Al₈Mn₅ in high pressure die cast AZ91: twinning, 2 morphology and size distributions

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14

15 Abstract

Manganese-bearing intermetallic compounds (IMCs) are important for limiting micro-16 galvanic corrosion of magnesium-aluminium alloys and can initiate cracks under tensile load. 17 Here we use electron backscatter diffraction (EBSD), deep etching, and focussed ion beam 18 (FIB) tomography to investigate the types of Al-Mn phases present, their faceted growth 19 20 crystallography, and their three-dimensional distribution at different locations in high pressure die cast (HPDC) AZ91D. The Al-Mn particle size distributions were well-described 21 by lognormal distributions but with an additional population of externally solidified crystals 22 (ESCs) formed in the shot chamber analogous to α -Mg ESCs. The large Al₈Mn₅ particles were 23 cyclic twinned. Differences in the particle size distributions and number density in the centre 24 compared with the HPDC skin are identified, and the spatial relationship between Mg₁₇Al₁₂ 25 26 and Al-Mn particles is explored.

27 Keywords AZ91, high pressure die casting, intermetallics

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29 Introduction

Automotive magnesium components are often Mg-Al-based alloys produced by high pressure die casting (HPDC). When conducted with an optimised die, process parameters and vacuum system ^[1,2], HPDC can mass produce large, thin-walled, complex shapes

containing microstructures with fine α -Mg grains (5-20 µm) ^[3,4], and a fine-scaled percolating eutectic Mg₁₇Al₁₂ network ^[5,6]. While a large body of research has investigated microstructure formation in Mg HPDC, including the formation of α -Mg grains ^[3,4,7], the surface 'skin' ^[4,8], the eutectic Mg₁₇Al₁₂ ^[5,9,10], and casting defects ^[11–17], less work has explored the formation of Al-Mn-(Fe) intermetallic particles ^[18–21]. These particles play an important role in determining micro-galvanic corrosion in HPDC Mg parts ^[22,23] and can initiate cracks under tensile loading ^[24,25].

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Most Mg-Al-based HPDC alloys (e.g. AM50A, AM60B, AZ91D^[26]) contain sufficient Mn and 41 42 Al that Al_8Mn_5 begins to form before α -Mg during solidification. For example, Figure 1 shows the sequence of phase formation assuming Scheil solidification of AZ91D with the 43 44 composition in Table 1, calculated with the Thermo-Calc TCMG magnesium database version 4 ^[27]. It can be seen that Al_8Mn_5 is the first solid phase to form, and becomes stable ~44K 45 46 above the α -Mg liquidus temperature for this composition. It has been confirmed by in-situ 47 X-ray imaging that Al₈Mn₅ forms at higher temperature (i.e. earlier on cooling) than α -Mg in a similar alloy ^[28,29]. A consequence of this in HPDC is that Al₈Mn₅ can form and settle in the 48 holding pot ^[29,30], for example during temperature drops when charging the furnace with 49 new ingots, leading to die casting sludge ^[30]. Furthermore, in cold chamber HPDC, heat loss 50 51 in the shot chamber can cause Al₈Mn₅ formation prior to injection as Al₈Mn₅ externally solidified crystals (ESCs) ^[20] in addition to the α -Mg ESCs that are widespread in HPDC Mg 52 components ^[3,14,31]. This occurs because a feature of Mg HPDC is partial solidification in the 53 shot chamber that leads to large α -Mg externally solidified crystals (ESCs) being injected into 54 the cavity ^[3,32]. The volume fraction of α -Mg ESCs has been shown to depend on the melt 55 superheat, the fill fraction and the temperature of the sleeve walls and plunger tip, and is 56 typically 10-30 vol.% ^[3,14,31,33]; similar factors might be expected to determine the formation 57 of Al₈Mn₅ ESCs. 58

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Table 1. Composition of the AZ91D alloy used (weight percent).

-	Mg	Al	Zn	Mn	Fe	Ni	Cu	Si	Be
_	bal.	8.95	0.72	0.19	<0.001	<0.001	0.001	0.039	0.0007

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Figure 1 shows that AI_8Mn_5 continues forming along with α -Mg below the α -Mg liquidus temperature until ~ 510°C when other Al-Mn IMCs start forming ($AI_{11}Mn_4$ and then AI_4Mn). Therefore, in HPDC, Al-Mn IMCs are expected to form in all stages of the process: in the shot chamber, during filling and during the intensification stage. According to calculations linked with Figure 1, at the end of Scheil solidification, the total mass fraction of Al-Mn IMCs (AI_8Mn_5 , $AI_{11}Mn_4$ and AI_4Mn) is 0.25% of which 95% is AI_8Mn_5 for the composition in Table 1.



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Figure 1: Phase formation during Scheil solidification up to 99% solid for Mg-8.95Al-0.72Zn-0.19Mn
(wt%). Calculated with Thermo-Calc TCMG magnesium database version 4 [27].

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Past work on AI-Mn particles in HPDC AZ91D has generally used TEM ^[18,19,21]. That work has 73 deduced that most Al-Mn particles in HPDC AZ91D are 100 nm to 1µm in size. The main 74 phase present has been found to be Al₈Mn₅ and another phase with higher Al content 75 (possibly $AI_{11}Mn_4$) has also been reported ^[18]. While these TEM studies enable high resolution 76 imaging, they did not explore the statistical variation in Al-Mn particle size and shape versus 77 position in the cross-section. This is an important question in HPDC parts since they usually 78 have highly non-uniform microstructures. They typically have a surface layer (a skin) of 79 distinctly different microstructure that is usually free of porosity and harder than more 80 central regions, one or more bands of porosity, various forms of macrosegregation, and ESCs 81 that tend to be concentrated towards the centre of cross-sections (e.g. ^[15,16,33,34]). 82

In this paper, we investigate the types of Al-Mn phases present, their faceted growth crystallography, and their three-dimensional distribution at different locations in high pressure die cast AZ91D. The specific aims are: (i) to compare the Al₈Mn₅ growth crystallography and twinning formed in HPDC with past work at sand casting cooling rates ^[35]; (ii) to quantify the 3D size, morphology and spatial distribution of Al-Mn particles in different locations in HPDC AZ91D: the skin, the defect band, and the centre; and (iii) to explore any correlations between Al-Mn particles and eutectic Mg₁₇Al₁₂ in 3D.

92 Methods

 93 ~6 kg of AZ91D Mg alloy with composition in Table 1 was melted in a mild steel crucible and held at 675°C (~ 75°C superheat) under a cover gas of ~3 vol% SF₆ in N₂. HPDC was conducted using a Frech DAK 450-54 cold chamber HPDC machine and the multi-cavity die that produces the casting in Figure 2. The die was preheated to 150°C, a portion of the melt was ladled into the shot chamber to a fill fraction of ~0.5, and the following set parameters were used: slow shot phase of 0.3 m.s⁻¹, fast shot phase of 4 m.s⁻¹, and intensification pressure of 36 MPa. The casting analysed in this work was made after six pre-shots.



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Figure 2 (a-b) Photographs of the HPDC part. The sectioning plane is indicated by superimposed lines.(c) as-polished optical micrograph.(d) the same section after etching.

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Samples for microstructural analysis were cut from the centre of the gauge length into slices of 10mm x 10mm x 0.5mm. Metallographic polishing was carried out down to 0.05µm colloidal silica by standard preparation methods. Some samples were etched in a solution of 200ml ethylene glycol, 68ml distilled water, 4ml nitric acid and 80 ml acetic acid. Both etched and polished samples were analysed in a Zeiss AURIGA field emission gun SEM (FEG-SEM)

with an Oxford Instruments INCA x-sight energy dispersive X-ray spectroscopy (EDX) 109 detector and a BRUKER e-Flash^{HR} electron backscatter diffraction (EBSD) detector. For EBSD 110 characterisation, the final step of preparation was Ar-ion milling for 40 min in a Gatan PECSII 111 112 instrument. The 4kV-accelerated beam hit the sample rotating at 2rpm, at a grazing incidence angle of 4°. Electron beam accelerating voltage of 20kV, working distance of 113 15mm, aperture size of 120mm, and beam current 80µA were used for EBSD measurements. 114 Bruker ESPRIT 2.1 software was used to index the obtained EBSD patterns. EBSD datasets 115 were analysed using MATLAB[™] 9.2 (Mathworks, USA) with the MTEX 5.1 toolbox ^[36]. 116 Accelerating voltage of 10kV, working distance of 5mm, aperture size of 60mm, and beam 117 118 current 80µA were used for EDS analysis. EDS spectrum was calibrated with a Si standard 119 sample prior to each electron microscopy session.

To investigate the 3-dimensional (3D) morphology of the Al-Mn intermetallics directly, α -Mg 120 was selectively etched using a solution of 4% nitric acid in ethanol. To quantify the 3D size 121 distribution of Al-Mn intermetallics, focussed ion beam (FIB) tomography was conducted in a 122 Zeiss AURIGA FG-SEM at 30 kV with 52° tilt angle. The slice distance was 90 nm and the 123 milling current was 200pA. Serial-sectioning secondary electron images were used. For FIB 124 tomography, 2D slices were aligned, cropped, and processed by an anisotropic diffusion filter 125 126 in ImageJ (US NIH, USA). 3D reconstruction and crystallographic analysis was performed using Avizo 9.2 (Visualization Science Group, France) and MATLAB 9.2™. The voxel size for 127 FIB tomography was bounded by the slice spacing of 90nm. Al₈Mn₅ particles with equivalent 128 diameter \geq 180nm were quantified. 129

To study porosity bands in 3D, X-ray micro-tomography was carried out on a North Star 130 Imaging (NSI) Micro-CT. The system is equipped with a 225 kV X-ray source with a minimum 131 focal spot size of 2 µm and a Perkin Elmer flat panel detector (2048×2048 pixels at 16bit 132 depth). During a CT scan, the sample was illuminated by cone beam X-rays which were 133 transmitted through the 360° rotating specimen and then illuminated on the flat panel 134 135 detector. The X-ray beam was filtered using a 0.25 mm Cu filter to reduce beam-hardening effects, and an acceleration voltage of 80kV and target current of 35µA was selected to 136 optimise image quality. 1440 two-dimensional projections were captured over 360° with an 137 exposure time of 1000ms. 3D reconstruction was performed in Avizo 9.2 and resulted in a 3D 138 spatial resolution with voxel size of 2.2 µm x 2.2 µm x 2.2 µm. 139

140 **3 Results and Discussion**

141 **3.1 General microstructural features**

At the centre of the gauge length, the AZ91D samples contained the typical microstructural 142 features and defects of HPDC reported in past work (e.g. [3,7,12,14,16,33,34,37]). For example, 143 annular rings of porosity can be seen in the as-polished condition in Figure 2(c), a dark band 144 of macrosegregation can be seen in the same location as the main porosity band in Figure 145 2(d) after light etching, and a high fraction (~30 vol%) of α -Mg ESCs can be seen throughout 146 147 much of the cross-section in Figure 2(d). However, the detail of these features differed significantly from casting to casting and between bars in the same casting as shown in the X-148 ray tomographs in Figure 3. The left-hand images are reconstructed volumes near the centre 149 of the gauge length showing the 3D distribution of porosity. The right-hand images are 150 viewed along the tensile rod axis to highlight the radial distribution of porosity. There are 151 major differences in the porosity in the two samples. The sample in Figure 3(b) has a 152 localised annular ring of porosity and a high fraction of porosity within this ring. The sample 153 154 in Figure 3(a) has more diffuse porosity and a less-well defined porosity ring but has the same trend of a higher fraction of porosity within the annular porosity band. Despite the 155 differences, in both samples, the main annular ring of porosity is at a similar radial position. 156 The projection images along the rod axes also reveal the surface 'skin' as an outer ring of 157 essentially zero porosity. This is particularly clear in Figure 3(a) where the abrupt change in 158 porosity demarcates the edge of the skin. 159



Figure 3 (a-b) X-ray tomograms of porosity near the centre of the gauge-length of typical castings.
Porosity is rendered as solid, material (Mg, Mg₁₇Al₁₂ and Al-Mn IMCs) is plotted as semi-transparent.
Left-hand side: perspective view. Right-hand side: projection view along the tensile rod axis.

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165 The typical α -Mg microstructure is shown in more detail in Figure 4(a)-(b). The micrograph 166 in Figure 4(a) shows the complex mixture of dendritic α -Mg ESCs, ESC fragments and in-167 cavity solidified grains. Figure 4(b) is an EBSD orientation map (IPF-y) of the α -Mg phase 168 from a similar region where the grains have been coloured by their mean-orientation. The 169 grains form a complex multimodal microstructure with, in this case, two large ESCs 170 surrounded by smaller α -Mg grains that are probably a mixture of α -Mg ESC fragments and 171 in-cavity solidified grains.

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173 The typical features of intermetallic compounds in the HPDC bars are overviewed in Figure 174 4(c) and (d). It can be seen that the eutectic Mg₁₇Al₁₂ phase appears as isolated regions in 2D 175 sections (Figure 4(c)) but actually forms a percolating Mg₁₇Al₁₂ network in 3D as revealed by 176 imaging after selective dissolution of the α -Mg in Figure 4(d). Figure 4(c) and (d) also 177 contains bright particles that are Al-Mn compounds. In the 2D section these appear both 178 within the α -Mg grains and near the Mg₁₇Al₁₂ phase (Figure 4(c)). After deep etching, it can 179 be seen that many Al-Mn particles are attached to the Mg₁₇Al₁₂ network (Figure 4 (d)).



Figure 4: Typical microstructural features in the HPDC AZ91 samples. (a) mixture of α -Mg ESCs and in-cavity solidified grains. (b) EBSD orientation map (IPF-Y) of the α -Mg phase. (c) 2D section of Mg₁₇Al₁₂ and Al₈Mn₅ phases. (d) 3D microstructure of Mg₁₇Al₁₂ network and attached Al₈Mn₅ particles, revealed after selective etching of α -Mg.

186 The remainder of this paper focuses on the Al-Mn intermetallic compounds and their 187 relationship to the microstructural features summarised in this section.

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189 **3.2 Twinned Al₈Mn₅ in HPDC AZ91D**

Al-Mn intermetallics were identified by combining EDS with EBSD. A typical EDS point 190 analysis from an Al-Mn particle is shown in Figure 5(a). The particle contains 59at%Al -191 40at%Mn and there are also small Mg, Si and Fe peaks, each present at less than 1 at%. 192 Since the solubility of Mg in Al-Mn intermetallics is negligible ^[38], the small Mg peak is likely 193 to be α -Mg in the interaction volume. The small Si peak is probably Si dissolved in the 194 particle, consistent with past work that has detected a small Si content in Al-Mn IMCs ^[18,39]. 195 The low Fe content in the particle is due to the high-purity AZ91D used in this study (with 196 <10ppm Fe, Table 1). 197

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An EBSD pattern from the Al-Mn particle is shown in Figure 5(b). This could be readily 199 distinguished as the rhombohedral Al_8Mn_5 phase ^[40,41] using the Hough transform-based 200 201 method in Bruker ESPRIT 2.1, and is indexed in Figure 5(c) in the hexagonal setting R3mH. Although various Al-Mn intermetallics are known to exist and three are expected to form 202 (Al₈Mn₅, Al₁₁Mn₄ and Al₄Mn) according to Scheil calculations using current thermodynamic 203 databases ^[27], the strong crystallographic differences between these phases enabled Al₈Mn₅ 204 to be clearly distinguished. Al₈Mn₅ is also consistent with the EDS measurement of 59at%Al -205 40at%Mn. Note that rhombohedral Al₈Mn₅ is also known as $\gamma_2^{[42]}$ and LT-AL₈Mn₅^[43], and is a 206 gamma brass with Strukturbericht designation D8₁₀. It is useful to index this crystal structure 207 in the non-standard body-centred rhombohedral (BCR) setting as discussed in refs. [35,41,44]. 208



Figure 5: (a) EDS spectrum from the particle in (b). (c) EBSD pattern from the same particle. (d) EBSD pattern indexed as rhombohedral AI_8Mn_5 (D8₁₀).

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Rhombohedral Al₈Mn₅ was the only Al-Mn intermetallic detected in the HPDC AZ91D 213 samples by SEM-based techniques in this work. This is reasonably consistent with Scheil 214 calculations within Thermo-Calc Software TCMG magnesium database version 4 [27] which 215 show that ~95% of all the Al-Mn phases formed during Scheil solidification are Al_8Mn_5 (using 216 217 the composition in Table 1). If Al₁₁Mn₄ and/or Al₄Mn were present in the HPDC samples, they were either too low in volume fraction or too small to be detected. The B2-Al(Mn,Fe) phase 218 identified in AZ91 in ref.^[35] was not detected in this work, most likely because the AZ91 used 219 here (Table 1) had a very low Fe content (<10 ppm). 220

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It was found that most HPDC Al₈Mn₅ particles were cyclic twinned containing up to four 222 orientations, similar to the Al₈Mn₅ particles at low cooling rate identified in ref. ^[35]. For 223 224 example, Figure 6(a) is a typical ~5 µm HPDC Al₈Mn₅ particle and Figure 6(b) is its EBSD orientation map showing the presence of three orientations within the particle. Note that 225 226 the grey pixels have unknown orientation due to low EBSD pattern quality in this region. The three orientations are plotted in pole figures in Figure 6(c) which show that all three 227 orientations share three common {100}_{BCR} planes and each orientation shares a common 228 $\{110\}_{BCR}$ plane with one of the other orientations. This orientation relationship between the 229 230 three Al₈Mn₅ orientations is shown geometrically in Figure 6(e) which is a plot of the BCR unit

cell wireframes using the EBSD-measured Euler angles and coloured consistent with Figure 231 6(b)-(c). The green orientation was not measured experimentally for this 2D section of the 232 particle but is likely to be present in the 3D particle based on the findings in our previous 233 work ^[35]. Note that the BCR unit cell of Al₈Mn₅ has rhombohedral angle ~89° ^[40,41] and so 234 appears as near-cubes in Figure 6(e). Figure 6(f) is a digital section through the geometrical 235 model in Figure 6(e). It can be seen that the Al₈Mn₅-Al₈Mn₅ interfaces in the sliced BCR 236 model have similar angular arrangement with the experimental interfaces in Figure 6(b), 237 consistent with the interfaces being $\{100\}_{BCR}$. The cyclic growth twinning of Al₈Mn₅ with 238 $\{100\}_{BCR}$ twin planes can be understood by noting that, with a rhombohedral angle of ~89° 239 ^[40,41], the crystal is pseudo-cubic which gives the possibility for growth twins with $\{100\}_{BCR}$ 240 interfaces by ~90° rotations around the three <100> $_{BCR}$ axes ^[35]. 241



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Figure 6. Cyclic growth twinning of Al_8Mn_5 particles in HPDC AZ91. (a) SEM image, (b) EBSD orientation map in RGBY colour scheme, (Grey region has unknown orientation due to low pattern quality). (c) {100}_{BCR} and {111}_{BCR} pole figures showing the three orientations. (d) band contrast map showing grain boundaries. The three BCR unit cell orientations (plus a green orientation that was not present in the cross-section). (f) {100}_{BCR} twin planes revealed by sectioning the BCR geometrical model.

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In the HPDC AZ91D sample studied here, it was found that all equiaxed polyhedral Al_8Mn_5 particles that were large enough for EBSD mapping were cyclic twinned. Comparing Figure 6 in this paper with the TEM images in Fig. 4(a) in ref ^[19] and Fig. 7(b) in ref. ^[21], it is likely that the HPDC Al_8Mn_5 particles in references ^[19,21] contain sector-twins and were also cyclic twinned, although those authors did not study or mention this.

Having confirmed that the majority of Al-Mn particles are Al_8Mn_5 by combined EDS and EBSD, Al_8Mn_5 could be distinguished in backscattered electron (BSE) images due to the much higher atomic-number of Mn compared with Mg and Al. For example, in Figure 2(c), the numerous bright particles are Al_8Mn_5 and the lighter grey particles are $Mg_{17}Al_{12}$.

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261 3.3 Al₈Mn₅ morphologies

262 The HPDC AZ91D bars contained a range of Al₈Mn₅ morphologies that could be broadly 263 classified into two categories: equiaxed-polyhedral and complex-branched particles. A representative selection is shown in Figure 7 where Figure 7(a) are equiaxed-polyhedral 264 265 morphologies, and Figure 7(b) are a range of complex-branched morphologies. Each column represents a different location in the test bars: the centre of the cross-section, the defect 266 band, and the skin. It can be seen that similar morphologies were present at each location of 267 the castings, although the size distributions were different as will be discussed in detail later 268 269 in this paper.



Figure 7: Typical range of Al_8Mn_5 morphologies in one HPDC AZ91 sample. SE-SEM images after selective etching of the α -Mg. (a) equiaxed polyhedral particles, (b) complex branched particles.

It has been shown by in-situ X-ray imaging of AZ91 solidification at low cooling rate ^[28,29], 273 that the Al₈Mn₅ particles that form in the early stages of solidification are equiaxed 274 polyhedral and it is likely, therefore, that the equiaxed-polyhedral particles in these HPDC 275 samples also formed in the earlier stages of solidification. The complex-branched particles in 276 the bottom row of Figure 7(b) may have formed relatively late during a eutectic-type 277 reaction when the remaining liquid regions were tortuous channels. This is consistent with 278 Figure 1 which shows that, for Scheil solidification, Al₈Mn₅ forms both as a primary phase 279 280 prior to α -Mg formation (the red line) and also by a eutectic-type reaction with α -Mg (the 281 green line), $L \rightarrow \alpha$ -Mg + Al₈Mn₅, over a range of temperature up to ~70% solid. However, further work is required to confirm that the complex-branched particles in the bottom row of 282 Figure 7(b) formed in this eutectic-type reaction. 283

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Past work on investment cast AZ91 reported dendritic Al₈Mn₅ near the surface ^[45]. In the 285 HPDC samples studied here, the complex-branched particles occasionally had dendritic 286 morphology (e.g. some in the top row of Figure 7(b)) but these were present at all locations 287 288 in the casting. FIB serial sectioning on one branched-faceted Al₈Mn₅ crystal with morphology similar to the top row of Figure 7(b) was conducted to explore its formation. 289 The FIB slices confirmed that, in this case, the branched structure grew from a common 290 centre. At the same time, it is also possible that other complex-branched Al₈Mn₅ similar to 291 the top row of Figure 7(b) are clusters of equiaxed-polyhedral particles that were swept 292 together during solidification. 293

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3.4 Al₈Mn₅ externally solidified crystals (ESCs)

The Al₈Mn₅ particles had a wide range of sizes spanning from <100 nm to >5 μ m, which is significantly broader than in previous work at sand-casting cooling rates. For example, in ref. ^[35], the Al₈Mn₅ particle size varied from 4-14 μ m for a cooling rate of ~1 K.s⁻¹. Figure 8(a) is a typical micrograph of a region containing Al₈Mn₅ particles with a wide size range in the HPDC samples. A ~4 μ m Al₈Mn₅ particle can be seen that is an order of magnitude larger than the numerous smaller Al₈Mn₅ particles in the surrounding material. It is likely that the large particle is an Al₈Mn₅ ESC that nucleated and grew in the shot chamber at low cooling

rate before being injected into the die cavity analogous to the α -Mg ESCs in Figure 2(d) and 303 4(a)-(b), whereas the smaller Al₈Mn₅ nucleated and grew at higher cooling rate. This can be 304 concluded based on three factors: (i) the larger (~5 μ m) Al₈Mn₅ particles in (e.g. Figures 5(a), 305 6(a) and 8(a)) are within the range of Al₈Mn₅ particle sizes reported for a cooling rate of ~1 306 K.s⁻¹ in past work ^[35], indicating that they did not form in the die cavity at high cooling rate; 307 (ii) as will be shown in the next section, the larger (~5 μ m) Al₈Mn₅ particles do not belong to 308 the same population as the smaller Al₈Mn₅ particles and the Al₈Mn₅ exhibit a multi-model 309 grain size distribution similar to α -Mg grains in HPDC parts containing α -Mg ESCs (e.g. ^[3]); 310 and (iii) Al₈Mn₅ ESCs are expected since these samples contain α -Mg ESCs (Figure 4) and 311 Al₈Mn₅ is stable above the α -Mg liquidus (Figure 1) for the composition in Table 1 ^[27]. Note 312 that abnormally large Al_8Mn_5 particles in HPDC parts can be even larger, with a 20 μ m Al_8Mn_5 313 particle found in HPDC AM50 in ref.^[20]. 314



Figure 8: (a) a typical large Al₈Mn₅ particle in HPDC AZ91D. (b-d) SE-SEM images of three Al₈Mn₅

317 particles after selective etching of α -Mg, and polyhedron models based on {100}, {110}, {112} facets 318 using a pseudo-cubic cell.

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In our previous work at sand-casting cooling rates $^{[35]}$, we identified the Al₈Mn₅ growth facets 320 using combined FIB-EBSD techniques as combinations of {100}, {110} and {112} using a 321 pseudo-cubic (pc) BCR unit cell. To explore whether the larger Al₈Mn₅ particles in these 322 HPDC samples had similar growth facets, deep etched images of Al₈Mn₅ particles were 323 explored using polyhedron models. It was found that the deep etched images could usually 324 be recreated from combinations of $\{100\}_{pc}$, $\{110\}_{pc}$ and $\{112\}_{pc}$ facets. Three such examples 325 are shown in Figure 8(b)-(d) where the models were generated by plotting the {100}, {110} 326 and {112} cubic facet families, and tuning the distance from the centroid to each facet to 327 best match the deep etched SEM images. Thus, the larger Al₈Mn₅ particles in HPDC AZ91D 328 have similar facets to sand cast AZ91 [35]. 329

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The wide range of polyhedral Al₈Mn₅ forms based on different combinations of the facet families indicates that these growth facets are sensitive to the local solidification conditions (thermal, solutal and/or kinetic) which are expected to vary substantially with time and location in the HPDC process. No simple trend of the polyhedral form of Al₈Mn₅ versus location in the HPDC part was identified in this work.

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337 3.5 3D size distributions of Al₈Mn₅ particles

Figure 9(a) shows typical 3D rendered images of Al₈Mn₅ particles from FIB tomography with 338 a 50nm slice step size. Each volume is \sim 13x13x13 μ m³ and comes from one of three 339 locations: the casting centre, the porosity band, and the skin. Figure 9 (b) show histograms 340 of the Al₈Mn₅ particle size distribution at each location. The histograms contain data from 341 342 multiple tomograms as summarised in Table 2. The size distributions are plotted in terms of the number of Al₈Mn₅ particles and in terms of the volume occupied by the Al₈Mn₅ particles, 343 separately. Two definitions of Al₈Mn₅ particle size are used: the equivalent sphere diameter 344 and the "3D length". The latter is defined as the longest Feret diameter. Note in Figure 9(a) 345 that the rendering causes the Al₈Mn₅ particles to appear rounded, but the particles are 346 actually faceted as can be seen in the typical images from FIB sectioning shown as insets in 347 the histograms of Figure 9(b). The volume fraction of Al-Mn IMCs varied from 0.11-0.22 vol. 348 % depending on the location (Table 2). This is similar to the 0.10 vol.% calculated with 349 Thermo-Calc TCMG4.0 ^[27] for the composition in Table 1, and 0.18% measured by Wang et 350

al. ^[46] for HPDC AZ91D, which shows that a sufficient volume of material has been sampled and the thresholding approach was reasonable. The particle size results in Figure 9 are in general agreement with past work using TEM on small volumes. For example, Wei et al. ^[18] reported that Al-Mn particles were 100 nm to ~ $I\mu m$ and usually less than 500 nm in AM and AZ Mg HPDC parts, and Wang et al. ^[19] reported Al₈Mn₅ to have polygonal morphology with size about 100 - 200 nm in HPDC AZ91D.



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Figure 9: Al_8Mn_5 particle size data in different locations in the HPDC cross-section based on FIBtomography. (a) Rendered images of Al_8Mn_5 particles in volumes of ~13x13x13 µm³. Each particle has a unique colour. (b) Al_8Mn_5 particle size histograms in terms of the number of particles and the volume occupied by particles. The inset micrographs are typical 2D SEM images of Al_8Mn_5 particles in each location. The scale bar is 200nm in each case.

Table 2: Summary of the Al₈Mn₅ particle size data at different locations in HPDC AZ91D extracted
 from the distributions in Figs. 9 and 10 from FIB-tomography. ESD= equivalent sphere diameter. IMC=
 intermetallic compound. (ESD>180nm particles calculated)

		Center	Band	Skin
Distance from surface	[µm]	2700-2900	1500-1600	10-20
Number of tomograms	[-]	6	3	4
Total volume sampled	[µm³]	24726	40203	25660

Number of IMCs measured	[-]	449	451	6547
Mean ESD	[nm]	432	453	163
Median ESD	[nm]	408	421	99.2
Standard deviation in ESD	[nm]	189	219	133
Maximum ESD	[nm]	2245	2245	1534
Volume of all Al-Mn IMCs	[µm ³]	35.9	44.3	55.8
Number density of particles	[µm ⁻]	0.0182	0.0112	0.2551
Volume Fraction of IMCs	[-]	0.0015	0.0011	0.0022

The distributions are analysed in more detail in Figure 10 and summarised in Table 2. Figure 10(a) are box plots showing the median, the 25th percentile and 75th percentile, and a significant tail at large size in each particle population. Note that the largest Al₈Mn₅ particle in Figure 10(a) and Table 2 is less than 2.3 μ m, which is significantly smaller than the Al₈Mn₅ particle in Figure 5(b), 6(a) and 8(a) (>4 μ m), so the tail to large size extends to even larger size than in Figure 10(a), even though the randomly-selected regions only contained particles up to ~2.3 μ m.

The particle size distribution data were compared with various distributions including 375 normal, lognormal and Weibull using probability test plots. At each location, the data were 376 best described by a lognormal distribution as shown in Figure 10(b). This is consistent with 377 378 many past studies that have shown grain size and particle size distributions are often welldescribed by a lognormal distribution (e.g. ^[3,47]) including Fe-bearing IMCs in cast aluminium 379 alloys ^[48,49]. In Figure 10(b), there is a negative deviation from the lognormal test line at large 380 particle size and at small particle size. At small size (<~200nm) this might be, at least partly, 381 due to measurement uncertainty caused by the 50nm FIB slice distance. At large particle size 382 (>~1 μ m at a cumulative probability >~99%), the negative deviation from the straight line 383 384 corresponds to Al₈Mn₅ particles larger than expected of this lognormal population. The 385 presence of this small number of abnormally large grains in the populations can also be seen in the volume occupied histograms in Figure 9(a), especially at the casting centre and near 386 the defect band. From this, and the observation of many abnormally large Al₈Mn₅ particles 387 such as that in Figure 8(a), it can be concluded that the larger Al₈Mn₅ particles do not belong 388 to the same population as the main lognormal distribution. The largest Al₈Mn₅ particles 389 390 were very likely present in the shot chamber but there may also be other size populations associated with the different cooling rate and flow regimes in the different stages of HPDC: 391 392 in the shot sleeve, the slow shot stage, the filling stage, and the intensification stage.



Figure 10: Analysis of the Al_8Mn_5 particle size data from FIB-tomography in Fig. 9 and Table 2. (a) Box plots showing the median, the 25th and 75th percentiles, and the outliers at large size. (b) lognormal probability plot to test for lognormality of the datasets at each location.

397

398 Considering now the distributions of Al₈Mn₅ particles in the three locations in the casting, it 399 can be seen in Figures 9 and 10 and Table 2, that the Al₈Mn₅ size distributions were similar in the centre and defect band regions. For example, the size distributions from the centre and 400 defect band overlap over most of the range from 1%-99% of the cumulative frequency plot 401 in Figure 10(b), and the median AI_8Mn_5 size was similar (at 414 \pm 7 nm) (Table 2). 402 Additionally, the tail at large size was similar in the centre and defect band, as can be seen in 403 the volume occupied histograms in Figure 9(a), and the similar maximum Al₈Mn₅ particle size 404 in the sampled volumes in Table 2. Thus, it is likely that the Al_8Mn_5 particle size distributions 405 are similar throughout the interior regions of the castings. 406

In contrast, the Al₈Mn₅ size distribution was markedly different in the skin with significantly 407 smaller and more numerous Al₈Mn₅ particles. For example, the Al₈Mn₅ distribution from the 408 skin is shifted to smaller size (to the left) in Figure 10(b) and the median size is smaller by a 409 410 factor of >4 in Table 2. There was also an order of magnitude higher number density (number per unit volume) of Al₈Mn₅ particles in the skin than in interior regions. This is 411 shown in Table 2 and can be seen by eye in the rendered images in Figure 9(a). This higher 412 number density is not simply due to the smaller Al₈Mn₅ size, but also because the volume 413 fraction of Al₈Mn₅ particles was higher in the skin by a factor of 1.5-2 (Table 2). 414

Although a large number of Al-Mn particles were sampled by FIB tomography in this work 415 (at least 449 in each region, Table 2), this technique is inherently limited by its small sampling 416 417 volume. To partially offset this issue, within each type of region (the skin, band or centre), we 418 selected each tomogram from different parts of the bar and sampled 3-6 tomograms (Table 2). For example, 4 tomograms were taken from randomly selected different parts of the skin, 419 and all showed a higher volume fraction and smaller size of Al₈Mn₅ than the other two 420 regions. Thus, the results in Figures 9 and 10 and Table 2 are likely to be generally valid 421 across the whole bar. Figure 3 showed large variation in porosity distribution from sample to 422 sample. From 2D backscatter electron imaging, there did not appear to be similarly large 423 424 differences in the distributions of intermetallic compounds. However, further detailed FIB 425 tomography work would be required to obtain quantitative detail on the variation in particle size distributions from sample to sample. Note that the most common porosity distribution 426 in the HPDC bars was similar to Figure 3(b), and we performed our FIB slice and view 427 characterisation and quantification on this type of sample. 428

429

430 **3.6 Correlations between Al₈Mn₅ particles and Mg₁₇Al₁₂**

In Figure 8(a), many bright Al₈Mn₅ particles appear close to eutectic regions on the 2D
section. Therefore, the 3D FIB-tomography datasets were further explored to investigate any
correlation between Al₈Mn₅ particles and eutectic Mg₁₇Al₁₂, noting from Figure 1 that Al₈Mn₅
forms before Mg₁₇Al₁₂.

435

436 In a previous FIB-tomography study on HPDC AZ91 ^[5], the eutectic $Mg_{17}AI_{12}$ was shown to 437 form an interconnected scaffold-like network in 3D. The eutectic $Mg_{17}AI_{12}$ network was more

438 profusely interconnected near the casting surface than at the casting centre which was 439 attributed to the higher fraction of large ESCs near the centre resulting in a larger length 440 scale of the Mg₁₇Al₁₂ network in the centre. A similar 3D Mg₁₇Al₁₂ microstructure was 441 measured by FIB tomography in this work as shown in Figure 11. The Mg₁₇Al₁₂ (rendered in 442 grey) forms a percolating network that is more intricately interconnected in the skin than in 443 the defect band and centre.

444

In Figure 11 the Al₈Mn₅ particles are rendered with colour, where a different colour has been 445 assigned to each distinct particle. It can be seen that some Al₈Mn₅ are in contact with 446 447 Mg₁₇Al₁₂ and many are a significant distance away from Mg₁₇Al₁₂. Noting that the 448 transparent phase is α -Mg, the numerous Al₈Mn₅ particles that are away from Mg₁₇Al₁₂ are 449 fully surrounded by α -Mg in 3D. For those Al₈Mn₅ particles that share an interface with $Mg_{17}AI_{12}$, it is not possible with the techniques used to conclude whether $Mg_{17}AI_{12}$ nucleates 450 451 on these pre-existing Al₈Mn₅ or whether Al₈Mn₅ particles are just pushed by the growth of α-Mg dendrites to the last liquid to solidify where they came into contact with Mg₁₇Al₁₂ during 452 the final eutectic solidification. Further work is required to distinguish between these 453 454 possibilities. A key finding from Figure 11 is that most Al₈Mn₅ particles do not contact Mg₁₇Al₁₂ in 3D. 455



457 Figure 11. Rendered Mg₁₇Al₁₂ eutectic (grey) and Al₈Mn₅ (colours) from the FIB-tomography datasets
458 in Figure. 9.

Comparing this HPDC study with past work at a controlled cooling rate of ~1 K s^{-1 [35]}, it can 459 be concluded that the growth crystallography and twinning of larger Al₈Mn₅ particles in 460 HPDC (Figure 6) is similar to slow cooled samples. However, the HPDC process generated a 461 much wider variation in Al₈Mn₅ size distribution, number density, and morphology due to the 462 wide range of cooling and flow conditions in the different stages of HPDC. This work has also 463 identified significant differences in the Al₈Mn₅ size distribution in the skin and interior 464 regions. The smaller particles, higher volume fraction and smaller interparticle spacing of 465 Al₈Mn₅ particles in the skin region may partially contribute to the increased hardness 466 reported in the skin ^[16]. In contrast, partial solidification in the shot sleeve ties up Mn in 467 larger Al₈Mn₅ ESCs which will reduce the number density of Al₈Mn₅ particles and reduce the 468 potential benefits that might be gained from smaller, more numerous particles. 469

470 **4** Conclusions

Al-Mn intermetallic compounds have been characterised and quantified in high pressure die cast (HPDC) AZ91D test bars to understand the types of Al-Mn phases present, their faceted growth crystallography, and their size distribution in relation to the other phases and the key microstructural features in HPDC: the skin, the defect band, and Mg₁₇Al₁₂. The following conclusion can be drawn.

- Similar to Al₈Mn₅ particles in slow cooled (~1 K/s) AZ91D samples studied previously
 ^[35], Al₈Mn₅ particles in HPDC were often cyclic twins containing four orientations with
 {100}_{BCR} twin planes. The facet morphology of large polyhedral Al₈Mn₅ particles could
 be described by combinations of {100}, {110}, and {112} facets.
- Al₈Mn₅ particles had a wide range of sizes and morphologies within the same HPDC
 component, but all could be broadly classified as equiaxed-polyhedral or complex branched.
- The great majority of Al_8Mn_5 particles were sub-micrometre in size but there was a significant population of much larger (~5 µm) polyhedral particles whose size is similar to Al_8Mn_5 particles solidified at low cooling rate (1-3 K/s). These particles are concluded to be externally solidified crystals (ESCs) that nucleated and grew in the shot chamber analogous to the α Mg ESCs.
- In all locations of the casting, the Al₈Mn₅ particle size distributions were reasonably
 well-described by lognormal distributions, accounting for the presence of an
 additional population(s) of larger grains associated with Al₈Mn₅ ESCs from the shot
 chamber.
- There were significant differences in the Al₈Mn₅ particle size and number density in the centre compared with the HPDC skin. The skin region had a median Al₈Mn₅ particle size (equivalent sphere diameter) of 99 nm, whereas the centre had a median Al₈Mn₅ size of 408 nm. The skin contained an order of magnitude higher number of Al₈Mn₅ particles per unit volume than interior regions
- 497 3D imaging showed that some Al₈Mn₅ particles were in contact with eutectic Mg₁₇Al₁₂ 498 but the majority of Al₈Mn₅ particles were surrounded by α-Mg.
- This study has shown that HPDC of AZ91D generates numerous Al_8Mn_5 particles with diameter 100-400nm and a small interparticle spacing. Partial solidification in the

501 shot sleeve ties up Mn in larger Al_8Mn_5 ESCs which reduces the number density of 502 Al_8Mn_5 particles.

503

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Al₈Mn₅ in high pressure die cast AZ91: twinning, 2 morphology and size distributions

3

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14

15 Abstract

Manganese-bearing intermetallic compounds (IMCs) are important for limiting micro-16 galvanic corrosion of magnesium-aluminium alloys and can initiate cracks under tensile load. 17 Here we use electron backscatter diffraction (EBSD), deep etching, and focussed ion beam 18 (FIB) tomography to investigate the types of Al-Mn phases present, their faceted growth 19 20 crystallography, and their three-dimensional distribution at different locations in high pressure die cast (HPDC) AZ91D. The Al-Mn particle size distributions were well-described 21 by lognormal distributions but with an additional population of externally solidified crystals 22 (ESCs) formed in the shot chamber analogous to α -Mg ESCs. The large Al₈Mn₅ particles were 23 cyclic twinned. Differences in the particle size distributions and number density in the centre 24 compared with the HPDC skin are identified, and the spatial relationship between Mg₁₇Al₁₂ 25 and Al-Mn particles is explored. 26

27 Keywords AZ91, high pressure die casting, intermetallics

28

29 Introduction

Automotive magnesium components are often Mg-Al-based alloys produced by high pressure die casting (HPDC). When conducted with an optimised die, process parameters and vacuum system ^[1,2], HPDC can mass produce large, thin-walled, complex shapes

containing microstructures with fine α -Mg grains (5-20 µm) ^[3,4], and a fine-scaled percolating eutectic Mg₁₇Al₁₂ network ^[5,6]. While a large body of research has investigated microstructure formation in Mg HPDC, including the formation of α -Mg grains ^[3,4,7], the surface 'skin' ^[4,8], the eutectic Mg₁₇Al₁₂ ^[5,9,10], and casting defects ^[11–17], less work has explored the formation of Al-Mn-(Fe) intermetallic particles ^[18–21]. These particles play an important role in determining micro-galvanic corrosion in HPDC Mg parts ^[22,23] and can initiate cracks under tensile loading ^[24,25].

40

Most Mg-Al-based HPDC alloys (e.g. AM50A, AM60B, AZ91D^[26]) contain sufficient Mn and 41 42 Al that Al_8Mn_5 begins to form before α -Mg during solidification. For example, Figure 1 shows the sequence of phase formation assuming Scheil solidification of AZ91D with the 43 44 composition in Table 1, calculated with the Thermo-Calc TCMG magnesium database version 4 ^[27]. It can be seen that Al_8Mn_5 is the first solid phase to form, and becomes stable ~44K 45 46 above the α -Mg liquidus temperature for this composition. It has been confirmed by in-situ 47 X-ray imaging that Al₈Mn₅ forms at higher temperature (i.e. earlier on cooling) than α -Mg in a similar alloy ^[28,29]. A consequence of this in HPDC is that Al₈Mn₅ can form and settle in the 48 holding pot ^[29,30], for example during temperature drops when charging the furnace with 49 new ingots, leading to die casting sludge ^[30]. Furthermore, in cold chamber HPDC, heat loss 50 51 in the shot chamber can cause Al₈Mn₅ formation prior to injection as Al₈Mn₅ externally solidified crystals (ESCs) ^[20] in addition to the α -Mg ESCs that are widespread in HPDC Mg 52 components ^[3,14,31]. This occurs because a feature of Mg HPDC is partial solidification in the 53 shot chamber that leads to large α -Mg externally solidified crystals (ESCs) being injected into 54 the cavity ^[3,32]. The volume fraction of α -Mg ESCs has been shown to depend on the melt 55 superheat, the fill fraction and the temperature of the sleeve walls and plunger tip, and is 56 typically 10-30 vol.% ^[3,14,31,33]; similar factors might be expected to determine the formation 57 of Al₈Mn₅ ESCs. 58

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Table 1. Composition of the AZ91D alloy used (weight percent).

-	Mg	Al	Zn	Mn	Fe	Ni	Cu	Si	Be
_	bal.	8.95	0.72	0.19	<0.001	<0.001	0.001	0.039	0.0007

61

Figure 1 shows that AI_8Mn_5 continues forming along with α -Mg below the α -Mg liquidus temperature until ~ 510°C when other Al-Mn IMCs start forming ($AI_{11}Mn_4$ and then AI_4Mn). Therefore, in HPDC, Al-Mn IMCs are expected to form in all stages of the process: in the shot chamber, during filling and during the intensification stage. According to calculations linked with Figure 1, at the end of Scheil solidification, the total mass fraction of Al-Mn IMCs (AI_8Mn_5 , $AI_{11}Mn_4$ and AI_4Mn) is 0.25% of which 95% is AI_8Mn_5 for the composition in Table 1.



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Figure 1: Phase formation during Scheil solidification up to 99% solid for Mg-8.95Al-0.72Zn-0.19Mn
(wt%). Calculated with Thermo-Calc TCMG magnesium database version 4 [27].

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Past work on AI-Mn particles in HPDC AZ91D has generally used TEM ^[18,19,21]. That work has 73 deduced that most Al-Mn particles in HPDC AZ91D are 100 nm to 1µm in size. The main 74 phase present has been found to be Al₈Mn₅ and another phase with higher Al content 75 (possibly $AI_{11}Mn_4$) has also been reported ^[18]. While these TEM studies enable high resolution 76 imaging, they did not explore the statistical variation in Al-Mn particle size and shape versus 77 position in the cross-section. This is an important question in HPDC parts since they usually 78 have highly non-uniform microstructures. They typically have a surface layer (a skin) of 79 distinctly different microstructure that is usually free of porosity and harder than more 80 central regions, one or more bands of porosity, various forms of macrosegregation, and ESCs 81 that tend to be concentrated towards the centre of cross-sections (e.g. ^[15,16,33,34]). 82

In this paper, we investigate the types of Al-Mn phases present, their faceted growth crystallography, and their three-dimensional distribution at different locations in high pressure die cast AZ91D. The specific aims are: (i) to compare the Al₈Mn₅ growth crystallography and twinning formed in HPDC with past work at sand casting cooling rates ^[35]; (ii) to quantify the 3D size, morphology and spatial distribution of Al-Mn particles in different locations in HPDC AZ91D: the skin, the defect band, and the centre; and (iii) to explore any correlations between Al-Mn particles and eutectic Mg₁₇Al₁₂ in 3D.

92 Methods

 93 ~6 kg of AZ91D Mg alloy with composition in Table 1 was melted in a mild steel crucible and held at 675°C (~ 75°C superheat) under a cover gas of ~3 vol% SF₆ in N₂. HPDC was conducted using a Frech DAK 450-54 cold chamber HPDC machine and the multi-cavity die that produces the casting in Figure 2. The die was preheated to 150°C, a portion of the melt was ladled into the shot chamber to a fill fraction of ~0.5, and the following set parameters were used: slow shot phase of 0.3 m.s⁻¹, fast shot phase of 4 m.s⁻¹, and intensification pressure of 36 MPa. The casting analysed in this work was made after six pre-shots.



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Figure 2 (a-b) Photographs of the HPDC part. The sectioning plane is indicated by superimposed lines.(c) as-polished optical micrograph.(d) the same section after etching.

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Samples for microstructural analysis were cut from the centre of the gauge length into slices of 10mm x 10mm x 0.5mm. Metallographic polishing was carried out down to 0.05µm colloidal silica by standard preparation methods. Some samples were etched in a solution of 200ml ethylene glycol, 68ml distilled water, 4ml nitric acid and 80 ml acetic acid. Both etched and polished samples were analysed in a Zeiss AURIGA field emission gun SEM (FEG-SEM)

with an Oxford Instruments INCA x-sight energy dispersive X-ray spectroscopy (EDX) 109 detector and a BRUKER e-Flash^{HR} electron backscatter diffraction (EBSD) detector. For EBSD 110 characterisation, the final step of preparation was Ar-ion milling for 40 min in a Gatan PECSII 111 112 instrument. The 4kV-accelerated beam hit the sample rotating at 2rpm, at a grazing incidence angle of 4°. Electron beam accelerating voltage of 20kV, working distance of 113 15mm, aperture size of 120mm, and beam current 80µA were used for EBSD measurements. 114 Bruker ESPRIT 2.1 software was used to index the obtained EBSD patterns. EBSD datasets 115 were analysed using MATLAB[™] 9.2 (Mathworks, USA) with the MTEX 5.1 toolbox ^[36]. 116 Accelerating voltage of 10kV, working distance of 5mm, aperture size of 60mm, and beam 117 118 current 80µA were used for EDS analysis. EDS spectrum was calibrated with a Si standard 119 sample prior to each electron microscopy session.

To investigate the 3-dimensional (3D) morphology of the Al-Mn intermetallics directly, α -Mg 120 was selectively etched using a solution of 4% nitric acid in ethanol. To quantify the 3D size 121 distribution of Al-Mn intermetallics, focussed ion beam (FIB) tomography was conducted in a 122 Zeiss AURIGA FG-SEM at 30 kV with 52° tilt angle. The slice distance was 90 nm and the 123 milling current was 200pA. Serial-sectioning secondary electron images were used. For FIB 124 tomography, 2D slices were aligned, cropped, and processed by an anisotropic diffusion filter 125 126 in ImageJ (US NIH, USA). 3D reconstruction and crystallographic analysis was performed using Avizo 9.2 (Visualization Science Group, France) and MATLAB 9.2[™]. The voxel size for 127 FIB tomography was bounded by the slice spacing of 90nm. Al₈Mn₅ particles with equivalent 128 diameter \geq 180nm were quantified. 129

To study porosity bands in 3D, X-ray micro-tomography was carried out on a North Star 130 Imaging (NSI) Micro-CT. The system is equipped with a 225 kV X-ray source with a minimum 131 focal spot size of 2 µm and a Perkin Elmer flat panel detector (2048×2048 pixels at 16bit 132 depth). During a CT scan, the sample was illuminated by cone beam X-rays which were 133 transmitted through the 360° rotating specimen and then illuminated on the flat panel 134 135 detector. The X-ray beam was filtered using a 0.25 mm Cu filter to reduce beam-hardening effects, and an acceleration voltage of 80kV and target current of 35µA was selected to 136 optimise image quality. 1440 two-dimensional projections were captured over 360° with an 137 exposure time of 1000ms. 3D reconstruction was performed in Avizo 9.2 and resulted in a 3D 138 spatial resolution with voxel size of 2.2 µm x 2.2 µm x 2.2 µm. 139

140 **3 Results and Discussion**

141 **3.1 General microstructural features**

At the centre of the gauge length, the AZ91D samples contained the typical microstructural 142 features and defects of HPDC reported in past work (e.g. [3,7,12,14,16,33,34,37]). For example, 143 annular rings of porosity can be seen in the as-polished condition in Figure 2(c), a dark band 144 of macrosegregation can be seen in the same location as the main porosity band in Figure 145 2(d) after light etching, and a high fraction (~30 vol%) of α -Mg ESCs can be seen throughout 146 147 much of the cross-section in Figure 2(d). However, the detail of these features differed significantly from casting to casting and between bars in the same casting as shown in the X-148 ray tomographs in Figure 3. The left-hand images are reconstructed volumes near the centre 149 of the gauge length showing the 3D distribution of porosity. The right-hand images are 150 viewed along the tensile rod axis to highlight the radial distribution of porosity. There are 151 major differences in the porosity in the two samples. The sample in Figure 3(b) has a 152 localised annular ring of porosity and a high fraction of porosity within this ring. The sample 153 154 in Figure 3(a) has more diffuse porosity and a less-well defined porosity ring but has the same trend of a higher fraction of porosity within the annular porosity band. Despite the 155 differences, in both samples, the main annular ring of porosity is at a similar radial position. 156 The projection images along the rod axes also reveal the surface 'skin' as an outer ring of 157 essentially zero porosity. This is particularly clear in Figure 3(a) where the abrupt change in 158 porosity demarcates the edge of the skin. 159



Figure 3 (a-b) X-ray tomograms of porosity near the centre of the gauge-length of typical castings.
Porosity is rendered as solid, material (Mg, Mg₁₇Al₁₂ and Al-Mn IMCs) is plotted as semi-transparent.
Left-hand side: perspective view. Right-hand side: projection view along the tensile rod axis.

164

165 The typical α -Mg microstructure is shown in more detail in Figure 4(a)-(b). The micrograph 166 in Figure 4(a) shows the complex mixture of dendritic α -Mg ESCs, ESC fragments and in-167 cavity solidified grains. Figure 4(b) is an EBSD orientation map (IPF-y) of the α -Mg phase 168 from a similar region where the grains have been coloured by their mean-orientation. The 169 grains form a complex multimodal microstructure with, in this case, two large ESCs 170 surrounded by smaller α -Mg grains that are probably a mixture of α -Mg ESC fragments and 171 in-cavity solidified grains.

172

173 The typical features of intermetallic compounds in the HPDC bars are overviewed in Figure 174 4(c) and (d). It can be seen that the eutectic Mg₁₇Al₁₂ phase appears as isolated regions in 2D 175 sections (Figure 4(c)) but actually forms a percolating Mg₁₇Al₁₂ network in 3D as revealed by 176 imaging after selective dissolution of the α -Mg in Figure 4(d). Figure 4(c) and (d) also 177 contains bright particles that are Al-Mn compounds. In the 2D section these appear both 178 within the α -Mg grains and near the Mg₁₇Al₁₂ phase (Figure 4(c)). After deep etching, it can 179 be seen that many Al-Mn particles are attached to the Mg₁₇Al₁₂ network (Figure 4 (d)).



Figure 4: Typical microstructural features in the HPDC AZ91 samples. (a) mixture of α -Mg ESCs and in-cavity solidified grains. (b) EBSD orientation map (IPF-Y) of the α -Mg phase. (c) 2D section of Mg₁₇Al₁₂ and Al₈Mn₅ phases. (d) 3D microstructure of Mg₁₇Al₁₂ network and attached Al₈Mn₅ particles, revealed after selective etching of α -Mg.

186 The remainder of this paper focuses on the Al-Mn intermetallic compounds and their 187 relationship to the microstructural features summarised in this section.

188

189 **3.2 Twinned Al₈Mn₅ in HPDC AZ91D**

Al-Mn intermetallics were identified by combining EDS with EBSD. A typical EDS point 190 analysis from an Al-Mn particle is shown in Figure 5(a). The particle contains 59at%Al -191 40at%Mn and there are also small Mg, Si and Fe peaks, each present at less than 1 at%. 192 Since the solubility of Mg in Al-Mn intermetallics is negligible ^[38], the small Mg peak is likely 193 to be α -Mg in the interaction volume. The small Si peak is probably Si dissolved in the 194 particle, consistent with past work that has detected a small Si content in Al-Mn IMCs ^[18,39]. 195 The low Fe content in the particle is due to the high-purity AZ91D used in this study (with 196 <10ppm Fe, Table 1). 197

198

An EBSD pattern from the Al-Mn particle is shown in Figure 5(b). This could be readily 199 distinguished as the rhombohedral Al_8Mn_5 phase ^[40,41] using the Hough transform-based 200 201 method in Bruker ESPRIT 2.1, and is indexed in Figure 5(c) in the hexagonal setting R3mH. Although various Al-Mn intermetallics are known to exist and three are expected to form 202 (Al₈Mn₅, Al₁₁Mn₄ and Al₄Mn) according to Scheil calculations using current thermodynamic 203 databases ^[27], the strong crystallographic differences between these phases enabled Al₈Mn₅ 204 to be clearly distinguished. Al₈Mn₅ is also consistent with the EDS measurement of 59at%Al -205 40at%Mn. Note that rhombohedral Al₈Mn₅ is also known as $\gamma_2^{[42]}$ and LT-AL₈Mn₅^[43], and is a 206 gamma brass with Strukturbericht designation D8₁₀. It is useful to index this crystal structure 207 in the non-standard body-centred rhombohedral (BCR) setting as discussed in refs. [35,41,44]. 208



Figure 5: (a) EDS spectrum from the particle in (b). (c) EBSD pattern from the same particle. (d) EBSD pattern indexed as rhombohedral AI_8Mn_5 (D8₁₀).

212

Rhombohedral Al₈Mn₅ was the only Al-Mn intermetallic detected in the HPDC AZ91D 213 samples by SEM-based techniques in this work. This is reasonably consistent with Scheil 214 calculations within Thermo-Calc Software TCMG magnesium database version 4 [27] which 215 show that ~95% of all the Al-Mn phases formed during Scheil solidification are Al_8Mn_5 (using 216 217 the composition in Table 1). If Al₁₁Mn₄ and/or Al₄Mn were present in the HPDC samples, they were either too low in volume fraction or too small to be detected. The B2-Al(Mn,Fe) phase 218 identified in AZ91 in ref.^[35] was not detected in this work, most likely because the AZ91 used 219 here (Table 1) had a very low Fe content (<10 ppm). 220

221

It was found that most HPDC Al₈Mn₅ particles were cyclic twinned containing up to four 222 orientations, similar to the Al₈Mn₅ particles at low cooling rate identified in ref. ^[35]. For 223 224 example, Figure 6(a) is a typical ~5 µm HPDC Al₈Mn₅ particle and Figure 6(b) is its EBSD orientation map showing the presence of three orientations within the particle. Note that 225 226 the grey pixels have unknown orientation due to low EBSD pattern quality in this region. The three orientations are plotted in pole figures in Figure 6(c) which show that all three 227 orientations share three common {100}_{BCR} planes and each orientation shares a common 228 $\{110\}_{BCR}$ plane with one of the other orientations. This orientation relationship between the 229 230 three Al₈Mn₅ orientations is shown geometrically in Figure 6(e) which is a plot of the BCR unit

cell wireframes using the EBSD-measured Euler angles and coloured consistent with Figure 231 6(b)-(c). The green orientation was not measured experimentally for this 2D section of the 232 particle but is likely to be present in the 3D particle based on the findings in our previous 233 work ^[35]. Note that the BCR unit cell of Al₈Mn₅ has rhombohedral angle ~89° ^[40,41] and so 234 appears as near-cubes in Figure 6(e). Figure 6(f) is a digital section through the geometrical 235 model in Figure 6(e). It can be seen that the Al₈Mn₅-Al₈Mn₅ interfaces in the sliced BCR 236 model have similar angular arrangement with the experimental interfaces in Figure 6(b), 237 consistent with the interfaces being $\{100\}_{BCR}$. The cyclic growth twinning of Al₈Mn₅ with 238 $\{100\}_{BCR}$ twin planes can be understood by noting that, with a rhombohedral angle of ~89° 239 ^[40,41], the crystal is pseudo-cubic which gives the possibility for growth twins with $\{100\}_{BCR}$ 240 interfaces by ~90° rotations around the three <100> $_{BCR}$ axes ^[35]. 241



242

Figure 6. Cyclic growth twinning of Al_8Mn_5 particles in HPDC AZ91. (a) SEM image, (b) EBSD orientation map in RGBY colour scheme, (Grey region has unknown orientation due to low pattern quality). (c) {100}_{BCR} and {111}_{BCR} pole figures showing the three orientations. (d) band contrast map showing grain boundaries. The three BCR unit cell orientations (plus a green orientation that was not present in the cross-section). (f) {100}_{BCR} twin planes revealed by sectioning the BCR geometrical model.

249

In the HPDC AZ91D sample studied here, it was found that all equiaxed polyhedral Al_8Mn_5 particles that were large enough for EBSD mapping were cyclic twinned. Comparing Figure 6 in this paper with the TEM images in Fig. 4(a) in ref ^[19] and Fig. 7(b) in ref. ^[21], it is likely that the HPDC Al_8Mn_5 particles in references ^[19,21] contain sector-twins and were also cyclic twinned, although those authors did not study or mention this.

Having confirmed that the majority of Al-Mn particles are Al_8Mn_5 by combined EDS and EBSD, Al_8Mn_5 could be distinguished in backscattered electron (BSE) images due to the much higher atomic-number of Mn compared with Mg and Al. For example, in Figure 2(c), the numerous bright particles are Al_8Mn_5 and the lighter grey particles are $Mg_{17}Al_{12}$.

260

261 3.3 Al₈Mn₅ morphologies

262 The HPDC AZ91D bars contained a range of Al₈Mn₅ morphologies that could be broadly 263 classified into two categories: equiaxed-polyhedral and complex-branched particles. A representative selection is shown in Figure 7 where Figure 7(a) are equiaxed-polyhedral 264 morphologies, and Figure 7(b) are a range of complex-branched morphologies. Each column 265 represents a different location in the test bars: the centre of the cross-section, the defect 266 band, and the skin. It can be seen that similar morphologies were present at each location of 267 the castings, although the size distributions were different as will be discussed in detail later 268 269 in this paper.



Figure 7: Typical range of Al_8Mn_5 morphologies in one HPDC AZ91 sample. SE-SEM images after selective etching of the α -Mg. (a) equiaxed polyhedral particles, (b) complex branched particles.

It has been shown by in-situ X-ray imaging of AZ91 solidification at low cooling rate ^[28,29], 273 that the Al₈Mn₅ particles that form in the early stages of solidification are equiaxed 274 polyhedral and it is likely, therefore, that the equiaxed-polyhedral particles in these HPDC 275 samples also formed in the earlier stages of solidification. The complex-branched particles in 276 the bottom row of Figure 7(b) may have formed relatively late during a eutectic-type 277 reaction when the remaining liquid regions were tortuous channels. This is consistent with 278 Figure 1 which shows that, for Scheil solidification, Al_8Mn_5 forms both as a primary phase 279 280 prior to α -Mg formation (the red line) and also by a eutectic-type reaction with α -Mg (the green line), L $\rightarrow \alpha$ -Mg + Al₈Mn₅, over a range of temperature up to ~70% solid. However, 281 further work is required to confirm that the complex-branched particles in the bottom row of 282 Figure 7(b) formed in this eutectic-type reaction. 283

284

Past work on investment cast AZ91 reported dendritic Al₈Mn₅ near the surface ^[45]. In the 285 HPDC samples studied here, the complex-branched particles occasionally had dendritic 286 morphology (e.g. some in the top row of Figure 7(b)) but these were present at all locations 287 288 in the casting. FIB serial sectioning on one branched-faceted Al₈Mn₅ crystal with morphology similar to the top row of Figure 7(b) was conducted to explore its formation. 289 The FIB slices confirmed that, in this case, the branched structure grew from a common 290 centre. At the same time, it is also possible that other complex-branched Al₈Mn₅ similar to 291 the top row of Figure 7(b) are clusters of equiaxed-polyhedral particles that were swept 292 293 together during solidification.

294

3.4 Al₈Mn₅ externally solidified crystals (ESCs)

The Al₈Mn₅ particles had a wide range of sizes spanning from <100 nm to >5 μ m, which is significantly broader than in previous work at sand-casting cooling rates. For example, in ref. ^[35], the Al₈Mn₅ particle size varied from 4-14 μ m for a cooling rate of ~1 K.s⁻¹. Figure 8(a) is a typical micrograph of a region containing Al₈Mn₅ particles with a wide size range in the HPDC samples. A ~4 μ m Al₈Mn₅ particle can be seen that is an order of magnitude larger than the numerous smaller Al₈Mn₅ particles in the surrounding material. It is likely that the large particle is an Al₈Mn₅ ESC that nucleated and grew in the shot chamber at low cooling

rate before being injected into the die cavity analogous to the α -Mg ESCs in Figure 2(d) and 303 4(a)-(b), whereas the smaller Al₈Mn₅ nucleated and grew at higher cooling rate. This can be 304 concluded based on three factors: (i) the larger (~5 μ m) Al₈Mn₅ particles in (e.g. Figures 5(a), 305 6(a) and 8(a)) are within the range of Al₈Mn₅ particle sizes reported for a cooling rate of ~1 306 K.s⁻¹ in past work ^[35], indicating that they did not form in the die cavity at high cooling rate; 307 (ii) as will be shown in the next section, the larger (~5 μ m) Al₈Mn₅ particles do not belong to 308 the same population as the smaller Al₈Mn₅ particles and the Al₈Mn₅ exhibit a multi-model 309 grain size distribution similar to α -Mg grains in HPDC parts containing α -Mg ESCs (e.g. ^[3]); 310 and (iii) Al₈Mn₅ ESCs are expected since these samples contain α -Mg ESCs (Figure 4) and 311 Al₈Mn₅ is stable above the α -Mg liquidus (Figure 1) for the composition in Table 1 ^[27]. Note 312 that abnormally large Al_8Mn_5 particles in HPDC parts can be even larger, with a 20 μ m Al_8Mn_5 313 particle found in HPDC AM50 in ref.^[20]. 314



Figure 8: (a) a typical large Al₈Mn₅ particle in HPDC AZ91D. (b-d) SE-SEM images of three Al₈Mn₅

317 particles after selective etching of α -Mg, and polyhedron models based on {100}, {110}, {112} facets 318 using a pseudo-cubic cell.

319

In our previous work at sand-casting cooling rates $^{[35]}$, we identified the Al₈Mn₅ growth facets 320 using combined FIB-EBSD techniques as combinations of {100}, {110} and {112} using a 321 pseudo-cubic (pc) BCR unit cell. To explore whether the larger Al₈Mn₅ particles in these 322 HPDC samples had similar growth facets, deep etched images of Al₈Mn₅ particles were 323 explored using polyhedron models. It was found that the deep etched images could usually 324 be recreated from combinations of $\{100\}_{pc}$, $\{110\}_{pc}$ and $\{112\}_{pc}$ facets. Three such examples 325 are shown in Figure 8(b)-(d) where the models were generated by plotting the {100}, {110} 326 and {112} cubic facet families, and tuning the distance from the centroid to each facet to 327 best match the deep etched SEM images. Thus, the larger Al₈Mn₅ particles in HPDC AZ91D 328 have similar facets to sand cast AZ91 [35]. 329

330

The wide range of polyhedral Al₈Mn₅ forms based on different combinations of the facet families indicates that these growth facets are sensitive to the local solidification conditions (thermal, solutal and/or kinetic) which are expected to vary substantially with time and location in the HPDC process. No simple trend of the polyhedral form of Al₈Mn₅ versus location in the HPDC part was identified in this work.

336

337 3.5 3D size distributions of Al₈Mn₅ particles

Figure 9(a) shows typical 3D rendered images of Al₈Mn₅ particles from FIB tomography with 338 a 50nm slice step size. Each volume is \sim 13x13x13 μ m³ and comes from one of three 339 locations: the casting centre, the porosity band, and the skin. Figure 9 (b) show histograms 340 of the Al₈Mn₅ particle size distribution at each location. The histograms contain data from 341 342 multiple tomograms as summarised in Table 2. The size distributions are plotted in terms of the number of Al₈Mn₅ particles and in terms of the volume occupied by the Al₈Mn₅ particles, 343 separately. Two definitions of Al₈Mn₅ particle size are used: the equivalent sphere diameter 344 and the "3D length". The latter is defined as the longest Feret diameter. Note in Figure 9(a) 345 that the rendering causes the Al₈Mn₅ particles to appear rounded, but the particles are 346 actually faceted as can be seen in the typical images from FIB sectioning shown as insets in 347 the histograms of Figure 9(b). The volume fraction of Al-Mn IMCs varied from 0.11-0.22 vol. 348 % depending on the location (Table 2). This is similar to the 0.10 vol.% calculated with 349 Thermo-Calc TCMG4.0 ^[27] for the composition in Table 1, and 0.18% measured by Wang et 350

al. ^[46] for HPDC AZ91D, which shows that a sufficient volume of material has been sampled and the thresholding approach was reasonable. The particle size results in Figure 9 are in general agreement with past work using TEM on small volumes. For example, Wei et al. ^[18] reported that Al-Mn particles were 100 nm to ~ μ m and usually less than 500 nm in AM and AZ Mg HPDC parts, and Wang et al. ^[19] reported Al₈Mn₅ to have polygonal morphology with size about 100 - 200 nm in HPDC AZ91D.



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Figure 9: Al_8Mn_5 particle size data in different locations in the HPDC cross-section based on FIBtomography. (a) Rendered images of Al_8Mn_5 particles in volumes of ~13x13x13 µm³. Each particle has a unique colour. (b) Al_8Mn_5 particle size histograms in terms of the number of particles and the volume occupied by particles. The inset micrographs are typical 2D SEM images of Al_8Mn_5 particles in each location. The scale bar is 200nm in each case.

Table 2: Summary of the Al₈Mn₅ particle size data at different locations in HPDC AZ91D extracted
 from the distributions in Figs. 9 and 10 from FIB-tomography. ESD= equivalent sphere diameter. IMC=
 intermetallic compound. (ESD>180nm particles calculated)

		Center	Band	Skin
Distance from surface	<mark>[µm]</mark>	2700-2900	1500-1600	10-20
Number of tomograms	[-]	6	3	4
Total volume sampled	<mark>[µm³]</mark>	24726	40203	25660

Number of IMCs measured	<mark>[-]</mark>	449	451	6547
Mean ESD	[nm]	432	453	163
Median ESD	[nm]	408	421	99.2
Standard deviation in ESD	[nm]	189	219	133
Maximum ESD	[nm]	2245	2245	1534
Volume of all Al-Mn IMCs	[µm ³]	35.9	44.3	55.8
Number density of particles	[µm ^{_3}]	0.0182	0.0112	0.2551
Volume Fraction of IMCs	<mark>[-]</mark>	0.0015	0.0011	0.0022

The distributions are analysed in more detail in Figure 10 and summarised in Table 2. Figure 10(a) are box plots showing the median, the 25th percentile and 75th percentile, and a significant tail at large size in each particle population. Note that the largest Al₈Mn₅ particle in Figure 10(a) and Table 2 is less than 2.3 μ m, which is significantly smaller than the Al₈Mn₅ particle in Figure 5(b), 6(a) and 8(a) (>4 μ m), so the tail to large size extends to even larger size than in Figure 10(a), even though the randomly-selected regions only contained particles up to ~2.3 μ m.

The particle size distribution data were compared with various distributions including 375 376 normal, lognormal and Weibull using probability test plots. At each location, the data were 377 best described by a lognormal distribution as shown in Figure 10(b). This is consistent with many past studies that have shown grain size and particle size distributions are often well-378 described by a lognormal distribution (e.g. ^[3,47]) including Fe-bearing IMCs in cast aluminium 379 alloys ^[48,49]. In Figure 10(b), there is a negative deviation from the lognormal test line at large 380 particle size and at small particle size. At small size (<~200nm) this might be, at least partly, 381 due to measurement uncertainty caused by the 50nm FIB slice distance. At large particle size 382 (>~1 μ m at a cumulative probability >~99%), the negative deviation from the straight line 383 384 corresponds to Al₈Mn₅ particles larger than expected of this lognormal population. The presence of this small number of abnormally large grains in the populations can also be seen 385 in the volume occupied histograms in Figure 9(a), especially at the casting centre and near 386 the defect band. From this, and the observation of many abnormally large Al₈Mn₅ particles 387 such as that in Figure 8(a), it can be concluded that the larger Al₈Mn₅ particles do not belong 388 to the same population as the main lognormal distribution. The largest Al₈Mn₅ particles 389 were very likely present in the shot chamber but there may also be other size populations 390

- associated with the different cooling rate and flow regimes in the different stages of HPDC:
- in the shot sleeve, the slow shot stage, the filling stage, and the intensification stage.



Figure 10: Analysis of the AI_8Mn_5 particle size data from FIB-tomography in Fig. 9 and Table 2. (a) Box plots showing the median, the 25th and 75th percentiles, and the outliers at large size. (b) lognormal probability plot to test for lognormality of the datasets at each location.

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Considering now the distributions of Al_8Mn_5 particles in the three locations in the casting, it can be seen in Figures 9 and 10 and Table 2, that the Al_8Mn_5 size distributions were similar in the centre and defect band regions. For example, the size distributions from the centre and defect band overlap over most of the range from 1%-99% of the cumulative frequency plot in Figure 10(b), and the median Al_8Mn_5 size was similar (at 414 ±7 nm) (Table 2). Additionally, the tail at large size was similar in the centre and defect band, as can be seen in the volume occupied histograms in Figure 9(a), and the similar maximum Al_8Mn_5 particle size in the sampled volumes in Table 2. Thus, it is likely that the AI_8Mn_5 particle size distributions are similar throughout the interior regions of the castings.

In contrast, the Al₈Mn₅ size distribution was markedly different in the skin with significantly 407 smaller and more numerous Al₈Mn₅ particles. For example, the Al₈Mn₅ distribution from the 408 409 skin is shifted to smaller size (to the left) in Figure 10(b) and the median size is smaller by a factor of >4 in Table 2. There was also an order of magnitude higher number density 410 (number per unit volume) of Al₈Mn₅ particles in the skin than in interior regions. This is 411 412 shown in Table 2 and can be seen by eye in the rendered images in Figure 9(a). This higher number density is not simply due to the smaller Al_8Mn_5 size, but also because the volume 413 414 fraction of Al_8Mn_5 particles was higher in the skin by a factor of 1.5-2 (Table 2).

Although a large number of Al-Mn particles were sampled by FIB tomography in this work 415 (at least 449 in each region, Table 2), this technique is inherently limited by its small sampling 416 volume. To partially offset this issue, within each type of region (the skin, band or centre), we 417 selected each tomogram from different parts of the bar and sampled 3-6 tomograms (Table 418 2). For example, 4 tomograms were taken from randomly selected different parts of the skin, 419 and all showed a higher volume fraction and smaller size of Al₈Mn₅ than the other two 420 regions. Thus, the results in Figures 9 and 10 and Table 2 are likely to be generally valid 421 422 across the whole bar. Figure 3 showed large variation in porosity distribution from sample to 423 sample. From 2D backscatter electron imaging, there did not appear to be similarly large differences in the distributions of intermetallic compounds. However, further detailed FIB 424 tomography work would be required to obtain quantitative detail on the variation in particle 425 size distributions from sample to sample. Note that the most common porosity distribution 426 in the HPDC bars was similar to Figure 3(b), and we performed our FIB slice and view 427 characterisation and quantification on this type of sample. 428

429

430 **3.6 Correlations between Al₈Mn₅ particles and Mg₁₇Al₁₂**

In Figure 8(a), many bright Al₈Mn₅ particles appear close to eutectic regions on the 2D
section. Therefore, the 3D FIB-tomography datasets were further explored to investigate any
correlation between Al₈Mn₅ particles and eutectic Mg₁₇Al₁₂, noting from Figure 1 that Al₈Mn₅
forms before Mg₁₇Al₁₂.

In a previous FIB-tomography study on HPDC AZ91^[5], the eutectic Mg₁₇Al₁₂ was shown to 436 form an interconnected scaffold-like network in 3D. The eutectic Mg₁₇Al₁₂ network was more 437 438 profusely interconnected near the casting surface than at the casting centre which was attributed to the higher fraction of large ESCs near the centre resulting in a larger length 439 scale of the Mg₁₇Al₁₂ network in the centre. A similar 3D Mg₁₇Al₁₂ microstructure was 440 measured by FIB tomography in this work as shown in Figure 11. The Mg₁₇Al₁₂ (rendered in 441 grey) forms a percolating network that is more intricately interconnected in the skin than in 442 the defect band and centre. 443

444

445 In Figure 11 the Al₈Mn₅ particles are rendered with colour, where a different colour has been 446 assigned to each distinct particle. It can be seen that some Al₈Mn₅ are in contact with Mg₁₇Al₁₂ and many are a significant distance away from Mg₁₇Al₁₂. Noting that the 447 448 transparent phase is α -Mg, the numerous Al₈Mn₅ particles that are away from Mg₁₇Al₁₂ are 449 fully surrounded by α -Mg in 3D. For those Al₈Mn₅ particles that share an interface with Mg₁₇Al₁₂, it is not possible with the techniques used to conclude whether Mg₁₇Al₁₂ nucleates 450 451 on these pre-existing Al₈Mn₅ or whether Al₈Mn₅ particles are just pushed by the growth of α-Mg dendrites to the last liquid to solidify where they came into contact with Mg₁₇Al₁₂ during 452 453 the final eutectic solidification. Further work is required to distinguish between these possibilities. A key finding from Figure 11 is that most Al₈Mn₅ particles do not contact 454 Mg₁₇Al₁₂ in 3D. 455

21



457 Figure 11. Rendered $Mg_{17}AI_{12}$ eutectic (grey) and AI_8Mn_5 (colours) from the FIB-tomography datasets 458 in Figure 9.

Comparing this HPDC study with past work at a controlled cooling rate of ~1 K s^{-1 [35]}, it can 459 be concluded that the growth crystallography and twinning of larger Al₈Mn₅ particles in 460 HPDC (Figure 6) is similar to slow cooled samples. However, the HPDC process generated a 461 much wider variation in Al₈Mn₅ size distribution, number density, and morphology due to the 462 wide range of cooling and flow conditions in the different stages of HPDC. This work has also 463 identified significant differences in the Al₈Mn₅ size distribution in the skin and interior 464 regions. The smaller particles, higher volume fraction and smaller interparticle spacing of 465 Al₈Mn₅ particles in the skin region may partially contribute to the increased hardness 466 reported in the skin ^[16]. In contrast, partial solidification in the shot sleeve ties up Mn in 467 larger Al₈Mn₅ ESCs which will reduce the number density of Al₈Mn₅ particles and reduce the 468 potential benefits that might be gained from smaller, more numerous particles. 469

470 **4** Conclusions

Al-Mn intermetallic compounds have been characterised and quantified in high pressure die cast (HPDC) AZ91D test bars to understand the types of Al-Mn phases present, their faceted growth crystallography, and their size distribution in relation to the other phases and the key microstructural features in HPDC: the skin, the defect band, and Mg₁₇Al₁₂. The following conclusion can be drawn.

- Similar to Al₈Mn₅ particles in slow cooled (~1 K/s) AZ91D samples studied previously
 ^[35], Al₈Mn₅ particles in HPDC were often cyclic twins containing four orientations with
 {100}_{BCR} twin planes. The facet morphology of large polyhedral Al₈Mn₅ particles could
 be described by combinations of {100}, {110}, and {112} facets.
- Al₈Mn₅ particles had a wide range of sizes and morphologies within the same HPDC
 component, but all could be broadly classified as equiaxed-polyhedral or complex branched.
- The great majority of Al_8Mn_5 particles were sub-micrometre in size but there was a significant population of much larger (~5 µm) polyhedral particles whose size is similar to Al_8Mn_5 particles solidified at low cooling rate (1-3 K/s). These particles are concluded to be externally solidified crystals (ESCs) that nucleated and grew in the shot chamber analogous to the α Mg ESCs.
- In all locations of the casting, the Al₈Mn₅ particle size distributions were reasonably
 well-described by lognormal distributions, accounting for the presence of an
 additional population(s) of larger grains associated with Al₈Mn₅ ESCs from the shot
 chamber.
- There were significant differences in the Al₈Mn₅ particle size and number density in the centre compared with the HPDC skin. The skin region had a median Al₈Mn₅ particle size (equivalent sphere diameter) of 99 nm, whereas the centre had a median Al₈Mn₅ size of 408 nm. The skin contained an order of magnitude higher number of Al₈Mn₅ particles per unit volume than interior regions
- 497 3D imaging showed that some Al₈Mn₅ particles were in contact with eutectic Mg₁₇Al₁₂ 498 but the majority of Al₈Mn₅ particles were surrounded by α-Mg.
- This study has shown that HPDC of AZ91D generates numerous Al_8Mn_5 particles with diameter 100-400nm and a small interparticle spacing. Partial solidification in the

501 shot sleeve ties up Mn in larger Al_8Mn_5 ESCs which reduces the number density of 502 Al_8Mn_5 particles.

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