

1 **Revision 1**

2 **Quasicrystals at extreme conditions: The role of pressure in stabilizing icosahedral**
3 **Al₆₃Cu₂₄Fe₁₃ at high temperature**

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21

22 **Abstract**

23 Icosahedrite, the first natural quasicrystal with composition $\text{Al}_{63}\text{Cu}_{24}\text{Fe}_{13}$, was discovered in
24 several grains of the Khatyrka meteorite, a CV3 carbonaceous chondrite. The presence of
25 icosahedrite associated with high-pressure phases like ahrensite and stishovite indicates
26 formation at high pressures and temperatures due to an impact-induced shock. Previous
27 experimental studies on the stability of synthetic icosahedral AlCuFe have either been limited to
28 ambient pressure, for which they indicate incongruent melting at ~ 1123 K, or limited to room
29 temperature, for which they indicate structural stability up to about 35 GPa. These data are
30 insufficient to experimentally constrain the formation and stability of icosahedrite under the
31 conditions of high pressure and temperature that formed the Khatyrka meteorite. Here we present
32 the results of room-temperature, high pressure diamond anvil cells measurements of the
33 compressional behavior of synthetic icosahedrite up to ~ 50 GPa. High P - T experiments were
34 also carried out using both laser-heated diamond anvil cells combined with in situ synchrotron
35 X-ray diffraction (at ~ 42 GPa) and multi-anvil apparatus (at 21 GPa) to investigate the structural
36 evolution and crystallization of possible coexisting phases. The results demonstrate that the
37 quasiperiodic order of icosahedrite is retained over the P - T range explored. We find that pressure
38 acts to stabilize the icosahedral symmetry at temperatures much higher than previously reported.
39 Direct solidification of AlCuFe quasicrystals from an unusual Al-Cu-rich melt is possible but it
40 is limited to a narrow temperature range. Alternatively, quasicrystals may form after
41 crystallization through solid-solid reactions of Al-rich phases. In either case, our results show
42 that quasicrystals can preserve their structure even after hypervelocity impacts spanning a broad
43 range of pressures and temperatures.

44 **Keywords:** Icosahedrite, Quasicrystals, CV3 chondrite, redox, Khatyrka meteorite, solar nebula.

45 **Introduction**

46 Quasicrystals (QC; Levine and Steinhardt 1984; Shechtman et al. 1984) represent a class
47 of solids characterized by quasiperiodic translational order and crystallographically-forbidden
48 rotational symmetries and are now observed in nature (Bindi et al. 2009). These symmetries
49 include the icosahedral (i) symmetry exhibited by $\text{Al}_x\text{Cu}_y\text{Fe}_z$ alloys, where x varies between 61
50 and 64, y is 24-26 and z 12-13 atomic% (Bancel 1999). This chemical interval corresponds to the
51 compositional range at which the i-QC solely is stable up to ~ 1023 K at ambient pressure. Above
52 this temperature the stability field of the quasicrystal decreases to a very narrow chemical
53 composition up to ~ 1123 K, where the i-QC with composition $\text{Al}_{62.5}\text{Cu}_{25}\text{Fe}_{12.5}$ coexists with a
54 liquid + (λ) monoclinic phase with composition $\text{Al}_{13}\text{Fe}_4$ (Tsai 2013; Zhan and Lück 2003a-e). At
55 temperatures above ~ 1200 K, i-QC has been shown to be unstable, such that the liquid only
56 coexists with the λ phase. At approximately 1300 K the system is totally molten and the liquid
57 composition reflects that of the starting QC.

58 Experimental studies of the phase relationships at ambient pressure have been the only
59 available tool to date to constrain the origin of the first natural quasicrystal, icosahedrite
60 $\text{Al}_{63}\text{Cu}_{24}\text{Fe}_{13}$ (Bindi et al. 2009, 2011). However, textural and petrographic evidence suggest that
61 the natural quasicrystal formed in outer space under pressures and temperatures considerably
62 higher than 1 bar and 1300 K (Bindi et al. 2012; Hollister et al. 2014). Recently, Stagno et al.
63 (2014) performed *in situ* high P - T X-ray diffraction studies and showed that the icosahedral
64 symmetry of the AlCuFe QC is retained at 5 GPa and temperatures up to 1673 K. Above this
65 temperature the synthetic icosahedrite used for the experiments was found to decompose to a
66 liquid in equilibrium with CuAl (corresponding to the mineral cupalite, an accessory phase also
67 found in the Khatyrka meteorite), and the cubic β phase (Bindi et al. 2011). Although this study

68 provided information on the compressional behavior of i-AlCuFe QC, the results gave only a
69 lower-bound on the P - T stability of natural icosahedrite. Previous studies that focused on the
70 compressional behavior and structural stability of QCs with compositions slightly different from
71 that of icosahedrite include measurements on i-Al₆₂Cu_{25.5}Fe_{12.5}, which was shown to be stable up
72 to 35 GPa (Sadoc et al. 1994, 1995; Lefebvre et al. 1995).

73 Here we present experimental results using both diamond anvil cell (DAC) and multi-
74 anvil techniques to investigate the stability of synthetic i-AlCuFe at higher pressures and
75 temperatures than those reported in previous studies. We show that the icosahedral structure is
76 stabilized at high temperature as pressure increases, which makes quasicrystal behavior similar
77 to that of most crystalline materials exposed to similar extreme conditions. Our results indicate
78 that icosahedrite could have formed within a large range of pressures and temperatures during
79 the formation of our solar system provided extremely reducing conditions. Hence, Khatyrka
80 might represent one of many QC-bearing meteorites in our solar system or elsewhere in the
81 cosmos.

82 **Experimental Methods**

83 The synthetic icosahedral quasicrystal used as starting material was characterized by
84 SEM and XRD measurements and shown to have the formula Al₆₃Cu₂₄Fe₁₃ (Bancel 1999) plus
85 minor amounts of cupalite, (Cu,Fe)Al. The synthetic material was first broken in several small
86 fragments. A small chip was, then, loaded in a diamond anvil cell with culet size of 600 μ m and
87 crushed to a fine powder by hand loading. A small portion of the powder with dimensions of
88 about 30 \times 30 μ m was picked with a needle and placed at the center of a symmetric diamond
89 anvil cell with 300 μ m culet size using a Re gasket as sample chamber with a 150 μ m diameter

90 hole. A couple of ruby spheres were placed next to the sample as pressure markers. One diamond
91 anvil was supported by a cubic boron nitride (c-BN) backing plate, and the other anvil by a
92 tungsten carbide (WC) backing plate. *In situ* angle-dispersive powder X-ray diffraction
93 measurements were performed at high pressure at the 16BM-D beamline, HPCAT (Advanced
94 Photon Source, APS, Argonne National Laboratory, ANL). The DAC was loaded with Ne, which
95 served as both a pressure medium and a pressure marker (Hemley et al. 1989), and then mounted
96 on a motor driven gearbox with the WC seat on the downstream side, and the c-BN seat on the
97 upstream side. Sample pressures were measured with the ruby luminescence method (Mao et al.
98 1986) through an on-line system. Monochromatic incident X-ray beams with wavelengths of
99 0.42460 Å and 0.5166 Å were used. The beam was focused to a spot of $5 \times 15 \mu\text{m}$ by using a
100 pair of Kirkpatrick-Baez mirrors. The MAR345 image plate detector was placed at a distance
101 approximately 478 mm from the sample in order to obtain high resolution and accuracy of the
102 Debye-Scherrer diffraction rings. Diffraction peaks were collected using a continuous ω -
103 oscillation scan mode over the range from -6° to $+6^\circ$ with an exposure time of 180 seconds.

104 Simultaneous high-pressure and temperature synchrotron powder X-ray diffraction
105 experiments were conducted at the 13ID-D beamline, GSECARS (APS, ANL) using a focused
106 monochromatic 30 keV X-ray beam with wavelength 0.4133 Å. Double-sided laser heating was
107 performed using two infrared laser beams focused to $15 \mu\text{m}$ flat-top spots on both sides of the
108 sample co-axially aligned with two optical paths for temperature measurements and visually
109 aligned with focused $4 \mu\text{m}$ X-ray beam using the X-ray induced luminescence of the sample
110 (Prakapenka et al. 2008). Laser power was adjusted independently on upstream and downstream
111 sides to control the sample temperature within $\pm 100\text{K}$. The target temperature was maintained
112 for about 10 min. Temperatures of the laser-heated sample were measured using thermal

113 radiation spectra fitted to the blackbody radiation function. Diffraction patterns were collected on
114 a MarCCD-165 detector with exposure time of 15 s. In these experiments, chips of synthetic
115 $\text{Al}_{63}\text{Cu}_{24}\text{Fe}_{13}$ were also crushed and then slightly pressed to form a platelet. A small platelet with
116 a diameter of approximately 60 μm was loaded into the sample chamber of a symmetric
117 diamond-anvil cell with flat anvils of 300 μm -size culet. Pressure was measured using the
118 thermal equation of state of Ne used as pressure medium. The *in situ* X-ray diffraction patterns
119 were processed using FIT2D software (Hammersley 1998), and the *d*-spacing relative to each
120 reflection was determined using PeakFit software.

121 Quench experiments were performed at 21 GPa and temperature between 1600 K and
122 2000 K using a 1500-ton Walker-type press available at the Carnegie Institution of Washington.
123 The starting material used in this study was a synthetic icosahedral AlCuFe ($\geq 99.9\%$)
124 quasicrystalline powder with nominal composition of $\text{Al}_{65}\text{Cu}_{23}\text{Fe}_{12}$, according to a previous
125 study (Stagno et al. 2014). Tungsten carbide anvils of 3 mm truncation edge length (TEL) were
126 used with 8 mm edge length MgO pressure media. Graphite and alumina capsules were used in
127 the attempt to prevent oxidation of the starting material. The capsules were then placed in the
128 central portion of a cylindrical Re furnace, surrounded by MgO sleeve and spacers. A LaCrO_3
129 sleeve was used as thermal insulator outside the heater.

130 Details of the pressure calibration of this type of assembly have been reported by Hirose
131 and Fei (2002). The temperature during the experiment at 1973 K was monitored with a W-
132 5%Re/W-26%Re (C-type) thermocouple inserted within an alumina sleeve, with the junction in
133 contact with the top of the capsule. From this run, a temperature versus power calibration curve
134 was obtained that was used for the additional runs. The sample was compressed to the target
135 pressure at a rate of 0.5 GPa/hr, and then heated to the target temperature and kept manually

136 constant within 10 K for a period of 15-60 minutes. The sample was quenched by turning off the
137 power to the furnace and, then, decompressed to ambient pressure.

138 All recovered samples from quench experiments were mounted in epoxy resin and
139 polished parallel to the axial furnace direction for textural observation and chemical composition
140 mapping by Field Emission Scanning Electron Microscope (JEOL JSM 6500F). Semi-
141 quantitative analyses using energy-dispersive X-ray spectroscopy, were performed at 15 kV and
142 1.1 nA using metals (Fe, Cu, Al,) and oxides (Al_2O_3) as standards. Phase identification of the
143 selected recovered run products was accomplished using an Oxford Diffraction Xcalibur PX
144 Ultra single-crystal diffractometer fitted with a 165 mm diagonal Onyx CCD detector (CuK α
145 radiation). The crystal-to-detector distance was 7 cm. Data were processed using the *CrysAlis*
146 software package version 1.171.31.2 (Oxford diffraction) running on the Xcalibur PX control
147 PC.

148

149 **Results and Discussion**

150 Compression behavior

151 Synthetic icosahedral quasicrystals with the formula $Al_{63}Cu_{24}Fe_{13}$ (Bancel 1999) were
152 used as starting materials for our *in situ* DAC experiments. An accurate textural and chemical
153 analyses of the synthetic material showed minor amounts of (Cu, Fe)Al. The first set of
154 experiments consisted of *in situ* powder angle-dispersive X-ray diffraction measurements on i-
155 QC using DACs up to ~50 GPa at room temperature. These experiments aimed to investigate the
156 evolution of the icosahedral structure under pressure, the determination of the lattice parameter
157 and the equation of state. A total of 50 diffraction patterns were collected during compression

158 and decompression. Figure 1 shows a characteristic spotty diffraction pattern constantly observed
159 during our experiments and resulting from heterogeneous size of the QC grains.

160 Figure 2 shows the variation of the d -spacing for 7 known diffraction peaks with
161 increasing pressure and after decompressing the sample to ambient pressure. The intensities of
162 the peaks appear strongly affected by the preferred orientation of the powder grains as can be
163 observed in Figure 1. The shift in d -spacing with increasing pressure can also be seen for most of
164 the peaks up to the target pressure. In addition, the peak broadening is apparent as the pressure
165 increases and can be attributed to an increase of the mosaic spread and local strains. After the
166 sample was decompressed to ambient pressure, peaks were still broadened in agreement with
167 what was reported by Sadoc et al. (1994) for $i\text{-Al}_{62}\text{Cu}_{25.5}\text{Fe}_{12.5}$.

168 We collected additional diffraction patterns at higher resolution by moving the detector to
169 a farther distance from the sample (~ 478 mm). This allowed us to investigate more accurately the
170 icosahedral structure, in particular the behavior in the high d -spacing region that included peaks
171 (12,16) at 5.53 Å and (8,4) at 8.94 Å, respectively [for indexing notation see Lu et al. (2001)].
172 Diffraction patterns collected up to 36 GPa clearly show a gradual broadening with increasing
173 pressure (Figure S1). Two additional peaks belonging to $i\text{-AlCuFe}$ were also observed with d -
174 spacings of ~ 6 Å and 9.7 Å that could be indexed as (24,15) and (6,3), respectively, using the
175 automated identification scheme described by Lu et al. (2001), or may be reflections due to
176 a minor unidentified phase. Our *in situ* X-ray diffraction measurements show that no peaks
177 appear or disappear up to the target pressure of ~ 50 GPa, which excludes possible pressure-
178 induced phase transformations, including amorphization, that has been found to occur for i -
179 AlLiCu quasicrystals (Itie et al. 1996). The observed peak broadening with pressure can be

180 interpreted as arising from the increasing atomic disorder, perhaps due to residual stress, without
181 changing the long-range quasicrystalline order.

182 We determined the pressure dependence of the lattice parameter a_{6D} up to the maximum
183 pressure of ~50 GPa (see Table S1 of Supporting Information). The parameter is defined as,

184

$$185 \quad a_{6D} = d \sqrt{\frac{N + M\tau}{2(2 + \tau)}} \quad (1),$$

186

187 where d is the d -spacing in Å, N and M the Cahn indices for which the d -spacing is
188 experimentally determined, and τ is the golden ratio, $(1+\sqrt{5})/2$ (Steurer and Deloudi 2009). The
189 six-dimensional lattice parameter is shown to gradually decrease with increasing pressure (Fig.
190 3). The slightly scattered data at 24-32 GPa is likely due to the less accurate pressure
191 determination caused by the overlap between the (111) peak of Ne and the (80, 128) peak of i-
192 AlCuFe. However, the lattice parameter calculated from compression experiments is in good
193 agreement with that calculated on decompression. The estimated reduction of the lattice
194 parameter from the ambient pressure value of 12.64 Å is about 8%. In addition, our a_{6D} value at
195 5 GPa and room temperature is consistent with that determined by Stagno et al. (2014) at similar
196 conditions. We conclude that the QC retains its icosahedral structure over the pressure range
197 investigated. Further studies are needed to distinguish whether the stability is thermodynamic or
198 kinetic.

199 The zero pressure bulk modulus K_0 and its pressure derivative K_0' were determined from
200 the least-squares fit to several equation of state (EOS) models. Fit to the first-order Murnaghan
201 EOS fit (Angel et al. 2014), which allows direct comparison with the results of previous studies,
202 resulted in $K_0 = 113.7(\pm 2.9)$ and $K_0' = 4.22(\pm 0.22)$. It can be seen from Figure 4 that K_0 and K_0'

203 obtained from our data are significantly lower and higher, respectively, than those obtained for
204 $\text{Al}_{62}\text{Cu}_{25.5}\text{Fe}_{12.5}$ from previous authors using the same EOS, i.e., $K_0 = 139(\pm 6)$ GPa and $K_0' = 2.7$
205 (Sadoc et al. 1994), and $K_0 = 155(\pm 10)$ GPa and $K_0' = 2$ (Lefebvre et al. 1995). Our EOS
206 parameters also differ from those determined for an approximant phase with composition
207 $\text{Al}_{64}\text{Cu}_{24}\text{Fe}_{12}$ that is close to our synthetic i-QC [$K_0 = 175(\pm 16)$ GPa and $K_0' = 2.00$ (Lefebvre et
208 al. 1995)]. Such differences in the compressional behavior can be interpreted as due to either
209 different composition of the QCs or distinct mechanical properties of the approximant
210 (crystalline) phase. It should be kept in mind, however, that in previous studies in which DACs
211 techniques were employed, silicon oil was used as a pressure medium, for which hydrostaticity is
212 limited to very low pressure (Angel et al. 2007). Moreover, the previously suggested EOSs have
213 been derived from data collected using energy-dispersive rather than angle-dispersive X-ray
214 diffraction with the lattice parameter calculated using a different peak than the (8,4) used in this
215 study without taking into consideration possible anisotropy of the material.

216 Our experimental data were also fit using both a third-order Birch-Murnaghan EOS [$K_0 =$
217 $110.4(\pm 2.9)$ and $K_0' = 4.79(\pm 0.28)$] and a Vinet et al. EOS [$K_0 = 109.4(\pm 2.9)$ and $K_0' =$
218 $5.06(\pm 0.29)$]. Whereas the resulting parameters deviate slightly from the parameters obtained
219 using a simple Murnaghan model, they confirm the lower bulk modulus of synthetic icosahedrite
220 compared to literature data. Figure 4 also shows the compressional behavior for pure *fcc*-Al, *fcc*-
221 Cu (Dewaele et al. 2004) and *hcp*-Fe (Mao et al. 1990) plotted using the Vinet et al. and Birch-
222 Murnaghan EOS models. For all these pure metals the structure has been reported to be stable
223 over a wide pressure range > 100 GPa. As observed by Sadoc et al. (1994), the compressional
224 behavior of our synthetic icosahedrite is much closer to that of pure Cu, although Al represents
225 the main constituent. We therefore expect a similar compressional behavior for i-AlCuFe

226 quasicrystals varying in compositions according to the phase diagram proposed by Bancel
227 (1999).

228

229 High P - T stability of icosahedral symmetry

230 Recently, the conditions for the formation of natural i-AlCuFe have been constrained on
231 the basis of textural and chemical analyses of the coexisting mineral phases within the CV3-like
232 chondritic grains of the Khatyrka meteorite (Hollister et al. 2014; MacPherson et al. 2013). The
233 observation of rare Al-Cu-Fe alloys associated with melt droplets and phases such as stishovite
234 and ahrensite imply that the meteorite was subjected to a combination high pressures and
235 temperatures resulting from a high-velocity impact-induced shock. A study of icosahedrite at
236 high pressures and temperatures enables more definitive understanding of the petrological
237 processes that formed the Khatyrka meteorite. A first study by Stagno et al. (2014) indicated that
238 synthetic icosahedrite is stable at 5 GPa at temperatures up to ~ 1673 K; at higher temperatures,
239 the sample was found to melt incongruently and produce two solid crystalline phases: β -phase
240 and cupalite. These experiments provided the first indication that pressure might act to stabilize
241 the QC at T higher than 1300 K.

242 *In situ* high P - T laser heating system diamond-anvil cell experiments were conducted to
243 further constrain the stability field of icosahedrite. The synthetic quasicrystalline powder was
244 first compressed to ~ 42 GPa, then heated to ~ 1830 K while collecting X-ray diffraction patterns
245 to monitor any possible transformation or melting (see details in the Experimental Methods
246 session). The sample was then cooled down to 1000 K before being quenched. The results show
247 that during heating at about 1560 K the intensity of most of the peaks decreases dramatically,
248 and new peaks belonging to the QC phase appear at d -spacings between 1.8 and 2.0 Å (Fig. 5).

249 Interestingly, during cooling of the sample at about 1500 K the main peaks re-appear and are
250 visible even after quenching the sample to room T . No amorphization or phase transformation
251 was evident, and the new peaks that are characteristic of the QC become visible as a result of a
252 strong preferential orientation. However, with increasing T the loss of quasicrystallinity (i.e.,
253 formation of crystal approximants) via a reversible process cannot be excluded.

254 Because of the small size of the laser spot, it is possible to heat different points of the
255 sample within the DAC. The results of the high T run performed at the same pressure for a
256 different point of the sample is shown in Fig. S2. In this case the QC was heated up to ~ 2110 K
257 and then quenched directly to room temperature. The results are similar to the previous
258 experiment at 1600 K, where peaks characteristic of synthetic icosahedrite were present. At
259 higher T , new unknown peaks appear that are preserved also after quenching the sample to room
260 temperature suggesting that the i-QC might have decomposed irreversibly. However, subsequent
261 *ex situ* single crystal X-ray diffraction measurements on a micrometer-sized grain handpicked
262 from the cell (see Fig. S3 of Supporting Information) showed that, at least for that fragment of
263 the sample, the 5-fold symmetry characteristic of i-QC is retained. Given the difficulty to
264 establish whether or not the *ex situ* fragment was heated all the way up to the target temperature,
265 we cannot extend with confidence the ex-situ results to the bulk heated sample.

266 We performed additional quench experiments using the multi-anvil technique to better
267 understand the nature of the unknown peaks discussed above. Experiments were performed at 21
268 GPa and temperatures between 1673 and 1993 K for icosahedral AlCuFe ($\geq 99.9\%$)
269 quasicrystalline powder with nominal composition of $\text{Al}_{65}\text{Cu}_{23}\text{Fe}_{12}$ previously characterized
270 (Stagno et al. 2014). SEM images using back-scattered electrons of the run products reveal the
271 compositions and textures of the recovered material (Fig. 6). The run quenched from 1673 K

272 consists of a single phase with composition $\text{Al}_{64.11(\pm 0.66)}\text{Cu}_{24.70(\pm 0.74)}\text{Fe}_{11.19(\pm 0.22)}$, which is
273 consistent with the starting composition of the synthetic QC. Several grains showed a patchy
274 texture that we believe are due to the onset of melting of the QC. At about 1773 K the recovered
275 sample shows the coexistence of β -phase ($\text{Al}_{64.73(\pm 0.36)}\text{Cu}_{20.23(\pm 0.91)}\text{Fe}_{15.04(\pm 0.68)}$), Fe-rich cupalite
276 ($\text{Al}_{48.77(\pm 0.35)}\text{Cu}_{36.10(\pm 0.93)}\text{Fe}_{15.13(\pm 0.70)}$) and a phase identified by the patchy texture with
277 composition $\text{Al}_{62.65(\pm 0.89)}\text{Cu}_{33.52(\pm 1.08)}\text{Fe}_{2.92(\pm 0.27)}$ that can be interpreted either as a Fe-poor
278 khatyrkite (CuAl_2) or a melt. Small grains of Al_2O_3 are also present and suggest possible
279 oxidation of the material during the experiment. Finally, the recovered sample from 1973 K
280 appears totally molten consisting of a Fe-rich liquid with composition
281 $\text{Al}_{12.07(\pm 0.50)}\text{Cu}_{21.34(\pm 1.04)}\text{Fe}_{65.48(\pm 0.73)}$ and exhibiting a characteristic quench texture that includes
282 “skeletal” Al metal (Fig. 6d).

283 The results of quench experiments thus, suggest 1) that the QC retains its stability at 21
284 GPa and 1673 K and its icosahedral symmetry is retained after quench; 2) as temperature
285 increases isobarically icosahedrite might melt congruently to, then, dissociate in a liquid with
286 composition very similar to khatyrkite + β -phase + cupalite similar to what was reported by
287 Stagno et al. (2014); 3) pure Al is the first phase crystallizing from a molten liquid with
288 icosahedrite-like composition.

289

290 **Implications**

291 Our experimental results reproduce key features associated with the presence of
292 icosahedrite that have been observed in the Khatyrka meteorite in terms of the phase assemblage
293 (cupalite, β -phase, khatyrkite and pure Al) and texture. The coexistence of icosahedrite with a
294 liquid phase having the same composition (i.e. congruent melting) than the ideal composition

295 reported by Bancel (1999; i.e., $\text{Al}_{63}\text{Cu}_{24}\text{Fe}_{13}$) has been never reported at such high pressures and
296 implies that the i-QC is kinetically stable and perhaps thermodynamically stable at high pressure.
297 This would confirm the hypothesis by Hollister et al. (2014) that the Al, Cu-rich assemblage in
298 the Khatyrka meteorite formed after an impact-induced shock followed by rapid cooling. (10^2 -
299 10^3 °C s⁻¹) from which the Al, Cu-rich assemblage formed. However, we point out that a similar
300 mechanism of formation for icosahedrite appears unlikely as we assume here thermodynamic
301 equilibrium between the QC and the liquid from which it forms. In fact, the coexistence of Al,
302 Cu rich-phases with icosahedrite in the Khatyrka meteorite would suggest an initial high
303 abundance of these elements, as confirmed by the finding of Al-rich Cu-bearing FeNi and sulfide
304 phases in proximity of the QC. The finding of pure Al in our experiments is a further element of
305 similarity with the natural assemblage (see Fig. 2 in Hollister et al. 2014) and can be explained in
306 light of its lower melting temperature with respect to Fe and Cu that would trigger its
307 mobilization by diffusion mechanisms and exsolution. At 1773 K the presence of khatyrkite
308 coexisting with pure Al would represent a further evidence of the high-temperature regime
309 occurred during crystallization of these phases that would require an unusual abundance of Al.

310 To date, two hypotheses have been proposed (Hollister et al. 2014) to explain the origin
311 of Al-Cu-Fe alloys in the Khatyrka meteorite: 1) formation by an impact-induced shock event
312 that causes the diffusion of Al and Cu out of FeNi initially enriched with Al and Cu; or 2)
313 formation during nebular processes before the impact occurred and re-melting and re-
314 solidification of these alloys after the impact. In either case, the impact event enables
315 mechanisms such as diffusion, solid-solid reactions and oxidation-reduction. Although the
316 processes occurring in the meteorite cannot be precisely reproduced in laboratory experiments, it
317 is notable that our experiments lead to the same phases even at pressures and temperatures higher

318 than those experienced by the meteorite, for which Hollister et al. (2014) proposed pressures
319 somewhat greater than 5 GPa and temperatures around 1200 °C.

320 The results of this study can be summarized as follows, 1) synthetic icosahedrite was
321 shown to retain its structure up to ~50 GPa at ambient temperature; 2) it was experimentally
322 demonstrated for the first time that pressure can stabilize the icosahedral AlCuFe quasicrystal
323 until it melts (or decomposes) with no evidence of direct structural change to a crystalline
324 phases; 3) congruent melting of icosahedrite might be limited to a very narrow temperature
325 interval at 21 GPa and 1673 beyond which Al, Cu-rich phases would form. Based on our results
326 the preservation of icosahedrite over cosmic time scales in a meteorite that formed at the early
327 stage of the solar system results from its unexpected stability at high temperatures and pressures.
328 Therefore, the discovery of icosahedrite in other meteorites exposed to extreme conditions and
329 with bulk compositions similar Khatyrka should be expected.

330

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343

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- 428

429 **Figure Captions**

430 **Figure 1.** Representative X-ray diffraction pattern of $i\text{-Al}_{63}\text{Cu}_{24}\text{Fe}_{13}$ collected at $P = 18.5$ GPa.
431 Diffraction rings and weak single spots are typical features of the collected patterns. The Debye-
432 Scherrer rings are labeled as in case of QCs using the two-integer indexes by Janot (1994). The
433 diffraction patterns were processed using Fit2D (Hammersley 1998) and Peakfit software.

434

435 **Figure 2.** Pressure dependence of the powder X-ray diffraction patterns of $i\text{-QC}$ collected at
436 room temperature in angle-dispersive mode (wavelength of 0.4246 Å). Filled circles indicate
437 peaks of ruby (pressure marker), grey circles for Au (pressure marker), empty circles for Re
438 (gasket) filled triangles for Ne (pressure medium). The diffraction peaks were indexed using
439 Cahn indices (N, M) following the scheme proposed by Janot (1994; see also Steurer and
440 Deloudi 2009). The diffraction pattern (in gray) at ambient pressure is relative to the sample after
441 decompression.

442

443 **Figure 3.** Variation of the lattice parameter as function of pressure: yellow square, ambient
444 pressure value (Bindi et al. 2011); black and white squares indicate lattice parameter determined
445 respectively from compression and decompression experiments; gray squares indicate
446 experiments at higher resolution.

447

448 **Figure 4.** Pressure-volume relations for $i\text{-Al}_{64}\text{Cu}_{23}\text{Fe}_{13}$ with experimental data (black squares)
449 fitted using a Murnaghan EOS (dashed black line). Results from Birch-Murnaghan and Vinet et
450 al. EOS fits are here omitted since both closely overlap the Murnaghan EOS fit. Uncertainties are
451 within the symbol size. Our fit is compared with previous studies by Sadoc et al. (1994; gray

452 line) and Lefbvre et al. (1995; blue line) both for icosahedral $\text{Al}_{62}\text{Cu}_{25.5}\text{Fe}_{12.5}$ and for one
453 approximant phase with composition $\text{Al}_{64}\text{Cu}_{24}\text{Fe}_{12}$ (Lefbvre et al. 1995; yellow line). The two
454 curves obtained from these previous studies are extrapolated up to 52 GPa assuming that no
455 structural phase transformation occurs. The EOS of pure Al, Cu (Dewaele et al. 2004) and Fe
456 (Mao et al. 1990) are also reported with green, blue and brown, respectively.

457

458 **Figure 5.** Representative angle-dispersive (wavelength of 0.4133 Å) X-ray diffraction patterns
459 for synthetic icosahedrite as function of temperature at ~ 42 GPa. Diffraction patterns were
460 collected with 15s exposure time after keeping the sample at approximately constant T for 180s.
461 Peaks are indexed as mentioned above. Filled black triangles indicate the (111) and (200) peaks
462 of Ne pressure medium. The figure demonstrates the stability of i-QC during heating up to about
463 1830 K followed by slow cooling down to 1000 K before quench.

464

465 **Figure 6.** Back-scattered electron (BSE) images of the recovered sample from runs at 21 GPa. a)
466 Run at 1673 K showing the presence of i-QC also confirmed by the five-fold symmetry in (b)
467 using single crystal X-ray diffraction. (c) Recovered sample from 21 GPa and 1773 K and d)
468 1973 K, respectively.

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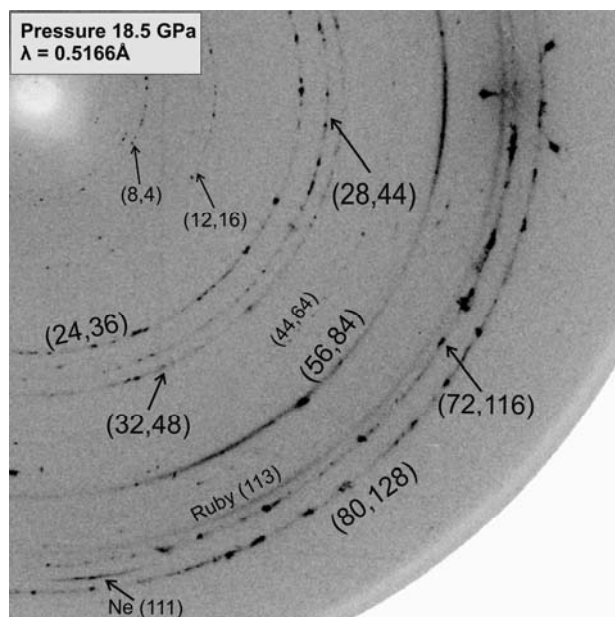
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474 **Figures**

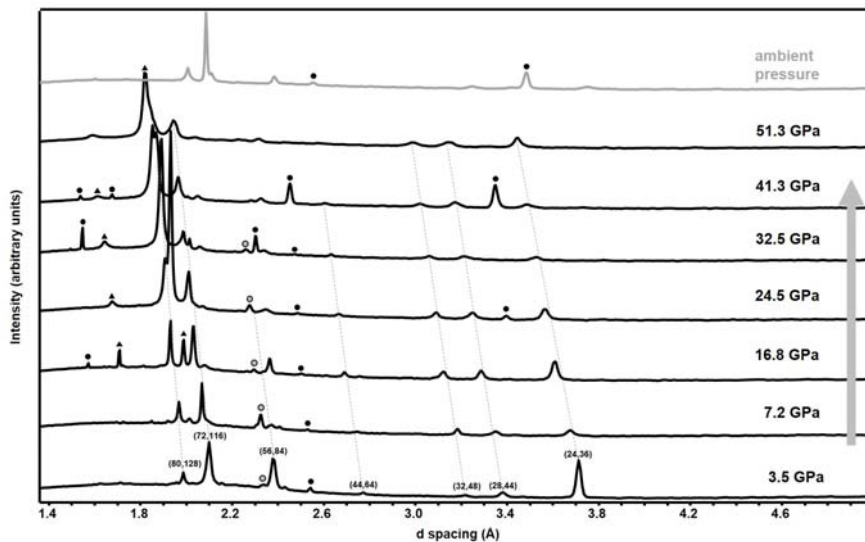


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Figure 1

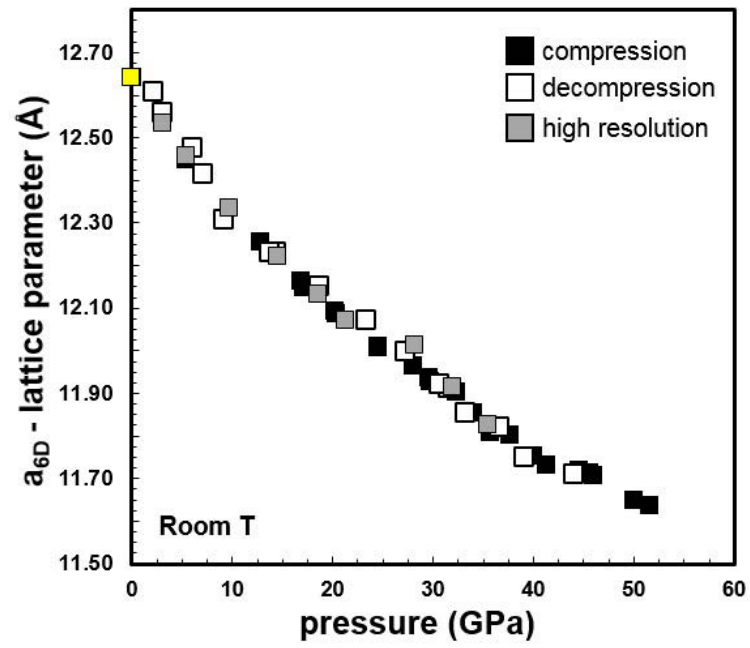


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Figure 2

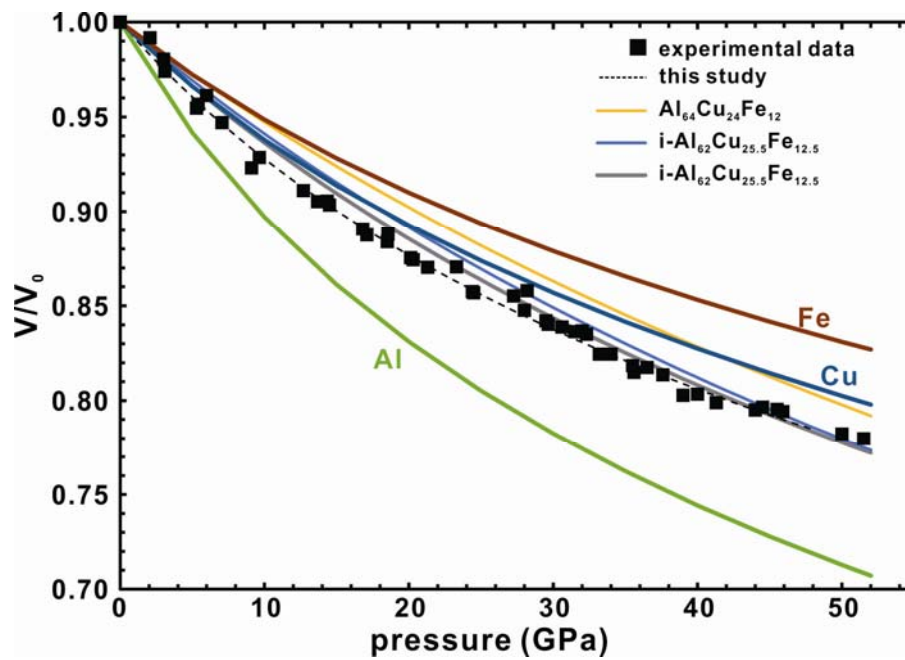


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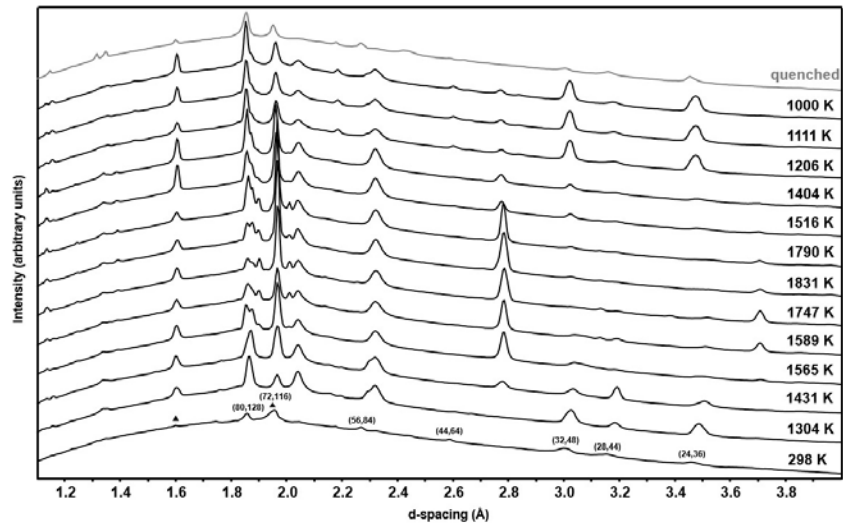
Figure 3



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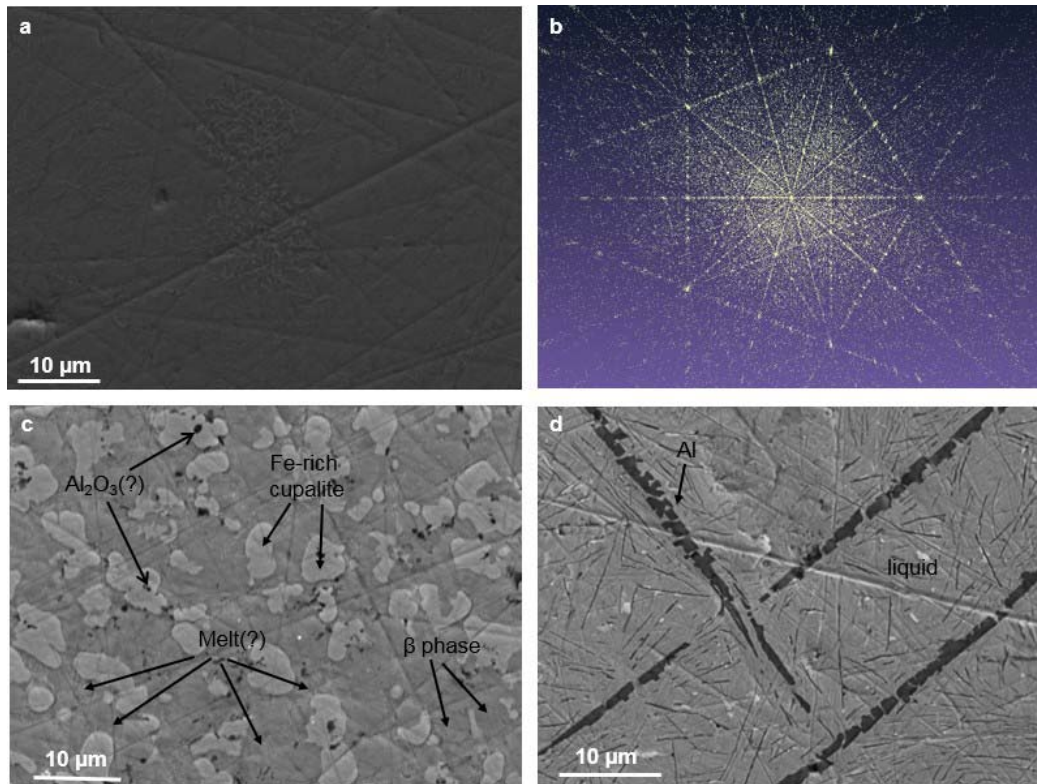
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Figure 4



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Figure 5



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Figure 6