantify the fabric type and
- 235 strength, respectively. Fabric strength (C) increase with the degree of anisotropy from zero
- 236 (isotropic) to infinity, and the shape parameter (K) ranges from zero to infinity and defines fabric 237
- shapes (lineation [K > 1], foliation [K < 1], triaxial [K = 1]). A description of the fabric 238 parameters and how to calculate them for each CT fabric quantification method are given in
- 239 Supplementary Material (Section F and G).
- 240
- 241 *Continuum fabrics*

242 Continuum fabrics were measured on the segmented 3D datasets using Quant3D software 243 (Ketcham and Ryan 2004, Ketcham 2005b). Olivine crystals in the olivine-phyric shergottites 244 that are cross-cut by the sample boundary were manually removed from the segmented dataset to 245 prevent measurement of these incomplete crystals. This software integrates data from thousands 246 of discrete measurements to quantify the fabric of the segmented phase. Analytical details and a 247 description of the method are provided in Supplementary Material (Section E). The output data 248 are summarized by a second-rank tensor which provides three orthogonal principal component 249 directions (eigenvectors) and magnitudes (eigenvalues) and a 3D rose diagram is created to aid in 250 detection of non-orthogonal sub-fabric components. Because test point placement is random in 251 Quant3D, the analysis was repeated thirty times for each phase to assess statistical variation. 252 Figure 2 shows a simplified 2D schematic illustrating the Quant3D workflow.

253 The first, second, and third eigenvectors from the thirty replicate analyses for each phase 254 were plotted using Stereonet11 software (Cardozo and Allmendinger 2013). Replicate analyses 255 were plotted to map out confidence regions for each eigenvector and to aid in interpretation of 256 fabric types. Clustered first eigenvectors with girdled second and third eigenvectors indicates a 257 lineation, whereas clustered third eigenvectors and girdled first and second eigenvectors 258 indicates a foliation. Triaxial fabrics are characterized by clustering of all eigenvectors. The 259 average orientation of the 30 replicate analyses was calculated for each eigenvector to compare 260 the CT fabric orientation of one phase to another within the same sample. A description of 261 calculating the angle between CT fabrics within a sample is provided in Supplementary Material 262 (Section H).

263

264 *Discrete fabrics*

265 Pyroxenes in basaltic and gabbroic shergottites feature a texture of intermixed, elongated crystals broken along multiple fractures (Fig. 1e-g). These could only be measured manually in 266 the 3D data, so we created a method for extracting a representative, non-biased sample. Using 267 Dragonfly, a crystal was randomly chosen and the CT data were reoriented along three 268 269 orthogonal views to expose its longest, intermediate, and shortest axes (Fig. 3b-d). We were only 270 interested in measuring long axis length and orientation, so instead of manually segmenting all 271 voxels comprising a single crystal, a rod was drawn through the crystal center along its length 272 using the 3D spherical brush tool (Fig. 3a). The diameter of the rod was smaller than the 273 diameter of the actual crystal to minimize touching of interlocking crystals. To prevent bias 274 based on crystal size or orientation, the 'cine' tool was used to randomly orient the data after 275 each crystal segmentation. The segmented data were imported into Blob3D software (Ketcham

276 2005a) where the few touching rods were manually separated, after which the length (BoxA dimension) and orientation of each crystal long axis was measured. Long axis orientations of

every measured crystal were plotted using Stereonet11 software, and their spatial distributions

quantified by three eigenvalues and eigenvectors using the Bingham axial distribution analysis

(Fisher et al. 1993). These eigenvalues parameterize discrete CT fabrics with the K and C values
 described above (see Supplementary Material (Section G) for calculating discrete CT fabric

parameters). This method, whereby a single 3D shape is used to approximate a phase's length

and orientation, is most effective with objects that have highly regular and anisotropic

284 morphologies (e.g., rods or plates). Measuring a phase's true shape, as in Fig. 3a, requires

segmenting all of its constituent voxels. This approach, or drawing at least three orthogonal
planes and fitting an ellipsoid (e.g., Hanna et al. 2015), is often required for phases with more

286 planes and fitting an ellipsoid (e.g., Hat287 equant or irregular morphologies.

288

289 Measuring highly attenuating phases

290 We also measured the size and orientation of every individual high-µ phase using Blob3D 291 software (Ketcham 2005a) to: (1) calculate their 3D crystal size distribution (CSD) and (2) 292 provide an additional fabric measurement to corroborate the other continuum and discrete CT 293 fabrics. Only interstitial high-µ phases within the groundmass were measured; high-µ phases 294 included within larger minerals (i.e., pyroxene oikocrysts and olivine megacrysts) were excluded 295 from measurement, as we are only interested in grains whose orientations could be influenced by 296 flow and may reflect late-stage growth during the final period of crystallization. We used the 297 unsegmented grayscale data for these analyses because these phases within CT data are small 298 with respect to the sample size, and thus affected by partial-volume and blurring effects (PVB) 299 (Ketcham and Mote 2019), which Blob3D corrects for by using grayscale rather than segmented 300 data, resulting in more accurate ellipsoid and caliper dimensions. A description of isolating only 301 interstitial phases for measurement as well as the Blob3D methods is found in Supplementary 302 Material (Section I).

Similar to the discrete pyroxene fabrics described above, the long, intermediate, and short axis orientation of each high-µ crystal was plotted using Stereonet11 and the distribution of each axis quantified by a single set of eigenvalues and eigenvectors. Eigenvalues were used to parameterize discrete CT fabrics with K and C. The primary eigenvector was used to represent the average orientation for each axis.

308 The BoxA caliper dimension (Ketcham and Mote 2019), which defines the crystal long 309 axis as the longest axis of the smallest rectangular box that fits the particle, was used for crystal 310 size distribution analyses. Because these measurements were made in 3D, there is no 311 requirement for stereological conversion from 2D to 3D measurements (e.g., Higgins 2000, 312 Morgan and Jerram 2006). CSD plots were created using the same bin sizes for all samples so 313 that direct comparisons can be made. The smallest crystal length from all samples was used for 314 the minimum bin size. Each subsequent bin size is 1.1 times as large as the previous one until the 315 bin size was larger than the largest crystal from all samples. The population density (units of 316 mm⁻⁴) was calculated by dividing the number of crystals from each bin size by the volume of 317 sample multiplied by the bin size. Plots of the natural logarithm of the population density versus 318 bin size are commonly used to interpret crystal size distributions. This method is not possible or 319 feasible with other phases in our samples because separating touching and intergrown grains of 320 the same mineral in CT data is not straightforward. 321

322

RESULTS

323 **3D petrographic descriptions**

324 Using CT data for petrographic analyses enables 3D observations of mineral shapes, 325 sizes, orientations, spatial distributions, internal compositional variations, and intergrowths, all 326 of which can be useful for interpreting petrogenesis. Further, CT data can be especially useful for 327 recording other macroscopic textures such as fractures, void spaces, and local melting (Liu et al. 328 2019). Preliminary "hand sample" observations can be made by viewing and manipulating the 329 sample using 3D volume rendering software, and the sample can be digitally sliced along any 330 orthogonal or oblique plane to view internal textures on 2D grayscale images. This is analogous 331 to viewing thousands of back-scattered electron (BSE) images of a serially-sectioned sample cut 332 from all possible angles, but more importantly without processing and physically cutting the 333 sample. In the sections below, we base most of our observations on 2D images obliquely sliced 334 from the 3D data along strategic orientations, e.g., parallel or perpendicular to foliation. Our 335 interpretations are based on what is visible in the studied samples. It is possible that larger or 336 different stones of the same meteorite may differ from ours. Finally, most samples have been 337 meticulously studied previously, and petrographic descriptions on 2D thin sections have been 338 adequate to describe most of the rock textures (see Supplementary Material (Section A)). Thus, 339 we only report new features that have not been recorded or ones that are better studied in 3D.

340

341 Basaltic and gabbroic

342 The basaltic (Zagami) and gabbroic (NWA 6963 and NWA 13134) shergottites are 343 predominantly pyroxene and maskelynite with minor high-µ phases (Fig. 1e-g). Zagami has the 344 most abundant pyroxenes (84 %) followed by NWA 6963 (78 %) then NWA 13134 (73 %) 345 (Table 2). Pyroxene crystals are euhedral and create an interlocking structure of elongated prisms 346 with interstitial maskelynite, which takes on a variety of habits, from blocky to elongated to 347 irregular. High-µ phases are euhedral to anhedral with some equant and blocky habits and some 348 with irregular and rounded habits. They are randomly distributed between crystal boundaries. Cross-cutting fractures are present in all samples. 349

350 Pyroxene crystals, when viewed along a random orientation, have a large range of sizes 351 (from < 1 mm up to 10 mm) and morphologies (from equant and rounded to highly elongated 352 and rod-like). However, when the CT data are strategically oriented along a single crystal's 353 longest, intermediate, and shortest axes, it becomes apparent that pyroxene crystals have a 354 slightly compressed rod-like morphology (Fig. 3). This is true for nearly every crystal in NWA 355 13134 and NWA 6963, whereas a small population of pyroxenes in Zagami are more equant and 356 blocky with sharp edges that meet at approximately 60° and 90°. We are confident the elongated 357 crystals are single, large crystals instead of linked, smaller crystals, because there are small (~50 358 µm thick) bright rims that are three-dimensionally continuous around the entire crystal. 359 Furthermore, fracture spacing, size, and orientation are consistent within a single crystal. Table 3 360 summarizes the length of measured pyroxene crystals. NWA 6963 and NWA 13134 are nearly 361 identical with crystal lengths up to around 13 mm (average = 5.8 mm and 6.4 mm, respectively) 362 and Zagami crystal lengths up to 6.7 mm (average = 2.3 mm). Only crystal lengths were 363 quantified by the rod analysis, but visual observations show that crystal length and diameter 364 scale with one another with NWA 6963 and NWA 13134 crystals up to ~1 mm diameter and 365 Zagami up to ~0.5 mm.

367 *Olivine-phyric*

368 Considerable textural diversity exists among the four studied olivine-phyric shergottites 369 (Fig. 1a-d). All are porphyritic with megacrysts of olivine surrounded by a fine- to medium-370 grained basaltic groundmass of predominantly maskelynite and pyroxene. Samples LAR 12011 371 and Tissint have the most abundant olivine megacrysts (27 % and 26 %, respectively) followed 372 by LAR 12095 (17 %) and EETA 79001 (14 %) (Table 2). Megacryst olivine are up to 4 mm and 373 their morphology/habit vary. Euhedral to subhedral crystals are commonly clustered and 374 touching in samples LAR 12011 and Tissint, whereas olivine in LAR 12095 and EETA 79001 375 are more isolated. LAR 12095 olivine are the most euhedral with flat faces and sharp vertices 376 and edges, whereas many olivines in EETA 79001 have non-linear edges and rounded vertices 377 and reentrant features, although most are still subhedral to euhedral. Olivine in LAR 12095, 378 Tissint, and EETA 79001 have mostly equant dimensions, whereas many of the olivine, 379 especially the larger ones, in LAR 12011 are more tabular.

380 Core-to-rim zoning from lower to higher CT #, which reflects increasing fayalite 381 content, is apparent in large olivine in all samples, with EETA 79001, LAR 12011 and Tissint 382 having well-defined rims that are usually around 50 µm thick) with a higher fayalite content and 383 LAR 12095 having a more subtle increase in favalite content from core to rim. Zoning is not 384 apparent in the smaller olivine within all samples and they have high CT #'s (i.e., more fayalitic) 385 comparable to the rims of the larger olivine. Olivine-hosted inclusions are common in LAR 386 12095, LAR 12011, and Tissint and rare in EETA 79001. Dark inclusions are interpreted to be 387 melt inclusions and brighter inclusions are interpreted to be primarily oxides (e.g., chromite). 388 Melt inclusions vary from spherical to irregularly shaped and from near the resolution limits of 389 the scan up to several hundred micrometers long. Oxide inclusions are generally equant and 390 about 50 µm long.

391 Groundmass phases are interpreted to be mostly maskelynite and pyroxene with 392 interstitial high-µ phases that are interpreted to be oxides and sulfides. The average abundance of 393 groundmass phases (i.e., pyroxene and maskelynite) is 78 % (range = 72 % to 86 %) (Table 2). 394 Tissint and EETA 79001 have the finest groundmass phases. LAR 12011 has slightly larger 395 groundmass maskelynite with a feathery and elongated morphology and LAR 12095 has the 396 largest groundmass phases with some pyroxene growing up to a few mm long and maskelynite 397 taking on a blocky and more equant habit. Systems of sub-parallel fractures are found in most 398 large olivine but do not propagate into surrounding phases. They are dominantly oriented 399 orthogonally to the crystal long axis. Cross-cutting fractures are found in all samples. A single 2 400 mm x 2 mm shock-melt pocket is found in EETA 79001 based on its unique CT #'s, rounded 401 shape, and internal texture (Fig. 1a – dark circular object in middle of image).

402403 *Poikilitic*

404 RBT 04261 is a coarse-grained assemblage of poikilitic pyroxenes surrounded by a non-405 poikilitic groundmass of olivine, pyroxene, and maskelynite (Fig. 1h). The pyroxene oikocrysts 406 are up to 15 mm long and enclose smaller (up to 2 mm) olivine chadacrysts and fine-grained 407 high-µ phases. The non-poikilitic texture contains coarse-grained olivine up to 6 mm long, 408 irregular pyroxene up to 1 mm long, blocky maskelynite (21 %; Table 2), and interstitial high-µ 409 phases. The non-poikilitic olivine are euhedral and form an interlocking framework with 410 interstitial maskelynite. These olivines also have higher fayalite content than the olivine 411 chadacrysts. Zoning is not apparent in either olivine type. Combined (poikilitic and non-412 poikilitic) olivine and pyroxene abundances are 54 % and 25 %, respectively (Table 2).

414 Modal abundances

415 Whole-rock and slice-by-slice modal abundances are shown in Fig. 4 and reported in 416 Table 2. We report the whole-rock value and the average, 25 standard deviation, and Min/Max of 417 the thousands of measured slices and include previously published values for comparison. For 418 each sample, the average of the thousands of measured slices is usually within 1% of the whole-419 rock value. However, individually, there is significant variation on a slice-by-slice basis, 420 especially for megacryst olivine (up to 31 % in EETA 79001) and pyroxene oikocrysts (up to 26 421 % in RBT 04261). The slice-by-slice variation is less for the basaltic and gabbroic samples. For 422 samples LAR 12095, EETA 79001, and Tissint, the published values are mostly within the total 423 range of slice-by-slice values, whereas the published values are outside this range for Zagami,

424 NWA 6963, and LAR 12011. 425

426 **Petrofabrics**

427 Continuum CT fabric parameters (K and C) of each measured phase (i.e., maskelynite, 428 pyroxene, olivine, and/or high-µ phases) are listed in Table 4 and illustrated on an eigenvalue 429 ratio plot (Woodcock and Navlor 1983) in Fig. 5. On average, basaltic and gabbroic shergottites 430 have the highest fabric strengths (C) (0.59 and 0.62, respectively) followed by the olivine-phyric 431 (0.49) then poikilitic (0.47) shergottites. Most phases display a triaxial (K = 1) to foliated (K < 1) 432 fabric with average K values of 0.64 (basaltic), 0.61 (gabbroic), 0.53 (olivine-phyric), and 0.41 433 (poikilitic). The orientations of continuum fabrics within a single sample are strongly to 434 moderately aligned with respect to one another (Fig. 6 and Table 5). All phases in Zagami, NWA 435 6963, LAR 12095, LAR 12011, and RBT 04261 are strongly aligned (i.e., within 15° of each 436 other) and moderately aligned (i.e., all phases within 25° of each other) in NWA 13134, Tissint, 437 and EETA 79001.

438 The fabric based on discrete pyroxene crystals measured with Blob3D in Zagami, NWA 439 13134, and NWA 6963 indicates foliations similar to the continuum results, but much stronger, 440 with C values higher by a factor of 4 (Figs. 7, S5; Tables 4, S2). NWA 6963 has a nearly uniaxial 441 girdle distribution (i.e., all orientations along the same plane), whereas NWA 13134 and Zagami 442 have around 10 % and 15 %, respectively, of crystals oriented away from the foliation plane. 443 Zagami has a mild preferred orientation within the foliation plane shown by a slight clustering of 444 pyroxene orientations. The pyroxene foliation planes measured with this discrete method 445 generally overlap with the pyroxene continuum measurement; they are within 13.9° of one 446 another in NWA 13134, 20.3° in NWA 6963, and 24.2° in Zagami (Fig. S8). These large 447 differences in fabric strength, and minor differences in orientation, are likely due to a 448 combination of the incomplete sampling of pyroxene crystals for the discrete method and the 449 continuum method merging pyroxene-pyroxene grain boundaries.

Like the pyroxene CT fabrics, the high- μ phase discrete fabrics indicate foliations similar to the high- μ phase continuum results, but have more comparable degrees of anisotropy, with C values only higher by a factor of around 1.5 (Fig. S6; Table S1). The high- μ phases' average long, intermediate, and short axis orientations measured with Blob3D (gray data in Fig. 6; Fig S2) are very similar to high- μ Quant3D results. See Supplementary Material (Section J) for a discussion comparing high- μ and pyroxene fabric measurements using Quant3D and Blob3D.

457 Crystal size distributions

458 Results from 3D crystal size distribution analyses for the high-µ phases are presented in 459 Fig. 8 and Table 6. Between 3,900 and 105,000 individual grains were measured per sample with 460 an average of 24 grains per mm³ (range 3.8 to 74.3 grains/mm³). The basaltic and gabbroic 461 samples have the largest average grain lengths (from 0.12 to 0.26 mm) followed by the poikilitic 462 sample RBT 04261 (0.13 mm) and the olivine-phyric samples (from 0.05 to 0.09 mm). The 463 average ratio of long to short axis (BoxA/BoxC in Table 6) range from 2.18 to 2.41 for the 464 gabbroic and basaltic samples and from 1.73 to 2.03 for the olivine-phyric samples and 1.67 for 465 the poikilitic sample. All samples have a mostly linear and negative CSD slope from their 466 maximum bin size to their turndown bin size. There is a slight increase in slope right before the 467 turndown at smaller bin sizes in all samples except RBT 04261. There is greater deviation from a linear slope at larger bin sizes due to smaller numbers of grains. The steep turndown at the 468 lowest bin sizes is likely a scanning resolution effect as there is a correlation ($r^2 = 0.83$) between 469 470 the scanning voxel size and the turndown bin size (Fig. S3), which is around four times as large 471 as the voxel size (slope = 0.27). The shortest axis (BoxC) is the controlling factor for visibility in 472 the CT data, which is usually around half the length of the longest axis (BoxA), implying that the 473 thinnest axis length where grains are beginning to be excluded from measurement is around two 474 voxels. We only use the values greater than the turndowns for linear regressions and interpreting 475 CSD patterns because data in bin sizes below the respective turndowns are likely to be in part a 476 resolution effect and not a petrogenetic feature. 477 Distinctly different CSD patterns correlate with lithology. The basaltic and gabbroic

478 samples have shallower slopes (-8 to -3 mm⁻¹) and extend to larger bin sizes (1.3 to 3.5 mm), 479 especially NWA 13134 and NWA 6963, which are nearly indistinguishable. The latter two have 480 a slight increase in slope at around 0.3 mm sizes before the turndowns. The olivine-phyric 481 samples have steeper slopes (-34 to -11 mm⁻¹) and smaller maximum sizes (<0.76 mm). LAR 482 12095, LAR 12011, and Tissint are nearly linear from the largest bin sizes down to the turndown 483 bin sizes, whereas EETA 79001 is linear from the largest bin sizes to around 0.24 mm and 484 slightly concave up from 0.24 mm to the turndown bin size. The poikilitic sample has a slope of -485 12 mm⁻¹ and a maximum bin size of 1.0 mm with a highly linear ($r^2 = 1.00$) CSD pattern down to 486 its turndown bin size.

487 488

DISCUSSION

489 Implications of grain-scale heterogeneity

490 The importance of measuring whole-rock versus slice-by-slice modal abundances is 491 illustrated in Figs. 4 and 9 and Table 2. The average modal abundance measured on thousands of 492 slices per sample is within ~1% of the whole-rock value, but any single slice can vary 493 significantly from the whole-rock modal abundance. This is especially true for samples with 494 isolated phases that are significantly larger than groundmass phases (e.g., phenocrysts). The 495 likelihood of olivine megacryst or pyroxene oikocryst abundance on a random slice being ± 1 % 496 of the whole volume value decreases from 33.9 % (LAR 12011) to 20.9 % (Tissint) to 17.5 % 497 (RBT 04261) to 17.0 % (LAR 12095) to 10.8 % (EETA 79001). Similarly, the 2RSD (relative 498 standard deviation) for megacryst phase abundance on the thousands of CT slices increases from 499 16.5 % (LAR 12011) to 30.5 % (Tissint) to 39.7 % (RBT 04261) to 60.5% (LAR 12095) to 500 105.5% (EETA 79001).

501 This divergence has significant implications when classifying samples and allocating 502 "representative" thin/thick sections or aliquots for petrological/geochemical analyses. Initial

503 meteorite classifications are sometimes based on observations from a single section. Fig. 9 shows 504 the minimum and maximum modal abundances of olivine megacrysts and pyroxene oikocrysts 505 that can be observed on a single slice from the same samples. For instance, if the minimum 506 olivine megacryst slice for EETA 79001 was sectioned for classification, this sample would be 507 classified as a basaltic lithology, and not olivine-phyric. Similarly, there are no intact pyroxene 508 oikocrysts in the minimum abundance slice for RBT 04261. Based on this slice, RBT 04261 509 would be classified as an olivine cumulate with little pyroxene. While it is unlikely to randomly 510 cut the sample on a plane with the minimum or maximum value for a specific phase, for a 511 sample like EETA 79001, which has a high megacryst 2RSD (>100%) and a low likelihood 512 (~10%) of a random slice being $\pm 1\%$ of the whole volume average, there is a higher likelihood of misinterpreting the lithology from a single slice. Based on our results, the samples we studied 513 514 are accurately classified. However, new samples, or volumetrically small samples where as few 515 as one thin section can be made, would benefit from being classified using X-ray CT. 516 Additionally, CT scanning can allow a specific plane to be selected for sectioning (e.g., Hanna et 517 al. 2015), or a representative and contextualized volume of sample be allocated for 518 geochemical/isotopic analyses via dissolution. For instance, the olivine megacrysts are evenly 519 distributed throughout LAR 12011, so a random volume from this sample should be 520 representative. However, the distribution of olivine megacrysts in LAR 12095, EETA 79001, and 521 Tissint is heterogeneous, requiring a larger volume of rock to obtain a similarly representative

522 523 aliquot.

524 Relative cooling rates from CSD analysis

525 There are distinct interstitial high-µ CSD patterns for the basaltic, gabbroic, olivine-526 phyric, and poikilitic lithologies. For the gabbroic and basaltic samples, it is interesting that 527 NWA 6963 and NWA 13134 have almost identical CSD patterns. Based on this and other 528 textural similarities (i.e., pyroxene sizes and modal abundances). NWA 13134 might be paired 529 with NWA 6963. NWA 13134 is a relatively new and unstudied sample (Burney et al. 2022), so 530 future geochemical and/or geochronological work may confirm this pairing. Extending this high-531 μ CSD method to additional shergottites may prove to be a relatively quick and quantitative tool 532 for lithologic classification and possibly aid in defining meteorite pairing groups.

533 Effective application of CSD theory requires that every crystal from a specific mineral 534 group (e.g., all pyroxenes or all olivines) is individually measured within a sample. The 535 interstitial high-µ phases measured here are interpreted to be both oxides and sulfides, which 536 have been commonly reported as interstitial opaque minerals in other petrographic studies and 537 are some of the last phases to crystallize (Barrat et al. 2002, Gattacceca et al. 2013, Goodrich 538 2003, Herd et al. 2002, Herd et al. 2001, McCoy et al. 1992, Sarbadhikari et al. 2009). These 539 authors found that pyrrhotite are smaller ($<20 \mu m$) and less abundant (<1 %) than the interstitial 540 oxides (1 - 2%), which are dominantly titanomagnetite and also include subordinate abundances 541 of ilmenite. While we cannot distinguish between the sulfides and oxides in the CT data, 542 previous investigations indicate that most sulfides are likely within bin sizes below the CSD 543 turndowns. A small population of the relatively larger sulfides added to the oxides in bin sizes 544 near the turndown might be the cause of the slight increase in slope near the turndown. 545 Therefore, the CSD patterns reflect mixing of different but overlapping mineral size classes. 546 However, due to the sulfides' small sizes, the patterns most likely reflect the growth of oxides,

547 mainly titanomagnetite.

548 Most shergottite CSD analyses are of olivine and/or pyroxene with the goal of 549 understanding: (1) physical processes within the parent magma such as crystal accumulation or 550 fractionation; (2) crystal growth rates; (3) residence times; and (4) nucleation histories (Ennis 551 and McSween 2014, Filiberto et al. 2018, Lentz and McSween 2000, Rahib et al. 2019). The goal 552 of our high-µ CSD analyses is to understand the final stages of crystallization as interstitial 553 oxides are some of the last phases to form. If we assume the high-µ phases and growth rates are 554 the same in all samples, the steeper CSD slopes are interpreted to reflect a shorter residence time 555 (i.e., faster cooling rate during parent melt solidification). To calculate residence times, the 556 crystal growth rate must be known and assumed to be constant. The generally linear form of the 557 CSD patterns is consistent with a constant growth rate under steady-state conditions. 558 Unfortunately, there have been no experiments measuring growth rates of titanomagnetite in 559 shergottitic melts, thus, quantifying residence times is not possible. However, we can infer 560 relative differences in residence times. The gabbroic samples (NWA 6963 and NWA 13134) had 561 the most prolonged crystallization period, followed by Zagami (basaltic) and RBT 04261 (poikilitic), then by the olivine-phyric samples (LAR 12011, LAR 12095, EETA 79001, and 562 563 Tissint). This interpretation is consistent with other groundmass mineral grain sizes with the 564 gabbroic samples being the coarsest followed by the basaltic and poikilitic samples then the 565 olivine-phyric samples. These results cannot be used to infer the specific solidification 566 environment for each sample but are useful for determining relative differences in cooling rates 567 during the final period of crystallization.

568

569 Interpreting fabrics

570 Continuum and discrete fabric analyses revealed that most phases preserve a foliated to 571 triaxial fabric and that CT fabrics of multiple phases measured within a single sample are 572 strongly to moderately aligned. Generally, foliated igneous fabrics are interpreted to form in 573 environments where the principal forces are normal to the foliation plane with limited simple 574 shear, possibly from compaction in a thicker lava flow or settling/compression in a sub-volcanic sill, dike, or chamber (Meurer and Boudreau 1998, Nicolas et al. 2009). Simple shear, which is 575 576 the dominant deformation regime in lava flows, is expected to impart a foliation with a preferred orientation (Jezzi and Ventura 2002, Merle 2000, Reed and Tryggvason 1974). However, a 577 578 complementary analogue study of CT fabrics in terrestrial mafic rocks with various geologic 579 contexts (dikes, sills, flows, and lava lakes) shows there is no correlation between the CT fabric 580 parameters and solidification environment because of several factors intrinsic to an individual 581 sample and extrinsic ones related to the magma itself (Eckley 2022). Interestingly, they do show 582 that the only samples (n = 3) with strong degrees of CT fabric alignment are from shallow 583 intrusive environments (i.e., dikes and sills) where the principal stresses associated with magma 584 emplacement in some dikes or sills may be more constant and uniformly align phases whereas 585 extrusive flows tend to be more chaotic during flow and solidification.

586 Comparing our CT fabric results to the analogue study by Eckley (2022) is complicated 587 by differences in minerals and their abundances and shapes, degrees of aggregation, and 588 presence and coalescence of vesicles, among other factors. However, we still posit that shear 589 forces are more consistent in subsurface emplacement environments than within lava flows on 590 Mars, which uniformly aligned the measured CT fabrics. It may also be possible that phases can 591 become strongly aligned at the base of lava flows, especially thicker ones, but we do not yet have 592 data to test this hypothesis. Therefore, we interpret the strong to moderate degrees of CT fabric alignment in the samples from this study to reflect uniform and consistent principal forces duringsolidification, most likely in subsurface environments.

595 Additional evidence for a subsurface emplacement environment for the basaltic and 596 gabbroic samples is borne out by the fabric analysis of individual pyroxene crystals. Pyroxene 597 crystals in gabbroic samples (NWA 13134 and NWA 6963) are almost exclusively large and rod-598 like and show a strongly foliated fabric with no preferred orientation. We interpret this fabric to 599 have likely formed from the settling, accumulation, and compaction of pyroxene crystals in a 600 subsurface magma chamber with no simple shear component (i.e., flow). Most of the pyroxene 601 crystals in the basaltic sample (Zagami) are also rod-like but have a slightly weaker pyroxene 602 foliation and a mild preferred orientation within the foliation plane. This fabric suggests there 603 was a mild component of simple shear during solidification, most likely from directional forces 604 associated with flow. 605

606 Models for shergottite emplacement

Here we describe models for the emplacement of shergottites in this study inferred from
 both the petrofabric analyses and high-μ phase crystal size distribution analyses in the context of
 previous studies.

610611 *Gabbroic*

612 The coarse-grained nature of the gabbroic shergottites (NWA 13134 and NWA 6963) is 613 interpreted to reflect an emplacement environment with slower cooling rates compared to other groups. This interpretation has been reiterated by Udry et al. (2017) for NWA 7320 through 614 615 geochemical and petrological analyses and by Filiberto et al. (2018) through geochemical, petrological, and quantitative 2D textural analyses of pyroxene in NWA 6963. Filiberto et al. 616 (2018) interpreted their CSD plots to reflect accumulation of crystals with relatively long 617 618 residence times. Interestingly, the CSDslice (Morgan and Jerram 2006) software used in their 619 study, which is commonly used to stereologically predict 3D shapes from 2D data, matched a 620 rectangular prism habit with an aspect ratio of 1.00:1.20:1.90 and a maximum length of around 621 3.25 mm. When viewed in 3D along strategic orientations, the pyroxene in NWA 6963 are rod-622 like with highly anisotropic aspect ratios (1.00:1.60:20.00 for the pyroxene in Fig. 3) and an average length of 5.8 mm (up to 13 mm). Their work followed the standard procedure for 2D 623 624 CSD analyses, but caution should be taken when measuring samples like NWA 6963 that have 625 highly anisotropic crystal habits. While the thin/thick section making process is often out of the 626 analyst's control, a single, random 2D thin section may not be adequate for accurate CSD 627 analyses and to identify rod-like shapes or other 3D shapes. A section oriented along the foliation plane (Figs. 10-column B and S9-left) would provide a more realistic representation of the rock. 628 629 Similarly, the 2D pyroxene shape preferred orientation (SPO) analyses resulted in a dominantly 630 clustered distribution, which is certainly an artifact of their section not being parallel to the 631 foliation plane. Nevertheless, based on their combined quantitative textural and geochemical 632 analyses, they infer that NWA 6963 was emplaced as a cumulate rock in the Martian subsurface. 633 Our results corroborate this interpretation. All evidence (relatively long late-stage residence time 634 from high-µ CSD analyses and a strongly foliated fabric of large and rod-like pyroxene crystals) 635 points toward emplacement in a subsurface environment with settling and accumulation of 636 pyroxene and no flow component, likely in a sill or magma chamber, for NWA 6963 and NWA 637 13134.

639 Basaltic

640 Zagami has long been recognized as having a relatively slow cooling rate during the final 641 crystallization period when compared to other basaltic and olivine-phyric shergottites (Becker et 642 al. 2011, McCoy et al. 1992, Stolper and McSween 1979). This was inferred from its coarse-643 grained nature, high crystal abundance, and thick pyroxene exsolution lamellae. They 644 hypothesized that these conditions are achievable in a thick (> 10 m) lava flow or thin intrusive 645 environment. Stolper and McSween (1979) and Becker et al. (2011) measured foliated pyroxene 646 SPO and crystallographic preferred orientation (CPO), respectively, and inferred that Zagami 647 formed in an intrusive cumulate environment with no directional flow. McCoy et al. (1992) 648 measured a similar foliated pyroxene SPO but stated that without confirmation of a lineation 649 associated with the foliation, then a cumulate environment is not substantiated. Here we confirm 650 that there is a foliated pyroxene SPO but with a mild preferred orientation along the foliation 651 plane. However, this may not indicate crystallization in a thick lava flow as injection of a magma 652 into a dike may impart a flow direction component onto the rock fabric. The strong alignment of CT fabrics in Zagami also suggests solidification in an environment where principal forces align 653 654 all phases similarly. Again, Eckley (2022) shows that alignment of phases does happen in a 655 subsurface environment but hypothesize it may also occur in a thick lava flow. The final line of 656 evidence against eruption to the surface and solidification as a lava flow is the high pyroxene 657 abundance (84%), which is drastically higher than the critical crystallinity (\sim 55%) that prevents a 658 basaltic magma from erupting and flowing (Marsh 1981). Therefore, we propose that Zagami 659 formed in an intrusive body, possibly injected as a dike, at a shallower depth, or in a smaller 660 volume magma body, than the gabbroic shergottites.

661

662 *Poikilitic*

Poikilitic sample RBT 04261 is hypothesized to have been emplaced in an environment 663 664 with relatively slow cooling rates, likely in a hypabyssal cumulate environment. This interpretation is based on quantitative 2D textural analyses of the non-poikilitic olivine in RBT 665 666 04261 and several other poikilitic shergottites (Combs et al. 2019, Howarth et al. 2014, Rahib et 667 al. 2019, Udry et al. 2020) that show accumulation of the non-poikilitic olivine. The strong 668 alignment of CT fabrics and high crystal content (olivine + pyroxene oikocrysts = 78 %) 669 measured here are consistent with emplacement in a subsurface cumulate environment. Based on 670 similar high-µ CSD patterns, RBT 04261 may have been emplaced at temperature and pressure 671 conditions similar to Zagami.

672

673 *Olivine-phyric*

674 The olivine-phyric shergottites have the finest-grained groundmass minerals and high-u CSD patterns which both reflect the quickest cooling rates during solidification. Olivine-phyric 675 shergottite Yamato 980459 is inferred to have erupted and quenched in a thin lava flow based on 676 677 its high abundance of glassy mesostasis (Usui et al. 2008) and dynamic crystallization 678 experiments (First and Hammer 2016). Aside from Yamato 980459, most shergottites are simply 679 interpreted to have experienced a relatively high cooling rate during the final period of 680 crystallization based on their fine-grained groundmass minerals and distinct Fe-rich olivine rims 681 (Dunham et al. 2019, Ennis and McSween 2014, Liu et al. 2013, Liu et al. 2016, Sarbadhikari et 682 al. 2009, Udry et al. 2020). Such a cooling rate is not only achievable during extrusion as a lava 683 flow, but also during emplacement of shallow intrusives (i.e., dikes and sills). Chilled margins, 684 quenched textures, and Fe-rich olivine rims are all common in terrestrial basaltic dikes and sills

685 (Petcovic and Dufek 2005). Balta et al. (2015) reported highly evolved late-stage minerals (Ti-

rich oxides, Fe-rich olivine, and Fe-rich merrillite) in Tissint that required a prolonged cooling

during the final stages of crystallization. They infer that slow final cooling requires emplacement

688 in a sill or dike rather than eruption at the surface. Here we show that LAR 12011, LAR 12095,

689 EETA 79001, and Tissint all have strong to moderately aligned CT fabrics which we interpret as

having formed during emplacement into a shallow intrusive environment, likely dikes and/or
 sills that are possibly associated with a volcanic edifice. The depth of emplacement is unknown,

but at shallower depths, and/or in thinner magma bodies, than the gabbroic, poikilitic, and

- 693 basaltic shergottites in this study.
- 694

695 Implications of subsurface emplacement

696 Determining where within the martian crust shergottites finally crystallized has broad 697 implications for better understanding possible sources of syn- and post-emplacement 698 modification, remote sensing observations, and the growth and evolution of martian volcanic 699 centers. Prolonged crystallization at depth versus rapid quenching at the surface vary 700 significantly in the amount of time the rock spent at elevated subsolidus temperatures. This can 701 have drastic effects on the thermally-activated diffusion of rapidly-diffusing elements (e.g. H, Li, 702 Be, B) whose in-situ geochemical profiles have been used to infer possible volatile outgassing 703 (Lentz et al. 2001, Udry et al. 2016) or presence of primordial volatile reservoirs using volatiles 704 in glasses from mineral-hosted melt inclusions (Gross et al. 2013a, Usui et al. 2012) or in 705 volatile-bearing minerals like apatite (Filiberto et al. 2016, Gross et al. 2013a, McCubbin et al. 706 2016, McCubbin et al. 2012). Shergottites that experienced rapid cooling are likely to be less affected by thermally-activated diffusion, making interpretations of their geochemical profiles 707 708 more robust.

709 While shergottites represent only a small and selective portion of the martian crust, they 710 are useful to help understand some remote sensing observations. For instance, McSween et al. 711 (2006) identified similarities between Gusev crater rocks and olivine-phyric rocks. Since olivine-712 phyric shergottites are more likely to have been emplaced closer to the surface when compared 713 to other shergottites in this study, this could provide an explanation for a high abundance of 714 olivine at the surface. Given the ancient ages of the Gusev crater olivine-rich rocks (~3.65 Ga (Greeley et al. 2005)) compared to the much younger shergottite ages (~165-600 Ma (Borg et al. 715 2003, Brennecka et al. 2014, Nyquist et al. 2001, Váci and Agee 2020)) this could imply a 716 717 tendency for solidification of olivine-rich samples in near surface environments on Mars over a 718 wide span of geologic time. Determining the solidification environments of shergottites is a 719 crucial step in trying to give these rocks geologic context.

720 Finally, the samples in this study are a small but representative suite spanning the dominant lithological and geochemical categories. Extending the analyses we presented here to a 721 722 statistically significant number of samples with various crystallization ages, ejection ages, and 723 geochemical signatures may provide an opportunity to understand the crystallization conditions 724 of individual volcanic systems on Mars. For instance, Lapen et al. (2017) identified a suite of 725 shergottites (n = 11) with the same ejection age and mantle source signature, but with 726 crystallization ages from 348 Ma to 2.4 Ga, suggesting the presence of a long-lived volcanic 727 center. A systematic petrofabric analysis of such a suite of samples could provide insight into 728 how volcanic centers have grown and evolved on Mars. 729

730

CONCLUSIONS

731 The analysis of 3D petrofabrics and crystal size distributions in shergottites using X-ray 732 computed tomography provides valuable insights into their final moments of crystallization, 733 which reflect the environments in which these martian igneous rocks solidified. We used a 3D 734 star volume distribution method to measure the strength, type, and direction of shape preferred 735 orientation of the phases (i.e., maskelynite, pyroxene, olivine, and/or oxides/sulfides) within 736 eight shergottite meteorites covering the known lithologic and geochemical subcategories. Most 737 phases preserve a foliated to triaxial petrofabric with variable degrees of anisotropy. Five out of 738 eight samples have strongly aligned petrofabrics where the eigenvector orientations for the 739 different phases within a sample are aligned to within 15° of each other, and the other three 740 samples have moderately aligned petrofabrics. This alignment is likely caused by uniform 741 stresses during solidification, which may be common in intrusive environments. These results 742 indicate that the samples in this study were probably emplaced in the subsurface, either as 743 cumulates or shallow intrusives. Measurements of individual pyroxene crystal orientations 744 provide further refinement of emplacement models for the gabbroic and basaltic shergottites in 745 this study. Gabbroic shergottites were emplaced in a subsurface cumulate environment with no 746 flow, possibly a large sill or magma chamber, whereas the basaltic shergottite was emplaced in a 747 subsurface environment with a minor flow component, possibly injected as a dike. The crystal 748 size distribution patterns of interstitial oxides and sulfides are unique for the different lithologic 749 types and reflect relative differences in cooling rates between different sample types during the 750 final stages of crystallization. Together, these results are used to evaluate the solidification 751 environments of the samples from this study. Gabbroic (NWA 6963 and NWA 13134), basaltic 752 (Zagami), and poikilitic (RBT 04261) shergottites were emplaced in the subsurface with 753 gabbroic samples solidifying at deeper depths, or in larger magma bodies. Olivine-phyric 754 shergottites (LAR 12011, LAR 12095, EETA 79001, and Tissint) were also emplaced in the 755 subsurface but as shallower intrusive bodies such as dikes or sills. The degree of geochemical 756 enrichment or depletion of each sample did not correlate directly to a sample's petrofabric or 757 emplacement as enriched shergottites existed across the full range of emplacement modes identified in this study. The absence of such a correlation between cooling history, depth of 758 759 emplacement, and degree of geochemical enrichment is consistent with the idea that the source 760 of enrichment within the enriched shergottites may not be, solely, the martian crust (Barnes et al. 2020, Brandon et al. 2012, Sarbadhikari et al. 2009, Sarbadhikari et al. 2011, Shearer et al. 2013, 761 762 Tait and Day 2018). Subsurface emplacement histories have important implications for 763 understanding the interaction of shergottite melts with upper crustal materials, cooling histories, 764 and remote sensing observations.

765 766

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783	
784	DATA AVAILABILITY
785	The data that support the findings of this study are available in the supplementary
786	material of this article which is openly available in a Mendeley Data Repository (DOI:
787	10.17632/mzgp37947y.1).
788	
789	CONFLICT OF INTERESTS
790	The authors declare that they have no known competing financial interests or personal
791	relationships that could have appeared to influence the work reported in this paper.
792	
793	References
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TABLES

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Table 1: CT scanning facilities and conditions

Sample	Facility	X-ray energy (kV)	X-ray power (W)	Beam filtering	Exposure (s)	Projections	Frame averaging	Voxel size (µm)	Dimensions (voxels)
Zagami	JSC	90	3.0	none	2.83	3141	2	11.49	1708 x 1436 x 1071
NWA 13134	JPL	170	7.7	none	0.64	2305	8	10.70	2737 x 2716 x 1792
NWA 6963	JSC	100	3.0	0.1 mm Cu	1.42	2121	2	18.03	1200 x 1134 x 2000
LAR 12095	JSC	95	3.0	0.1 mm Al	2.00	2521	2	8.57	1634 x 1396 x 1360
LAR 12011 EETA	JSC	95	3.0	0.1 mm Al	2.00	2831	4	12.49	1364 x 1456 x 1140
79001	JSC	95	3.0	0.1 mm Al	2.00	3141	4	13.14	1362 x 1590 x 775
Tissint	JSC	95	3.0	0.1 mm Al	2.00	1891	2	12.69	836 x 979 x 1622
RBT 04261	UTCT	190	34.0	Al	1.00	3000	2	19.40	1494 x 1589 x 1880

1041 **Table 2:** Whole-rock and slice-by-slice modal abundances

Sample/PhaseAbundancesMean(26)Min.Max.# of SlicesZagami	Whole-rock Individual Slice Abundances						nces
ZagamiMaskelynite14.014.02.311.019.02453Pyroxene83.783.72.378.786.91High-µ2.32.30.51.33.11NWA 13134 $IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII$	Sample/Phase	Abundances	Mean	(2σ)	Min.	Max.	# of Slices
Maskelynite14.014.02.311.019.02453Pyroxene83.783.72.378.786.9High- μ 2.32.30.51.33.1NWA 13134 </td <td>Zagami</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>	Zagami						
Pyroxene83.783.72.378.786.9High-μ2.32.30.51.33.1NWA 13134 $Maskelynite$ 23.723.63.518.529.22353Pyroxene73.373.43.368.178.1High-μ3.03.00.72.04.8NWA 6963 $Maskelynite$ 19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High-μ1.91.90.50.72.8LAR 12095 $Maskelynite$ 15.915.83.211.421.02753Pyroxene66.065.68.652.376.0 $Megacrysts$ 17.317.710.76.733.8High-μ0.80.80.40.41.6 IAR 2428 $Pyroxene$ 62.162.14.057.067.5Megacrysts27.327.34.521.832.7 $IIgh-μ$ 0.60.60.20.41.1EETA 79001Maskelwaite12.014.14.09.110.22002	Maskelynite	14.0	14.0	2.3	11.0	19.0	2453
High-μ2.32.30.51.33.1NWA 13134Maskelynite23.723.63.518.529.22353Pyroxene73.373.43.368.178.1High-μ3.03.00.72.04.8NWA 6963Maskelynite19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High-μ1.91.90.50.72.8LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High-μ0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High-μ0.60.60.20.41.1EETA 79001Maskelynite12.014.14.09.410.22002	Pyroxene	83.7	83.7	2.3	78.7	86.9	
NWA 13134Maskelynite23.723.63.518.529.22353Pyroxene73.373.43.368.178.1High- μ 3.03.00.72.04.8NWA 6963 </td <td>High-µ</td> <td>2.3</td> <td>2.3</td> <td>0.5</td> <td>1.3</td> <td>3.1</td> <td></td>	High-µ	2.3	2.3	0.5	1.3	3.1	
Maskelynite23.723.63.518.529.22353Pyroxene73.373.43.368.178.1High- μ 3.03.00.72.04.8NWA 6963Nmaskelynite19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High- μ 1.91.90.50.72.8LAR 12095Naskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.076.0Megacrysts17.317.710.76.733.816.6LAR 12011Naskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.52428Pyroxene62.162.14.051.410.22003High- μ 0.60.60.20.41.12428Pyroxene62.162.14.057.067.52428Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.52428Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.52023Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.0 <td< td=""><td>NWA 13134</td><td></td><td></td><td></td><td></td><td></td><td></td></td<>	NWA 13134						
Pyroxene73.373.43.368.178.1High-μ3.03.00.72.04.8NWA 6963 $Maskelynite$ 19.920.13.814.528.42653Pyroxene78.278.03.670.383.711.9High-μ1.91.90.50.72.8LAR 12095 $Maskelynite$ 15.915.83.211.421.02753Pyroxene66.065.68.652.376.076.0Megacrysts17.317.710.76.733.814.22428Pyroxene62.162.14.057.067.52428Pyroxene62.162.14.057.067.567.5Megacrysts27.327.34.521.832.711.1EETA 79001Maskelynite12.014.14.09.110.22002	Maskelynite	23.7	23.6	3.5	18.5	29.2	2353
High- μ 3.03.00.72.04.8NWA 6963Maskelynite19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High- μ 1.91.90.50.72.8LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001Maskelwrite12.014.14.08.110.22002	Pyroxene	73.3	73.4	3.3	68.1	78.1	
NWA 6963Maskelynite19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High- μ 1.91.90.50.72.8LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 7900120022002	High-µ	3.0	3.0	0.7	2.0	4.8	
Maskelynite19.920.13.814.528.42653Pyroxene78.278.03.670.383.7High- μ 1.91.90.50.72.8LAR 12095Naskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Naskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.521.832.7High- μ 0.60.60.20.41.121.120.2EETA 79001Maskelynite12.014.14.09.110.22002	NWA 6963						
Pyroxene78.278.03.670.383.7High-μ1.91.90.50.72.8LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High-μ0.80.80.40.41.6LAR 12011Μaskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.54.521.832.7High-μ0.60.60.20.41.120022002	Maskelynite	19.9	20.1	3.8	14.5	28.4	2653
High- μ 1.91.90.50.72.8LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.54.521.832.7High- μ 0.60.60.20.41.120022002	Pyroxene	78.2	78.0	3.6	70.3	83.7	
LAR 12095Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001	High-μ	1.9	1.9	0.5	0.7	2.8	
Maskelynite15.915.83.211.421.02753Pyroxene66.065.68.652.376.0Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001	LAR 12095						
Pyroxene 66.0 65.6 8.6 52.3 76.0 Megacrysts 17.3 17.7 10.7 6.7 33.8 High- μ 0.8 0.8 0.4 0.4 1.6 LAR 12011Maskelynite 10.0 9.9 1.6 7.7 14.2 2428 Pyroxene 62.1 62.1 4.0 57.0 67.5 Megacrysts 27.3 27.3 4.5 21.8 32.7 High- μ 0.6 0.6 0.2 0.4 1.1 EETA 79001	Maskelynite	15.9	15.8	3.2	11.4	21.0	2753
Megacrysts17.317.710.76.733.8High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001	Pyroxene	66.0	65.6	8.6	52.3	76.0	
High- μ 0.80.80.40.41.6LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001Meakelymite12.014.14.09.110.22002	Megacrysts	17.3	17.7	10.7	6.7	33.8	
LAR 12011Maskelynite10.09.91.67.714.22428Pyroxene62.162.14.057.067.5Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 79001Maskelymite12.014.14.09.110.22002	High-µ	0.8	0.8	0.4	0.4	1.6	
Maskelynite10.09.91.67.714.22428Pyroxene 62.1 62.1 4.0 57.0 67.5 Megacrysts 27.3 27.3 4.5 21.8 32.7 High- μ 0.6 0.6 0.2 0.4 1.1 EETA 79001 Meakelynite 12.0 14.1 4.0 9.1 10.2 2002	LAR 12011						
Pyroxene 62.1 62.1 4.0 57.0 67.5 Megacrysts 27.3 27.3 4.5 21.8 32.7 High- μ 0.6 0.6 0.2 0.4 1.1 EETA 79001Maskalumita 12.0 14.1 4.0 9.1 10.2 2002	Maskelynite	10.0	9.9	1.6	7.7	14.2	2428
Megacrysts27.327.34.521.832.7High- μ 0.60.60.20.41.1EETA 7900114.14.09.110.22002	Pyroxene	62.1	62.1	4.0	57.0	67.5	
High- μ 0.6 0.6 0.2 0.4 1.1 EETA 79001 Maskedwrite 12.0 14.1 4.0 9.1 10.2 2002	Megacrysts	27.3	27.3	4.5	21.8	32.7	
EETA 79001 Maakadamita 12.0 14.1 4.0 9.1 10.2 2002	High-µ	0.6	0.6	0.2	0.4	1.1	
Maskalumite 12.0 14.1 4.0 9.1 10.2 2002	EETA 79001						
Maskelymle 15.9 14.1 4.0 8.1 19.2 2003	Maskelynite	13.9	14.1	4.0	8.1	19.2	2003
Pyroxene 72.2 72.7 11.2 58.4 83.2	Pyroxene	72.2	72.7	11.2	58.4	83.2	
Megacrysts 13.5 12.8 13.5 0.0 30.9	Megacrysts	13.5	12.8	13.5	0.0	30.9	
High-μ 0.4 0.4 0.3 0.0 1.0	High-µ	0.4	0.4	0.3	0.0	1.0	
Tissint	Tissint						
Maskelynite 13.2 13.3 2.1 10.8 18.1 2333	Maskelynite	13.2	13.3	2.1	10.8	18.1	2333
Pyroxene 60.2 60.5 6.5 49.5 70.7	Pyroxene	60.2	60.5	6.5	49.5	70.7	
Megacrysts 25.9 25.5 7.8 13.2 38.5	Megacrysts	25.9	25.5	7.8	13.2	38.5	
High-μ 0.7 0.7 0.2 0.3 1.1	High-µ	0.7	0.7	0.2	0.3	1.1	
RBT 04261	RBT 04261						
Maskelynite 20.6 21.0 3.5 13.2 26.6 3603	Maskelynite	20.6	21.0	3.5	13.2	26.6	3603
Pyroxene 24.8 23.9 9.5 11.1 37.1	Pyroxene	24.8	23.9	9.5	11.1	37.1	
Olivine 53.6 54.3 8.3 44.9 66.0	Olivine	53.6	54.3	8.3	44.9	66.0	
High-μ 1.0 1.0 0.3 0.6 1.7	High-µ	1.0	1.0	0.3	0.6	1.7	

	Somplo	Pyroxen	Count			
к 	Sample	Average	1sd	Min	Max	Count
2	Zagami NWA	2.3	1.2	0.6	6.7	121
1	3134	6.4	2.4	1.8	13.1	110
1	NWA 6963	5.8	2.4	0.7	13.0	117
46						

1045
Table 3: Length of pyroxene long axis from Blob3D analysis

Sample/Phase	K	1σ	С	1σ
Zagami				_
Maskelynite	0.96	0.13	0.71	0.04
Pyroxene	0.51	0.09	0.49	0.03
High-µ	0.44	0.08	0.57	0.03
NWA 13134				
Maskelynite	0.54	0.10	0.57	0.03
Pyroxene	0.25	0.05	0.65	0.03
High-µ	0.42	0.09	0.46	0.03
NWA 6963				
Maskelynite	1.26	0.17	0.57	0.03
Pyroxene	0.93	0.12	0.86	0.04
High-µ	0.28	0.06	0.61	0.04
LAR 12095				
Maskelynite	0.23	0.08	0.40	0.03
Pyroxene	0.13	0.05	0.38	0.02
Olivine	0.91	0.10	0.70	0.03
High-µ	0.19	0.08	0.37	0.03
LAR 12011				
Maskelynite	0.96	0.15	0.64	0.04
Pyroxene	0.12	0.05	0.57	0.03
Olivine	0.20	0.05	1.26	0.06
High-µ	0.15	0.06	0.50	0.04
EETA 79001				
Maskelynite	1.38	0.46	0.28	0.03
Pyroxene	0.42	0.13	0.19	0.01
High-µ	0.22	0.06	0.34	0.02
Tissint				
Maskelynite	0.62	0.14	0.43	0.04
Pyroxene	0.29	0.09	0.31	0.02
Olivine	1.01	0.17	0.54	0.03
High-µ	1.06	0.17	0.37	0.03
RBT 04261				
Maskelynite	0.46	0.09	0.56	0.02
Olivine	0.56	0.07	0.44	0.03
High-µ	0.21	0.07	0.41	0.02

Table 4: Continuum CT fabric shape parameters, K and C, from Quant3D analysis1050

1053 **Table 5:** Angle between CT fabric orientations (°)

Sample	Maskelynite - Pyroxene	Maskelynite - Olivine	Maskelynite - High-µ	Pyroxene - Olivine	Pyroxene - High-µ	Olivine - High-µ
Zagami	8.6		2.0		10.5	
NWA 13134	11.3		16.9		23.4	
NWA 6963	8.7		7.5		4.8	
LAR 12095	4.9	6.5	8.0	9.2	11.6	6.8
LAR 12011	8.9	12.7	14.3	5.9	6.6	6.4
Tissint	12.2	20.4	12.9	11.7	2.8	11.8
EETA 79001	22.9		18.6		8.9	
RBT 04261		8.6	8.3			9.6

Table 6: High-μ phase crystal size distribution analysis results

Sample	# of Grains	Measured Volume (mm ³)	Average Length (mm)	±1 SD (mm)	BoxA/BoxC	±1 SD	Bin Interval	Slope (mm ⁻¹)	Y-intercept	R ²
Zagami	16336	858	0.12	0.11	2.41	0.92	0.04 - 1.34	-7.95	5.18	0.98
NWA 13134	17357	4564	0.26	0.25	2.18	0.79	0.07 - 3.48	-3.27	2.59	0.97
NWA 6963	17639	3335	0.22	0.22	2.32	0.85	0.06 - 2.87	-3.86	3.14	0.97
LAR 12095	23877	322	0.05	0.04	1.73	0.51	0.03 - 0.52	-21.16	7.60	0.97
LAR 12011	12348	1039	0.06	0.03	2.03	0.86	0.04 - 0.32	-33.51	7.23	0.99
EETA 79001	3874	450	0.09	0.07	1.92	0.69	0.06 - 0.76	-11.19	4.74	0.94
Tissint	41028	705	0.07	0.05	1.83	0.60	0.04 - 0.63	-18.59	7.51	0.99
RBT 04261	105233	11348	0.13	0.08	1.67	0.61	0.06 - 1.01	-11.77	5.31	1.00















1084 Figure 6









1092 Figure 9 1093 0% megacryst olivine 31% megacryst olivine 1000 100



1095 Figure 101096



1097

- 1098 1099
- 1099
- 1100 1101
- 1101

FIGURE CAPTIONS

- **Figure 1**: Grayscale CT slices for the olivine-phyric (a d), basaltic (e f), gabbroic (g), and poikilitic (h) shergottites in this study. The main phases are maskelynite, pyroxene, olivine (in
- 1105 olivine-phyric and poikilitic samples), and oxides/sulfides in order of increasing brightness.
- 1106 Scale bars are 1 mm.
- 1107 Figure 2: 2D schematic showing the Quant3D workflow. (A) Grayscale CT data are segmented
- 1108 into their constituent phases (B). (C) Inset from (B) showing points randomly placed through
- 1109 phase-of-interest (i.e., white material) and cones emanating from each point along a set number
- 1110 of orientations. (D) The volume of each cone along every orientation is combined to create a
- 1111 tensor ellipsoid, from which eigenvectors and eigenvalues can be derived. The data can also be 1112 visually assessed with a 3D rose diagram. This process is shown here in 2D but is actually
- 1112 visually assessed with 1113 performed in 3D.
- 1114 **Figure 3**: (A) 3D volume rendering of an elongated pyroxene crystal (transparent yellow) in
- 1115 NWA 13134 (10.8 mm x 0.8 mm x 0.5 mm) and a rod (light blue) drawn to represent the length
- 1116 and orientation of pyroxene crystal. A colored block illustrates 3 orthogonal orientations. (B) CT

- slice oriented orthogonally to the long axis of the same crystal highlighted in yellow (red
- 1118 orientation on block). (C and D) CT slices oriented along the long axes of the same crystal
- 1119 highlighted in yellow (blue and green orientations on block). Scale bars are 1 mm.
- 1120 Figure 4: Whole-rock and slice-by-slice modal abundances for the various phases within
- 1121 shergottites.
- 1122 Figure 5: Plot of CT fabric eigenvalue ratios after Woodcock and Naylor (1983) for different
- 1123 shergottite phases measured with Quant3D. Foliated fabrics are defined as K < 1, lineations
- 1124 defined as K > 1, triaxial defined as K = 1. Higher values of C correspond to stronger fabrics.
- 1125 Error bars represent 1σ standard deviation of 30 replicate analyses.
- 1126 Figure 6: Stereonets (lower hemisphere; equal area projection) showing the eigenvector
- 1127 orientations for each phase measured using Quant3D and the average orientation of high-µ
- 1128 phases measured with Blob3D (gray data). Each Quant3D data point represents one of the thirty
- replicate analyses. In general, a tight cluster of the third or first eigenvector, and a spread of the
- 1130 other two around a great circle, reflects a foliation or lineation, respectively. The high-µ Blob3D
- 1131 axes are not orthogonal because the average vector for each axis was calculated individually
- 1132 using a Bingham axial distribution (Fisher et al., 1993).
- 1133 Figure 7: Contoured stereonets (lower hemisphere; equal area projection) showing the
- 1134 orientation of individual pyroxene long axes. Each black dot represents a single crystal. The
- 1135 density of datapoints is reported in percent per 1% of stereonet area. Darker contours indicate a
- 1136 higher density of datapoints. Yellow lines represent foliation plane. See Figure 5 for meaning of
- 1137 K and C.
- **Figure 8**: Crystal size distribution (CSD) patterns for shergottite high-µ phases. The same bin
- sizes were used for all samples but the X- and Y-axes are scaled differently for each group. The
- 1140 gray columns represent data from bin sizes less than the turndown bin and are excluded from
- 1141 interpretations.
- 1142 **Figure 9**: CT slices with the minimum (left) and maximum (right) megacryst olivine or pyroxene
- 1143 oikocryst abundances for olivine-phyric and poikilitic shergottites, respectively. Scale bars are 2 1144 mm.
- 1145 **Figure 10:** (column A) 3D image showing whole-rock (transparent blue) with rods (magenta)
- that represent long axis of individual pyroxene crystals in NWA 6963, NWA 13134, and
- 1147 Zagami. (column B) CT slice oriented parallel to foliation plane (yellow line in [a]). (columns C
- and D) are CT slices oriented orthogonally to (B). All scale bars are 1 cm.