

Scandium effect on undercooling and dendrite morphology of Al-4.5wt% Cu droplets

J. Valloton^{1,*}, A.-A. Bogno¹, H. Henein¹, D.M. Herlach² and D. Sediako³

¹ Advanced Materials and Processing Laboratory, University of Alberta, Edmonton, Canada, T6G 1H9

² Institut für Materialphysik im Weltraum, Deutsches Zentrum für Luft- und Raumfahrt, Cologne, Germany, 51170

³ UBC Okanagan, Kelowna, BC, Canada, V1V 1V7.

* Corresponding author: valloton@ualberta.ca

Abstract

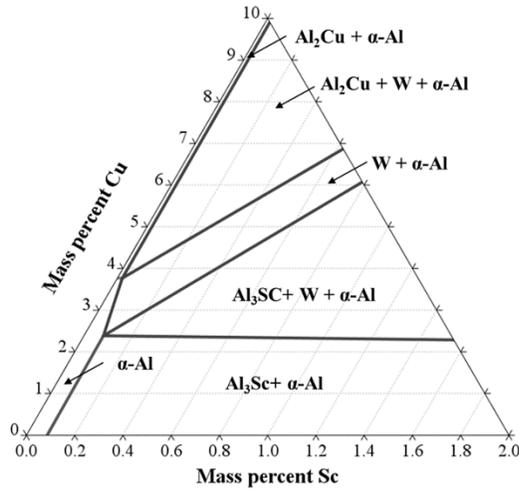
This paper reports on the undercooling and growth morphology of Al-4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc with a focus on the effect of Sc addition. It is found that the addition of Sc reduces the undercoolings of both primary phase and eutectic. In addition, the morphology of the Al-4.5wt%Cu-0.4wt%Sc dendrites is less favored in the $\langle 111 \rangle$ direction at similar undercoolings as with Al-4.5wt%Cu. The development of Solidification Continuous Cooling Transformation diagrams that relate the solidification paths to the inherent solidification microstructures is also introduced. The Solidification Continuous Cooling Transformation diagrams are obtained, based on the measurement of phase fractions of a solidified microstructure. The quantitative data is combined with well-established solidification models and phase diagrams to yield undercooling temperatures of individual phases. The thermal history and undercooling of different phases in the solidified alloy are estimated for a wide range of cooling rates (from 10^{-2} K/s to 10^4 K/s). It is found that a minimum cooling rate of about 1 K/s is required to avoid the nucleation of the detrimental intermetallic, W-phase in hypo-eutectic Al-Cu-Sc.

30 Introduction

31 Aluminum is the 3rd most abundant element on the planet, accounting for about eight percent of
32 the Earth's crust. Aluminum alloys occupy an important place in various industrial applications,
33 such as automotive and aerospace. This is due to their good mechanical properties coupled with
34 low density (i.e. excellent strength-to-weight ratio), corrosion resistance and castability [1].
35 Aluminum alloys are typically strengthened via precipitation of secondary intermetallic phases
36 from alloying elements in solid solution. Copper is one of the most widely used alloying element
37 due to its well-known age hardening effect characterized by precipitation of finely dispersed
38 Guinier–Preston (GP) zones, θ' and followed by the stable θ phase through heat treatment [2].

39 Lately, the development of commercial age-hardenable aluminum alloys with improved
40 performances has focused on systems forming Al_3X precipitates, such as Al-Sc, Al-Zr or Al-V [3].
41 Of these, Al-Sc alloys have garnered the most attention [4][5]. Age hardening leads to the
42 formation of a dense and homogeneous dispersion of approximately spherical Al_3Sc particles.
43 These nanosized precipitates effectively block the movement of dislocations and grain boundaries
44 and thus stabilize fine-grained structures [5][6][7]. Besides precipitation hardening, addition of Sc
45 to Al-alloys can also act as a grain refiner during casting using hypereutectic additions of Sc.
46 Indeed, Al_3Sc has an FCC structure and a lattice parameter close to that of α -Al [8]. For
47 hypereutectic compositions of Sc, Al_3Sc precipitates will thus act as nuclei for the formation of the
48 aluminum phase [9].

49 Ternary Al-Cu-Sc alloys have been scarcely studied. In addition to the traditional Al_2Cu and Al_3Sc
50 intermetallics, Kharakterova reported one ternary compound, $Al_{8-x}Cu_{4+x}Sc$ ($0 \leq x \leq 2.6$), that can be
51 in equilibrium with α -Al [10][11]. This phase, dubbed W-phase, is found in the Al-rich corner of the
52 Al–Cu–Sc phase diagram, as shown in Figure 1. Recently, Bo et al. carried out a thermodynamic
53 analysis of this system [12]. A good agreement was found between calculated phase equilibria
54 and the reported experimental data from Kharakterova. The W-phase was further evidenced in
55 the work of Bogno et al. during solidification of Al-4.5wt%Cu-0.4wt%Sc at low cooling rates [7].
56 Their work showed that the addition of 0.4wtSc to Al-4.5wt%Cu did not demonstrate any
57 significant benefit since most of the Sc precipitated as the W-phase. Only a slight hardness
58 increase was observed after heat treatment, which was attributed to the precipitation of the
59 remaining Sc in solid solution within the matrix to form Al_3Sc . In general, the formation of the W-
60 phase is detrimental as it consumes part of the Sc and Cu atoms in the Al matrix. As a
61 consequence, both the precipitation of Al_3Sc and the occurrence of Cu strengthening phases are
62 reduced and this minimizes the positive effect of Sc on the mechanical properties of the alloy [13].



63

64 **Figure 1:** Isothermal section of the Al-rich corner of the Al-Cu-Sc system at 500°C computed
 65 with Thermocalc [14] using the ALDEMO database.

66

67 Our previous study of rapidly solidified Al-4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc samples
 68 showed that with hypoeutectic additions of Sc in Al-4.5wt% Cu, no grain refining effect is observed
 69 [15]. Furthermore, rapid solidification supersaturates copper and scandium in the aluminum matrix
 70 and in the interdendritic regions, and thus prevents the formation of the W-phase. A dramatic
 71 improvement in mechanical properties is observed when Al-4.5wt%Cu-0.4wt%Sc samples are
 72 aged, with the microhardness increasing from about 75 HV as-atomized to 120 HV after heat
 73 treatment. This is attributed to the precipitation of nanosized Al₃Sc and Al₂Cu particles. Thus, by
 74 rapidly solidifying Al-4.5wt%Cu-0.4wt%Sc, the solutionizing and quenching step can be omitted
 75 from the regular heat treatment process.

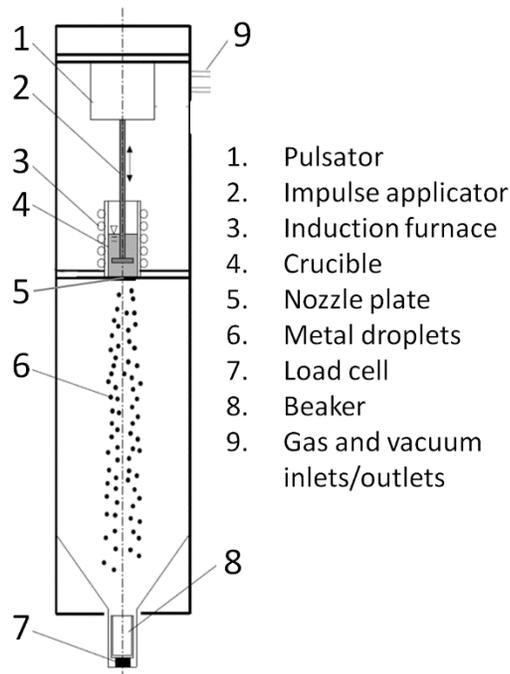
76 As scandium remains a very expensive alloying element, keeping its level low is economically
 77 warranted. This work thus reports on the solidification of Al-4.5wt%Cu containing 0, 0.2 and
 78 0.4wt%Sc under cooling rates varying from 10⁰ K/s to 10⁴ K/s by Differential Scanning Calorimetry
 79 (DSC), Electro-Magnetic Levitation (EML) and Impulse Atomization (IA). This paper focuses on
 80 the effect of Sc on the undercooling and the morphology of the solidified samples and introduces
 81 Solidification Continuous Cooling Transformation (SCCT) diagrams for Al-4.5wt%Cu and Al-
 82 4.5wt%Cu-0.4wt%Sc.

83

84

85 **Experimental methods**

86 Impulse atomization (IA) is a drop tube-type containerless solidification technique where
87 solidifying droplets experience high cooling rates and nucleation undercoolings [16]. It consists in
88 the transformation of a bulk liquid into a spray of liquid droplets that solidify rapidly during free fall
89 by losing heat to a surrounding gas of choice (usually N₂, Ar or He). The base material is melted
90 using an induction furnace and the atomization is achieved by the application of a mechanical
91 pressure (impulse) to the melt in order to push it through a nozzle plate with one or several orifices
92 of known size and geometry. A liquid ligament emanates from each orifice, which in turn breaks
93 up into droplets due to Rayleigh-type instabilities. The solidified powders are then collected in a
94 beaker at the bottom of the tower. IA generates a range of droplets sizes per run, giving a range
95 of cooling rates and undercoolings. The cooling rate is a function of both the droplet size and the
96 gas in the atomization tower and can reach up to ~10⁵ K/s. However, no direct measurement of
97 temperature has been feasible to date. The cooling rates of individual droplets are estimated using
98 a solidification model for atomization developed by Wiskel et al. [17][18] while primary and
99 secondary phase nucleation undercoolings are determined using a new, novel methodology
100 described in details in [19], with a summary given below.



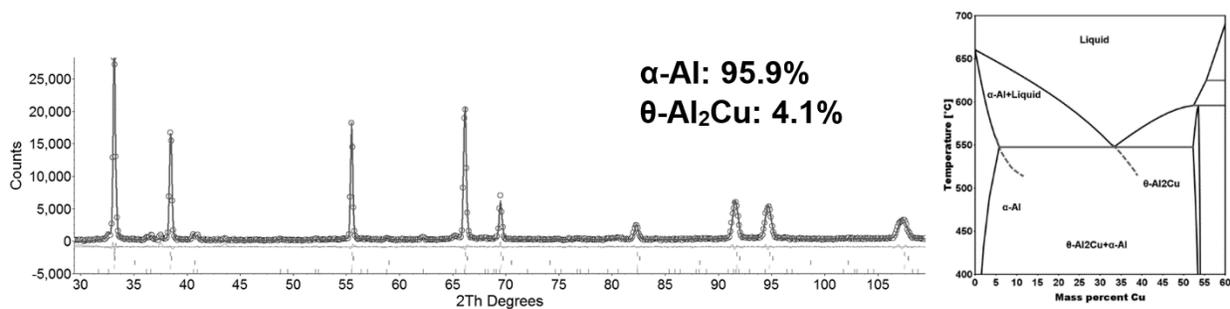
101

102

Figure 2: Schematic view of an impulse atomization apparatus.

103

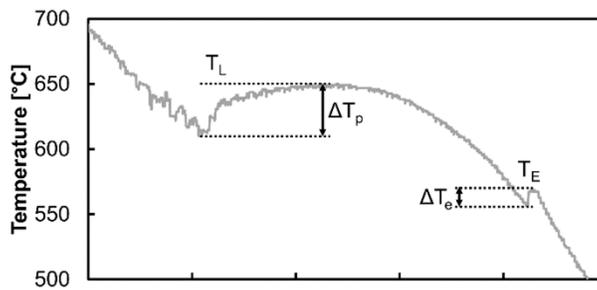
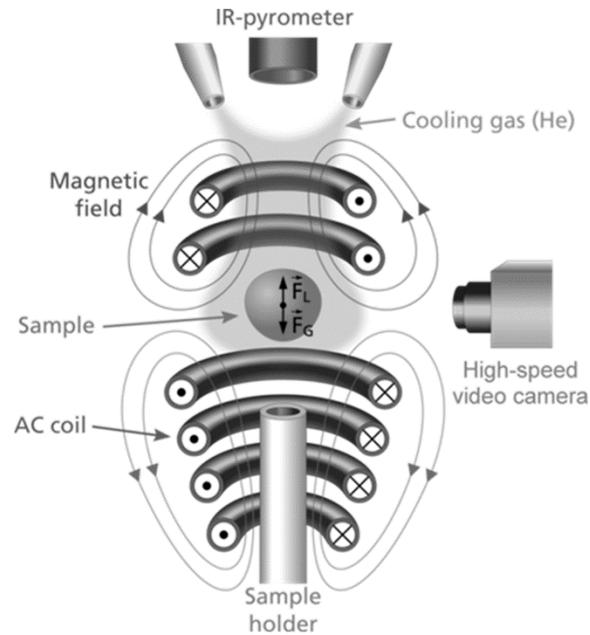
104 The methodology is based on experimental determination of phase fraction using Neutron
 105 Diffraction. The eutectic fraction is determined from the fraction of Al_2Cu obtained by Rietveld
 106 refinement of the diffraction spectra for an Al-Cu alloy (Figure 3 left) [20]. The corresponding
 107 eutectic nucleation undercooling is then evaluated from the metastable extension of the solidus
 108 and liquidus of the phase diagram of the alloy (Figure 3 right). The primary dendritic nucleation
 109 undercooling is subsequently determined using semi-empirical coarsening models of secondary
 110 dendrite arms. In the case of Al-4.5wt%Cu-0.4wt%Sc, the same methodology was used with a
 111 pseudo-binary phase diagram generated with ThermoCalc.



112
 113 **Figure 3:** Left: Neutron diffraction diagram of IA droplets of Al-4.5wt% Cu. The phase fractions
 114 are obtained by Rietveld refinement analysis of the diffraction pattern. Right: Al-richer corners of
 115 Al-Cu binary phase diagrams. The dashed lines represent the extensions of the solidus and
 116 liquidus lines obtained with ThermoCalc using the TTAL7 database.

117
 118 Electromagnetic levitation (EML) is a powerful containerless solidification technique for the
 119 processing of metallic and semiconductor samples with a large range of undercoolings. A
 120 schematic view of the apparatus is shown in Figure 4. An alternating current flowing through a
 121 water-cooled levitation coil produces an alternating electromagnetic field. A conducting sample
 122 placed within this field is levitated by the Lorentz force F_L which compensates for the gravitational
 123 force F_G . Simultaneously, the eddy currents induced in the sample heat and melt the sample by
 124 ohmic losses. To solidify the sample, cooling jets of inert gas are used. The temperature of the
 125 sample is monitored continuously with a two-color pyrometer (Impac IGA10-LO) with an accuracy
 126 of ± 5 K. As shown in Figure 4, this allows for a direct measurement of undercoolings, as well as
 127 cooling rates. Detailed information on the EML technique can be found in [21].

128



129

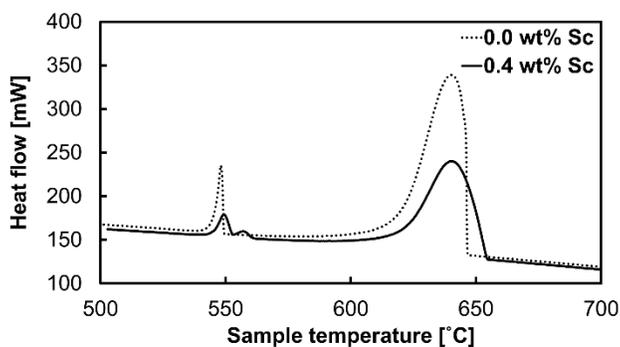
130 **Figure 4:** Schematic view of an EML apparatus and typical temperature-time profile obtained
 131 during EML solidification of an Al-4.5wt%Cu sample. Primary and eutectic solidification are clearly
 132 identified by the corresponding recalescence events of undercoolings $\Delta T_p = 34$ K and $\Delta T_e = 13$ K
 133 respectively.

134

135 DSC is a thermal analysis technique used to determine the amount of energy absorbed or
 136 released by a sample as it is heated or cooled in a controlled manner inside a crucible. As a non-
 137 containerless solidification technique, it yields low nucleation undercooling and its cooling rate
 138 (measurable) is limited to a narrow range (50 K/s maximum). In this work Al-Cu and Al-Cu-Sc
 139 alloys were solidified under low cooling rates and low undercooling conditions in a Setaram
 140 Labsys Evo 1600 DSC using alumina crucibles. A typical solidification curve using DSC is
 141 presented in Figure 5. The nucleation temperatures can be inferred from the onset of the
 142 exothermic peaks. Along with the primary and eutectic peaks, the Al-4.5wt%Cu-0.4wt%Sc sample

143 shows a third peak corresponding to the formation of the W-phase. Note that the difference in
144 heat flows stems from the different masses of the two samples analyzed.

145



146

147 **Figure 5:** DSC solidification curves of Al-4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc cooled at 5
148 K/min.

149

150 For metallographic analysis, DSC, EML and IA samples were first mounted in epoxy resin.
151 Grinding was carried out using silicon carbide papers up to grit 1000 (P2500), followed by
152 mechanical polishing with 3 and 1 μm diamond particles on soft cloths. Final polishing was
153 performed with a 0.05 μm colloidal silica. Microstructural characterization was carried out using
154 scanning electron microscopy (SEM) with a Zeiss Sigma FE-SEM running at 20 kV. Cell spacing
155 measurements were obtained using the line intercept method on selected SEM micrographs
156 according to ASTM E112-13.

157 Neutron diffraction measurements on IA samples were performed on the C2 neutron
158 diffractometer located at the Canadian Neutron Beam Centre in Chalk River, Canada.
159 Measurements were performed using a wavelength of 1.33 \AA from a Si531 monochromator at
160 $92.7^\circ 2\theta$.

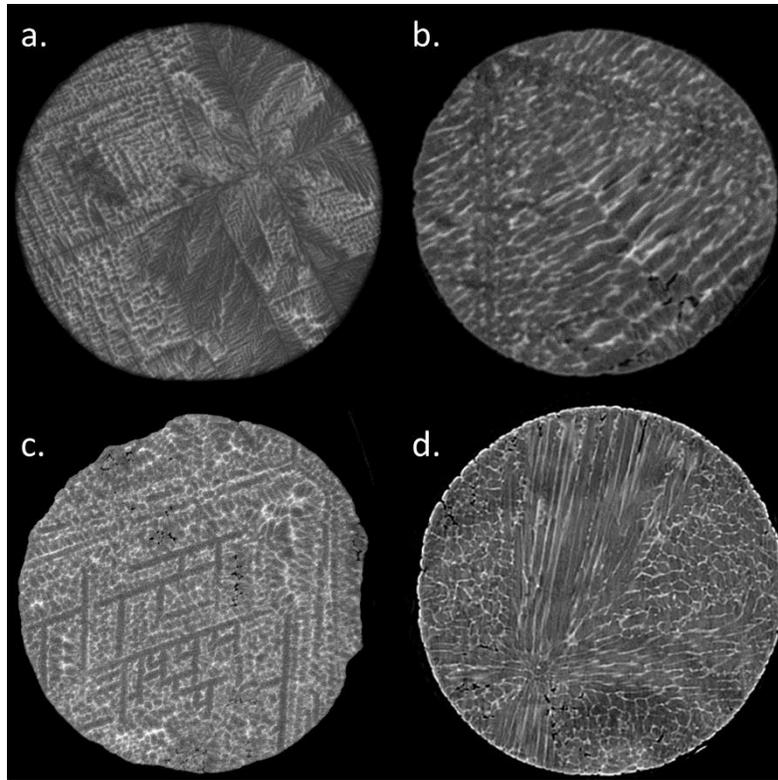
161 Synchrotron X-ray micro-tomography was carried out post-mortem at ESRF (European
162 Synchrotron Radiation Facility, Grenoble, France) on the ID19 beamline. Two pixel resolutions
163 were used: a high resolution of 0.18 $\mu\text{m}/\text{voxel}$ (field of view of 369 μm side cube) to analyze in
164 detail the fine microstructure of small droplets, and a medium resolution of 0.56 $\mu\text{m}/\text{voxel}$ (field of
165 view of 1146 μm side cube) to scan several small droplets at the same time to derive statistical
166 data.

167

168

169 **Results and discussion**

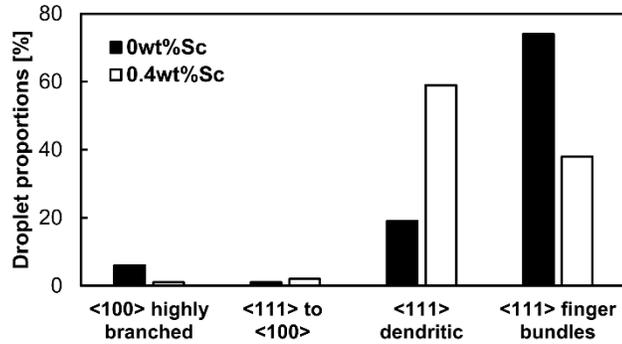
170 Cubic type crystals of metallic alloys such as Al-Cu, Al-Fe and Al-Ni generally grow along $\langle 100 \rangle$
171 directions due to the anisotropy of the solid-liquid interfacial energy. Under certain conditions (e.g.
172 high undercooling) dendrites growth deviates from $\langle 100 \rangle$ and unusual and complex morphologies
173 can develop. For example, in Al-0.6wt%Fe and Al-1.9wt%Fe impulse atomized droplets, a change
174 in dendrite growth direction from $\langle 100 \rangle$ to $\langle 111 \rangle$ was observed [22]. For IA Al-4.5wt%Cu droplets,
175 an earlier in-depth investigation by Bedel et al. [23] revealed four distinct dendritic morphologies
176 using X-Ray microtomography (Figure 6). The highly branched morphology (a) shows dendrite
177 growing along the usual $\langle 100 \rangle$ while microstructural features indicate that dendrite arms develop
178 mostly along $\langle 111 \rangle$ directions in the other three morphologies (b-d). The transition from $\langle 100 \rangle$ to
179 $\langle 111 \rangle$ is attributed to an increase in the solidification growth velocity. At the slowest solidification
180 growth velocity, $\langle 100 \rangle$ arms develop (a). At higher cooling rates and/or undercooling, primary
181 arms start growing along $\langle 111 \rangle$ but higher level arms forming after recalescence are slower and
182 thus grow along $\langle 100 \rangle$ (b). At even higher solidification rates, the droplet solidifies completely
183 with a $\langle 111 \rangle$ growth direction, as illustrated in (c). Finally, at the highest speed, a competition
184 between different $\langle 111 \rangle$ arms originating from the same nucleation point leads to the formation
185 of so-called finger bundles (d). Our collaborative work also showed that as the cooling rate
186 increases, $\langle 111 \rangle$ dendrites are favored and the number of droplets with a finger bundle
187 morphology increases.



188

189 **Figure 6:** Typical dendrite morphologies observed in Al-4.5wt%Cu droplets solidified in IA: (a)
 190 <100> highly branched dendrites; (b) <111> to <100> dendrite transition; (c) <111> dendritic
 191 morphology; (d) <111> finger bundle morphology.

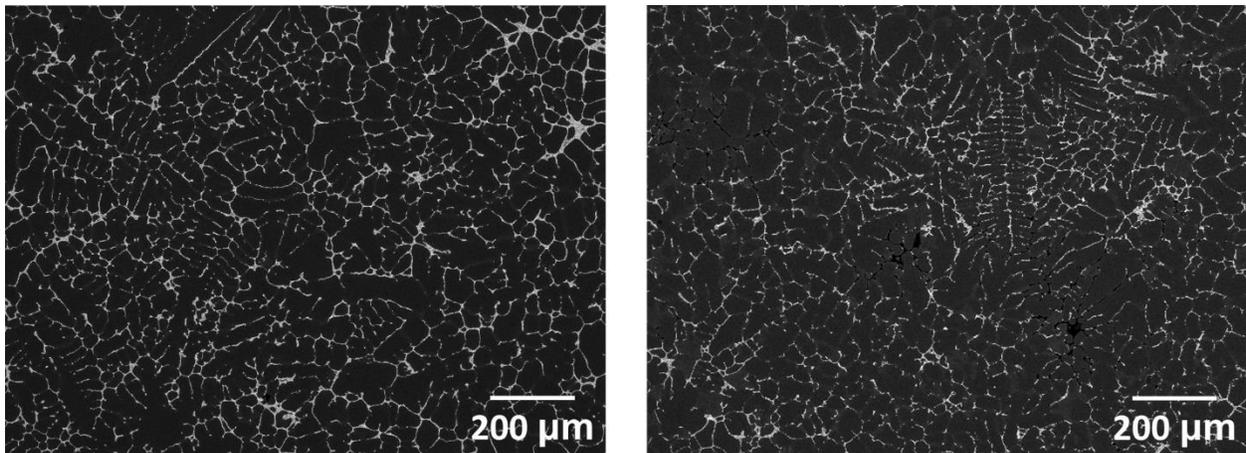
192 Figure 7 shows the statistical distribution of the four dendrite morphologies observed in IA droplets
 193 of composition Al-4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc solidified in argon with diameters in the
 194 range of 0 to 212 μm . A total of 69 droplets were analyzed for Al-4.5wt%Cu and 91 for Al-
 195 4.5wt%Cu-0.4wt%Sc. When Sc is added, it is observed that the number of droplets with finger
 196 bundles decreases in favor of lower speed morphologies. As the processing gas and the droplet
 197 size range are the same in both cases, the droplet cooling rates are expected to be similar,
 198 regardless of the presence of scandium. Thus, this shift in morphology is attributed to a change
 199 in the droplet undercoolings induced by the addition of Sc, as will be discussed later.



200

201 **Figure 7:** Statistical distribution of the four morphologies in Al-4.5wt%Cu and Al-4.5wt%Cu-
 202 0.4wt%Sc droplets Impulse Atomized in Ar with a diameter range $0 < d < 212 \mu\text{m}$.

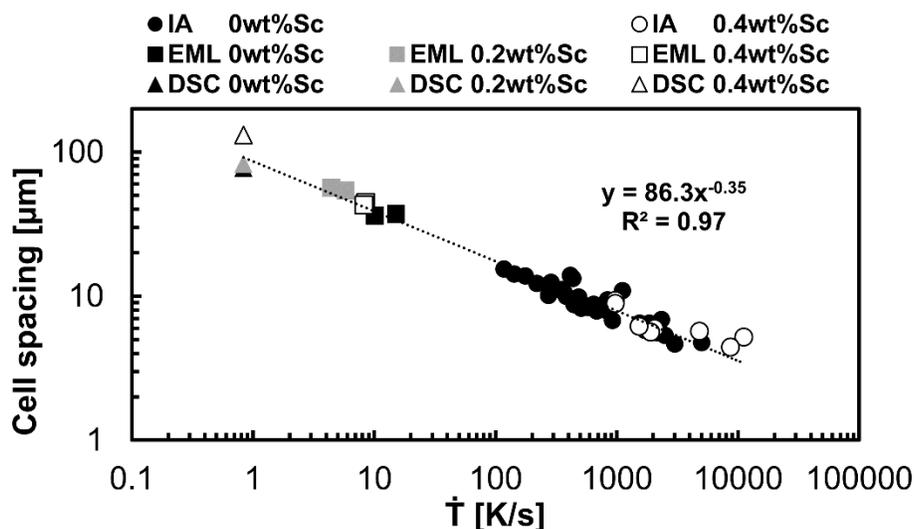
203 Figure 8 shows a representative solidification microstructure of an Al-4.5wt%Cu sample solidified
 204 in EML with a solidification cooling rate of $\sim 9 \text{ K/s}$ and primary and eutectic undercoolings of $\Delta T_p = 34 \text{ K}$
 205 and $\Delta T_e = 13 \text{ K}$, respectively, as well as an Al-4.5wt%Cu-0.4wt%Sc sample solidified at
 206 $\sim 12 \text{ K/s}$ with $\Delta T_p = 6 \text{ K}$ and $\Delta T_e = 4 \text{ K}$. EML samples typically show a cellular microstructure with
 207 no obvious growth direction. Some dendritic remnants can be observed in Figure 8 and in other
 208 EML samples. When present, these dendrites exhibit $\langle 100 \rangle$ growth direction in all cases. This
 209 suggests that the growth velocity in EML samples is slower than in IA. This is supported by the
 210 scale of the microstructure, which is one order of magnitude coarser in EML than in IA samples.
 211 Finally, it is observed that the addition of scandium does not alter the morphology of the solidified
 212 EML samples.



213

214 **Figure 8:** Representative solidification microstructure of Al-4.5wt%Cu solidified in EML at $\sim 9 \text{ K/s}$
 215 with $\Delta T_p = 34 \text{ K}$ and $\Delta T_e = 13 \text{ K}$ (left) and Al-4.5wt%Cu-0.4wt%Sc solidified in EML at $\sim 12 \text{ K/s}$
 216 with $\Delta T_p = 6 \text{ K}$ and $\Delta T_e = 4 \text{ K}$ (right). In both cases, the structure is mostly cellular with some
 217 dendrite remnants showing $\langle 100 \rangle$ growth directions.

218 Figure 9 shows the cell spacing measured with the line intercept methods as a function of cooling rate for DSC, EML and IA samples [15]. Regardless of the solidification technique, the effect of
 219 rate for DSC, EML and IA samples [15]. Regardless of the solidification technique, the effect of
 220 Sc addition on the microstructures scale is found to be negligible. However, microstructure
 221 refinement is shown to be very dependent on the cooling rate. This relationship follows a power
 222 law of the type $\lambda_2 = A\dot{T}^{-n}$, where λ_2 is the cell spacing, \dot{T} the cooling rate and A and n are constants,
 223 as described by Eskin et al. [24]. The values of A and n found in this study are in the range of
 224 values published by Mullis and co-workers in the estimation of cooling rates during close-coupled
 225 gas atomization of Al-4wt%Cu using secondary dendrite arm spacing measurements [25].
 226 Furthermore, the exponent value being very close to the theoretical value of 1/3 [26], the decrease
 227 in cell spacing observed with increasing cooling rate indicates that the final scale of the
 228 microstructure is governed mainly by coarsening.



229
 230 **Figure 9:** Variation of cell spacing with cooling rate for Al-4.5 wt% Cu with different Sc levels.
 231

232 As EML experiments allow the direct measurement of the temperature profile during solidification,
 233 it is easy to assess the effect of scandium addition on the nucleation behavior of the alloy studied.
 234 Table 1 compiles the average of the measured undercoolings obtained during EML solidification
 235 of Al-4.5wt%Cu samples with the addition of 0, 0.2 and 0.4wt%Sc (9, 7 and 12 solidified samples
 236 respectively). Nucleation being a stochastic event, a range of undercoolings is obtained for each
 237 composition, which is reflected in the standard deviation for each composition in Table 1. It is
 238 clear that the addition of Sc promotes the nucleation of the α -Al and θ -Al₂Cu phases as both the
 239 primary and eutectic undercoolings decrease when Sc is added to the alloy. The primary
 240 undercooling decreases significantly only with a 0.4wt%Sc addition, while both 0.2 and 0.4wt%Sc

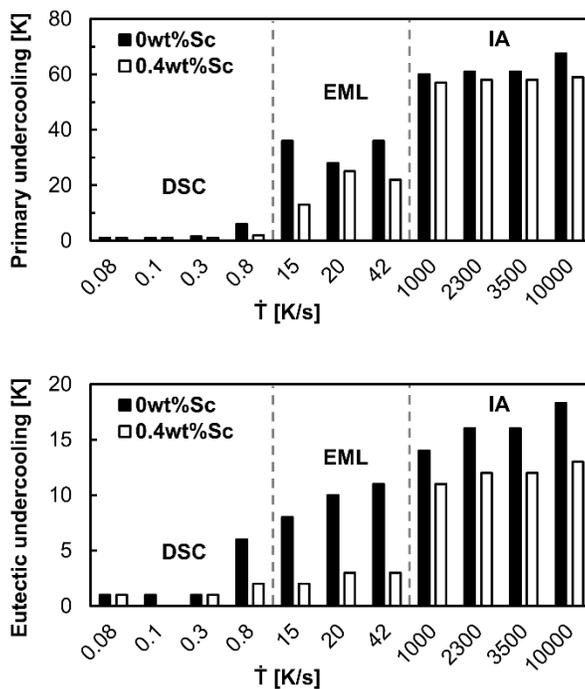
241 exhibit the same change in eutectic undercooling. This suggests that the presence of scandium
 242 in the melt is sufficient to alter the interfacial energy between the Sc containing liquid and Al₂Cu
 243 and to promote nucleation of the intermetallic phase. The results of cooling rate of the liquid
 244 samples, as well as the primary and eutectic undercoolings for each method used is presented in
 245 Figure 10. The same trend is observed, regardless of the cooling rate or solidification technique.

246

247 **Table 1:** Average primary and eutectic nucleation undercoolings during EML solidification.

	ΔT_p [K]	ΔT_e [K]
0 wt% Sc	35.7 ± 6.5	10.1 ± 1.9
0.2 wt% Sc	30.9 ± 6.4	3.6 ± 0.9
0.4 wt% Sc	13.8 ± 5.7	3.7 ± 1.8

248



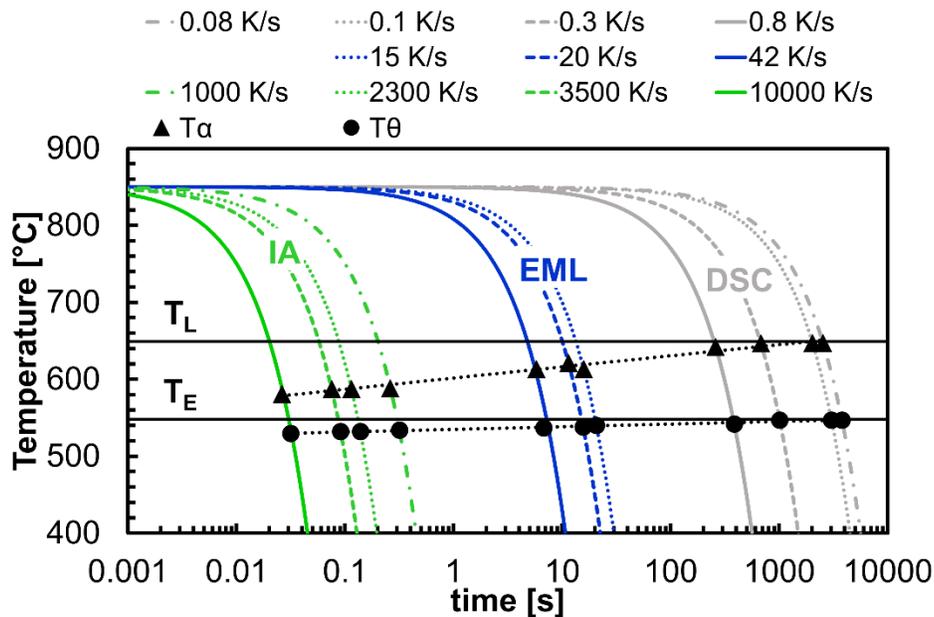
249

250 **Figure 10:** Variation of nucleation undercooling with solidification cooling rate for Al-4.5wt%Cu
 251 and Al-4.5wt%-0.4wt%Sc

252 As reported in a previous contribution, X-ray and Neutron Diffraction of EML and IA samples did
 253 not show the presence of the W-phase. Only in the DSC solidified samples was the W-phase
 254 observed [7]. Knowing this, and with the results presented above, the quantification of the
 255 solidification path of Al-4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc alloys is possible. In order to

256 represent the resultant microstructure and relate it to macro-solidification conditions, solidification
 257 continuous cooling transformation (SCCT) curves were developed. To construct these maps, the
 258 liquid cooling rates of the samples were used (imposed for DSC, measured on the temperature-
 259 time profile for EML, and estimated with the atomization model for IA) and the corresponding
 260 undercoolings from Figure 10 were plotted on a CCT diagram (Figure 11 and 12). Also plotted on
 261 these diagrams are the equilibrium liquidus, T_L , and eutectic, T_E , of the respective alloys. From
 262 these results, it is observed that both primary phase and eutectic nucleation undercoolings
 263 increase as the cooling rate is increased. Also, to avoid the formation of the detrimental ternary
 264 W-phase, a cooling rate of the order of 1 K/s is necessary.

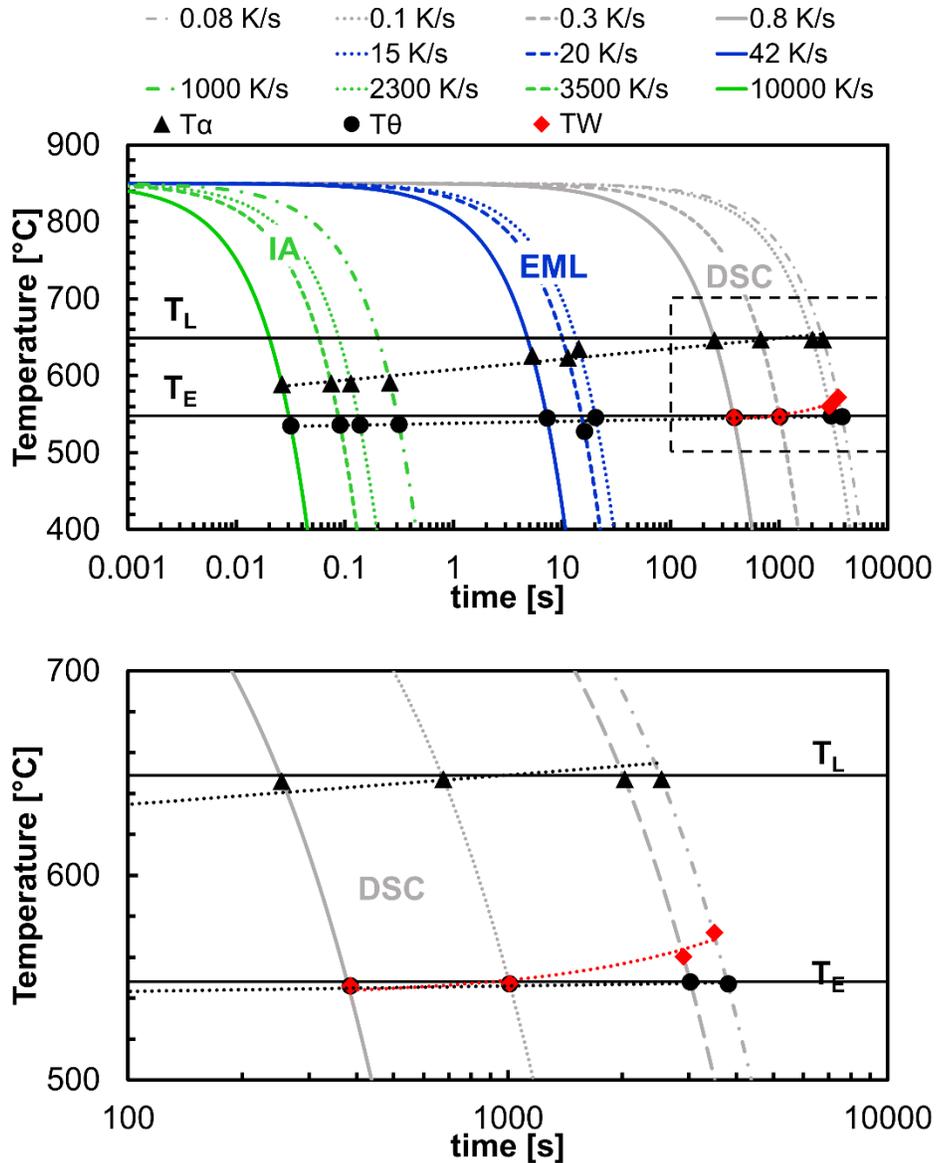
265 The use of the SCCT diagrams is not restricted to solidifying liquid droplets but should apply to
 266 any liquid of the Al-4.5wt%Cu-(0.4wt%Sc) composition solidifying in any given solidification
 267 process. Limitations to the use of this diagram will occur when there is significant segregation of
 268 Cu during solidification. However, in these instances, similar SCCT diagrams may be derived
 269 using droplet cooling rates for alloys with different Cu compositions, to trace the path of
 270 solidification of a given alloy in a particular process. This is the subject of further research.



271

272

Figure 11: Solidification Continuous Cooling Transformation curves of Al-4.5wt%Cu



273

274 **Figure 12:** Top: Solidification Continuous Cooling Transformation curves of Al-4.5wt%Cu-

275 0.4wt%Sc. Bottom: Magnified view of the nucleation temperatures of the ternary W-phase.

276

277 Conclusions

278 Al-4.5wt% Cu with different level of Sc addition (0, 0.2 and 0.4wt%) were generated under low,

279 medium and high cooling rate conditions respectively by DSC, EML and IA. No refining effect of

280 Sc is found. Cell spacing variation with cooling rate for the investigated alloys is found to follow

281 an empirical coarsening law of secondary dendrite arms spacing commonly found in literature. Sc

282 addition is shown to reduce both the primary and eutectic undercoolings in all three types of

283 solidification experiments carried out. This in turns promotes the formation of lower speed
284 morphologies as evidenced by the decrease of droplets exhibiting finger bundles in IA samples.

285 Using solidification continuous cooling transformation maps, the solidification path of Al-
286 4.5wt%Cu and Al-4.5wt%Cu-0.4wt%Sc has been charted over a wide range of cooling rates. An
287 increase in cooling rate leads to an increase of the primary and eutectic undercoolings. During
288 DSC experiments at low cooling rates, scandium induces the precipitation of the W-phase, which
289 is detrimental to the mechanical properties of the alloy. At higher cooling rates in EML and IA, the
290 precipitation of the W-phase is suppressed. The SCCT diagram shows that a minimum cooling
291 rate of about 1 K/s is required to avoid the nucleation of the detrimental intermetallic, W-phase.

292

293 **Acknowledgements**

294 Financial support from the Natural Sciences and Engineering Research Council of Canada
295 (NSERC) and the European Space Agency (ESA) within the frame of the NEQUISOL project is
296 gratefully acknowledged. The authors are grateful to the Canadian Nuclear Laboratories (CNL)
297 and the European Synchrotron Radiation Facility (ESRF) for beam time and expert support during
298 the measurement campaigns. The assistance of Daniel Auras with morphology analysis is
299 appreciated.

300

301 **References**

- 302 [1] J.R. Davis (ed.): *ASM Specialty Handbook: Aluminum and Aluminum Alloys*, ASM
303 International 1993.
- 304 [2] P. Haasen: *Physical Metallurgy*, 3rd ed., Cambridge University Press, Cambridge, UK,
305 1996.
- 306 [3] Z.C. Sims, O.R. Rios, D. Weiss, P.E.A Turchi, A. Perron, J.R.I. Lee, T.T. Li, J.A.
307 Hammons, M. Bagge-Hansen, T.M. Willey, K. An, Y. Chen, A.H. King and S.K. McCall:
308 *Mater. Horiz.*, 2017, vol. 4 (6), pp. 1070-1078.
- 309 [4] N. Blake and M.A. Hopkins: *J. Mater. Sci.*, 1985, vol. 20 (8), pp. 2861-2867.
- 310 [5] D.N. Seidman, E.A. Marquis and D.C. Dunand: *Acta Mater.*, 2002, vol. 50 (16), pp. 4021-
311 4035.
- 312 [6] L.S. Toporova, D.G. Eskin, M.L. Kharakterova and T.V. Dobatkina: *Advanced*
313 *aluminum alloys containing scandium: structure and properties*, Gordon &
314 Breach, Amsterdam, 1998.

- 315 [7] A.-A. Bogno, J. Valloton, H. Henein, D.G. Ivey, A.J. Locock, and M. Gallerneault: *Can.*
316 *Metall. Quart.*, 2018, vol. 57 (2), pp. 148-159.
- 317 [8] J. Røyset and N. Ryum: *Int. Mater. Rev.*, 2005, vol. 50 (1), pp. 19-44.
- 318 [9] J. Røyset: *Metall. Sci. Technol.*, 2007, vol. 25 (2), pp. 11-21.
- 319 [10] M.L. Kharakterova and T.V. Dobatkina: *Russ. Metal.*, 1988, vol. 6, pp. 175-178.
- 320 [11] M.L. Kharakterova: *Russ. Metal.*, 1991, vol. 4, pp. 191-194.
- 321 [12] H. Bo, L.B. Liu and Z.P. Jin: *J. Alloy. Compd.*, 2010, vol. 490 (1-2), pp. 318-325.
- 322 [13] S. Riva, K.V. Yusenko, N.P. Lavery, D.J. Jarvis and S.G.R. Brown: *Int. Mater. Rev.*,
323 2016, vol. 61 (3), pp. 203-228.
- 324 [14] J.O. Andersson, T. Helander, L. Höglund L, S. Pingfang and B. Sundman: *Calphad.*
325 2002, vol. 26 (2), pp. 273-312.
- 326 [15] A.-A. Bogno, J. Valloton, H. Henein, M. Gallerneault and D.M. Herlach: in *Proc. 3rd Pan*
327 *American Materials Congress*, Berlin, Springer, 2017, pp. 355-363.
- 328 [16] H. Henein: *Mater. Sci. Eng.*, 2002, vol. A326, pp. 92-100.
- 329 [17] J. Wiskel, H. Henein and E. Maire: *Can. Metall. Quart.*, 2002, vol. 41 (1), pp. 97-110.
- 330 [18] J. Wiskel, K. Navel, H. Henein and E. Maire: *Can. Metall. Quart.*, 2002, vol. 41 (2), pp.
331 193-204.
- 332 [19] A.-A. Bogno, P.D. Khatibi, H. Henein and Ch.-A. Gandin: *Metall. Mater. Trans. A*, 2016,
333 vol. 47 (9), pp. 4606-4615.
- 334 [20] R. A. Young (ed.): *The Rietveld Method*. International Union of Crystallography/Oxford
335 University Press, Oxford, UK, 1993.
- 336 [21] D. M. Herlach and D. M. Matson (eds.): *Solidification of Containerless Undercooled*
337 *Melts*, 1st ed, Wiley-VCH, Weinheim, 2012, pp. 9-16.
- 338 [22] J. Chen, U. Dahlborg, C.M. Bao, M. Calvo-Dahlborg, and H. Henein: *Metall. Mater.*
339 *Trans. B*, 2011, vol. 42 (3), pp. 557-567.
- 340 [23] M. Bedel, G. Reinhart, A.-A. Bogno, Ch.-A. Gandin, S. Jacomet, E. Boller, H. Nguyen-
341 Thi and H. Henein: *Acta Mater.*, 2015, vol. 89, pp. 234-246.
- 342 [24] D. Eskin, Q. Du, D. Ruvalcaba and L. Katgerman: *Mater. Sci. Eng.*, 2005, Materials
343 Science and Engineering A, 405 (1-2), pp. 1-10.
- 344 [25] A.M. Mullis, L. Farrell, R.F. Cochrane and N.J. Adkins: *Metall. Mater. Trans. B*, 2013,
345 vol. 44 (4), pp. 992-999.
- 346 [26] J. A. Dantzig and M. Rappaz: *Solidification*, EPFL Press, Lausanne, 2009.