Nucleation Behavior of a Single Al-20Si Particle Rapidly Solidified in a Fast Scanning Calorimeter

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Abstract: Understanding the rapid solidification behavior characteristics, nucleation undercooling, and nucleation mechanism is important for modifying the microstructures and properties of metal alloys. In order to investigate the rapid solidification behavior in-situ, accurate measurements of nucleation undercooling and cooling rate are required in most rapid solidification processes, e.g., in additive manufacturing (AM). In this study, differential fast scanning calorimetry (DFSC) was applied to investigate the nucleation kinetics in a single micro-sized Al-20Si (mass%) particle under a controlled cooling rate of 5000 K/s. The nucleation rates of primary Si and secondary α-Al phases were calculated by a statistical analysis of 300 identical melting/solidification experiments. Applying a model based on the classical nucleation theory (CNT) together with available thermodynamic data, two different heterogeneous nucleation mechanisms of primary Si and secondary α-Al were proposed, i.e., surface heterogeneous nucleation for primary Si and interface heterogeneous nucleation for secondary α-Al. The present study introduces a practical method for a detailed investigation of rapid solidification behavior of metal particles to distinguish surface and interface nucleation.

Keywords: nucleation kinetics; rapid solidification; differential fast scanning calorimetry (DFSC); Al-20Si particle

1. Introduction

In the production chains of metal components, e.g., in spray forming, rapid solidification of metal particles is a critical process. Recently, further interest has been gained in rapid solidification as an important step in additive manufacturing (AM) [1–5]. As important engineering materials, aluminum-based alloys, e.g., Al-Si alloys, are frequently used in AM [6–9]. In AM processes, the cooling rates are typically from ca. 100 K/s up to 10⁶ K/s [5,10–13]. For example, in laser assisted powder bed fusion (LPBF), the cooling rate of Al-12Si (mass%) alloy was estimated via simulations, which was above 10⁶ K/s for most parts of the melt pool [11].

Undercooling and nucleation in solidification determine many important aspects of the relevant microstructural features, e.g., the grain size, and the final material properties [14–17]. Studies regarding the nucleation undercooling of Al-Si alloy particles during rapid solidification processes has been reported [15,18–21]. However, in-situ observation of the rapid solidification behavior of metal particles is not easily available due to limitations of...
the measurement capability. For an accurate description of the microstructure evolution during rapid solidification, the nucleation undercooling and nucleation mechanisms are crucial, as they provide access to the optimization of the process parameters. Moreover, accurate measurements of nucleation undercooling and cooling rate, as well as accurate descriptions of rapid solidification mechanisms are required for a better understanding of rapid solidification processes of metal particles at high cooling rates.

In this study, the rapid solidification of Al-20Si (mass%) alloy was investigated by differential fast scanning calorimetry (DFSC) [22] at 5000 K/s cooling, which is in the range of the cooling rate in AM. Micro-sized Al-20Si particles with a diameter of 25 µm were used. A series of 300 nucleation events (300 heating–cooling cycles) were observed using DFSC, and the nucleation kinetics in a single particle was studied by statistical analysis. By combining microstructure and DFSC investigation, the nucleation mechanisms of the primary Si and secondary α-Al phases in a rapidly solidified single Al-20Si particle is discussed. Assuming surface and interface nucleation in a spherical cavity for Si and α-Al respectively, a modified theoretical model based on classical nucleation theory (CNT) is developed.

2. Materials and Methods

In the present work, gas-atomized Al-20Si (mass%) powder particles were used, which is purchased from Nanoval GmbH & Co. KG (Berlin, Germany). The mass fractions of alloying elements of the gas-atomized powder particles was analyzed by X-ray fluorescence spectroscopy (XRF, Bruker, Billerica, MA, USA). The composition is given in Table 1, which is close to the nominal composition.

Table 1. Mass fractions of alloying elements in the gas-atomized Al-20Si powder particles.

| Alloy | Si  | Fe  | Mg  | Ag  | Ti  | Cu  | Al   |
|------|-----|-----|-----|-----|-----|-----|------|
| Al-20Si | 20% | 0.27% | 0.07% | 0.03% | 0.02% | 0.02% | Bal. |

DFSC based on the thin film sensor (UFH1, Mettler Toledo, Greifensee, Switzerland) was used to characterize the melting and solidification temperatures in-situ. As indicated in Figure 1, film-thermopiles and resistive film-heaters are located at the center of the sensor membrane (amorphous silicon-nitride membrane). One particle (Particle I) of near-spherical shape with a diameter of 25 ± 0.5 µm was chosen and deposited in the center of the measurement area (with a diameter of 90 µm) by a thin copper wire with the help of a stereomicroscope (SZX16, Olympus, Tokyo, Japan) for DFSC measurements, as shown in Figure 1. Before and after the measurements, the particle size was characterized by optical microscopy (Olympus BX41). In order to improve the heat contact between the sensor membrane and the sample, a tiny amount of silicone oil was used. Due to the high heating rate (5000 K/s) and consequent short times (less than 0.12 s) at high temperatures, it was expected that the silicone oil was stable up to high temperatures, even though the boiling temperature of silicone oil is about 250 °C at low heating rates. After the first melting/solidification cycle, a slight flattening of the particle occurred. Thus, improved thermal contact between the sensor membrane and the sample was established. Because of a static temperature offset and a dynamic temperature offset due to thermal lag, the temperature calibration and correction were performed by the eutectic temperature (577 °C) of the Al-Si alloy upon heating at different heating rates before the DFSC measurements. Details of temperature calibration and correction can be found in [23]. For the investigation of the rapid solidification of this particle (Particle I) by DFSC, the particle was heated from 37 °C to 777 °C at a rate of 5000 K/s and held for 0.01 s. This maximum heating temperature ensures a certain overheating at the high heating rate, which is ca. 93 °C higher than the liquidus temperature of Al-20Si (ca. 684 °C [24]). Afterward, the particle was quenched to 37 °C at the same cooling rate. All the DFSC measurements were performed under argon (100 kPa constant pressure).
After rapid solidification by DFSC at 5000 K/s cooling, the particle (Particle I) was transferred from the sensor membrane into an epoxy block. After careful grinding and polishing of the epoxy block, a cross-section of the particle was obtained. After that, the specimen was etched with Weck’s reagent for 4 s [25]. Details of the specimen preparation of one single micro-sized metallic particle for microstructure characterization can be found in [26]. Scanning electron microscopy (SEM) (Zeiss Merlin VP Compact, Oberkochen, Germany) was used to characterize the microstructure (cross-section) of the particle.

To fully characterize the solidification behavior of the particle undergoing rapid solidification at 5000 K/s, another particle (Particle II) with the same size (25 ± 0.5 μm in diameter) as the particle for microstructure characterization (Particle I) was heated and cooled with the same time–temperature profile for 300 times. Note that the temperature calibration and correction were performed before these repeated heating–cooling measurements. Then, the nucleation undercooling, ΔT, for each scan was calculated by using the following equation:

\[
\Delta T = T_l - T_{s,\text{onset}}^{\text{correct}}
\]  

(1)

where \(T_l\) is the liquidus temperature and \(T_{s,\text{onset}}^{\text{correct}}\) is the solidification onset temperature (nucleation temperature). The evaluation of \(T_{s,\text{onset}}^{\text{correct}}\) can be found in [23].

3. Results and Discussion

The microstructure of the cross-section of a single Al-20Si particle (Particle I) rapidly solidified at 5000 K/s cooling by DFSC was characterized by SEM, as shown in Figure 2. The grey and bright areas in Figure 2 are Al and Si, respectively. The microstructure is comprised of the primary Si, secondary α-Al dendrites, and (α-Al + Si) eutectics. It seems very likely that the primary Si with polygonal morphology solidified from the surface of the particle (see Figure 2). Unlike the microstructure of gas-atomized particles, only two Si grains were observed because of the lower cooling rate than that of gas-atomized particles.

Figure 3 shows the corresponding DFSC curve at the heating–cooling rate of 5000 K/s for Particle I shown in Figure 2. The primary Si phase starts to solidify first during rapid solidification of an Al-20Si alloy, followed by the formation of the α-Al phase, and then the (α-Al + Si) eutectic. Two solidification onsets upon cooling (as indicated by the arrows in Figure 3) correspond to the starts of the nucleation of the primary Si and the secondary α-Al. The corresponding nucleation temperatures of the Si and α-Al phases for Particle I cooled at 5000 K/s are 627 °C and 521 °C, respectively. Note that the nucleation temperature of the α-Al + Si eutectics cannot be evaluated precisely because of the overlapping solidification peaks (at ca. 500 °C) between α-Al dendrites and α-Al + Si eutectics, as shown in Figure 3. For the nucleation of the primary Si, the nucleation undercooling can be evaluated by Equation (1) with the liquidus temperature of the Al-20Si alloy (684 °C), which is 57 K.
After the nucleation of primary Si, primary Si grows and rejects Al into the melt with decreasing temperature. At 521 °C (lower than the eutectic temperature 577 °C [24]), Al nucleates on the interface of primary Si and starts to grow in the melt. The melt is depleted of Si because of the solidification of primary Si. Assuming that the concentration of Si in the melt in front of the solidifying Si follows the metastable extension of the Al liquidus line in the Al-Si phase diagram [24] during the rapid solidification of primary Si, the concentration of Si is 9% (mass%) at 521 °C. The liquidus temperature at this concentration is 597 °C [24], and thus, the nucleation undercooling of α-Al is 76 K.

![Image](image-url)

**Figure 2.** SEM images (secondary electron images) of the cross-section of an Al-20Si particle (Particle I) solidified at 5000 K/s, etched with Weck’s reagent.

![Image](image-url)

**Figure 3.** DFSC heat flow curve of a single Al-20Si particle (Particle I) at the cooling rate of 5000 K/s. The melting and solidification onset temperatures of different phases are indicated by arrows.

For a sound statistical basis, another particle (Particle II) with the same size (25 ± 0.5 μm in diameter) as Particle I was heated and cooled at 5000 K/s for 300 times. Figure 4 shows the nucleation temperatures and undercoolings of the Si and α-Al phases for each of the 300 nucleations in chronological order. Both nucleations occur in bands of approximately 30 K width because of the stochastic nature of nucleation. The nucleation temperatures of the primary Si and α-Al for Particle I are also in these bands.
For a sound statistical basis, another particle (Particle II) with the same size (25 ± 0.5 μm in diameter) as Particle I was heated and cooled at 5000 K/s for 300 times. Figure 4 shows the nucleation temperatures and undercoolings of the Si and α-Al phases for each event in chronological order. Both nucleations occur in bands of approximately 30 K width because of the stochastic nature of nucleation. The nucleation temperature intervals, α, μ, and β, are used as their solidification enthalpy at ca. 627 °C. Thus, the mass of the solidified Si, \( m_{\text{Si}} \), is calculated by the following expression:

\[
\Delta H_{s,\text{DFSC}} = \frac{\Delta H_{m,\text{Si}}}{\Delta H_{s,\text{DFSC}}}
\]

where \( \Delta H_{s,\text{DFSC}} \) is the solidification enthalpy measured by DFSC (2.4 ± 0.2 μJ for the DFSC curve shown in Figure 3) and \( \Delta H_{m,\text{Si}} \) is the melting enthalpy of Si (1726 J/g) \[31\]. It should be pointed out that \( \Delta H_{s,\text{DFSC}} \) was evaluated by integrating the Si solidification shoulder.
(peak) until α-Al starts to solidify, as shown in Figure 3. It is assumed that the solidification of Si is limited by the nucleation events of α-Al. Thus, the mass of the solidified Si, \( m_{\text{Si,}_{\text{s}}} \), can be calculated and equals 1.4 ± 0.1 ng, corresponding to a volume of 600 ± 40 \( \mu \text{m}^3 \).

Then, assuming that the nuclei for α-Al form on the primary Si (interface nucleation on the primary Si), whose volume is 7581 ± 480 \( \mu \text{m}^3 \), the nucleation rate for α-Al is calculated by Equation (3), as shown in Figure 5b. As expected, both nucleation rates increase with increasing undercooling. It is supposed that the size of the primary Si and α-Al decrease with increasing undercooling.

![Figure 5](image)

**Figure 5.** Nucleation rates as a function of undercooling for the (a) primary Si and (b) α-Al dendrites. The dashed curves indicate the best fit to the data for the surface and interface heterogeneous nucleation models. Distributions of nucleation events versus undercooling are shown in the insets.

To understand the tendency between the nucleation rate and undercooling in terms of the spherical cap model of CNT, the expressions for the surface and interface heterogeneous nucleation rates of a single particle during rapid solidification are derived in [32,33] as:

\[
J_S = n_S D_l \frac{\sigma_{ls}}{a_0 k(T_{m,\text{onset}} - \Delta T)} \exp \left( - \frac{\Delta G^*_{S}}{k(T_{m,\text{onset}} - \Delta T)} \right)
\]  

(6)

and:

\[
J_V = n_V D_l \frac{\sigma_{ls}}{a_0 k(T_{m,\text{onset}} - \Delta T)} \exp \left( - \frac{\Delta G^*_{V}}{k(T_{m,\text{onset}} - \Delta T)} \right)
\]

(7)

with:

\[
\Delta G^* = \frac{16 \pi \sigma_{ls}^3}{3 \Delta G_{V}}
\]

(8)

where \( n_S \) and \( n_V \) are the densities of surface and interface heterogeneous nucleation sites (the numbers of potential surface and interface heterogeneous nucleation sites per unit surface area and unit volume), respectively, \( D_l \) is the liquid diffusivity, \( a_0 \) is the atomic spacing, \( \sigma_{ls} \) is the solid-liquid interfacial energy (surface tension), \( k \) is the Boltzmann constant, \( f_S \) and \( f_V \) are the shape factors for surface and interface heterogeneous nucleation, and \( \Delta G_{V} \) is the free energy difference between the melt and the solidified phase. The shape factors describe the potency of the heterogeneous nuclei.

During the DFSC measurements, an inherent Al\(_2\)O\(_3\) layer always exists, while the primary Si is solidified from the surface of the particle, as shown in Figure 2. Therefore, it is assumed that the surface nuclei for Si form in a spherical cavity at the surface of the
Materials 2021, 14, 2920

= 0

\[ f_{S,V} = \frac{1}{4\pi S_{S,V}^3} \left( -1 + 2\rho_{S,V} + \sqrt{1 + \rho_{S,V}^2 + 2\rho_{S,V} \cos \theta_{S,V}} \right) \left( 1 + \rho_{S,V} - \sqrt{1 + \rho_{S,V}^2 + 2\rho_{S,V} \cos \theta_{S,V}} \right)^2 \]  

(9)

with:

\[ \rho_{S,V} = \frac{r^*}{R_{S,V}} \]  

(10)

where subscript \( S,V \) designates the surface and interface heterogeneous nucleation, respectively, \( \theta_{S,V} \) is the contact angle between the heterogeneous nucleus and the undercooled melt, \( R_{S,V} \) is the radius of the spherical cavity, and \( r^* \) is the critical radius of a nucleus. Based on CNT, \( r^* \) is given by:

\[ r^* = \frac{2\sigma_s}{AG_V} \]  

(11)

Due to the lack of specific heat capacity data for each phase, especially in the undercooled region, \( AG_V \) is estimated by a parallel tangent construction [35]. Based on a thermodynamic assessment of the Al-Si system, the Gibbs energies of solid Al, solid Si, and liquid phases are represented by polynomial expansions [24]:

\[ G^i = 0G_{Al}^i(1 - x) + 0G_{Si}^i x + RT[x \ln x + (1 - x) \ln(1 - x)] + x(1 - x) \left[ A^i + B^i(1 - 2x) + C^i \left( 1 - 6x + 6x^2 \right) \right] \]  

(12)

where \( i \) designates the phase, \( x \) is the atomic fraction of Si (0.19 and 0.09 for the nucleation of Si and Al, respectively), \( R \) is the gas constant, \( T \) is absolute temperature, \( 0G_{Al}^i \) and \( 0G_{Si}^i \) are the lattice stability terms of the pure Al and Si, and \( A^i, B^i, \) and \( C^i \) are the interaction parameters that are linearly dependent on temperature. Table 2 lists the parameters for Equation (12).

| Table 2. Thermodynamic model parameters, J/mol, T in K [15,24]. |
|-----------------|-----------------|
| \( 0G_{Al}^{fcc} \) | \( -10,792 + 11.56T \) |
| \( 0G_{Si}^{fcc} \) | \( 12.12T \) |
| \( 0G_{Al}^{dia} \) | \( -50.600 + 30.00T \) |
| \( 0G_{Si}^{dia} \) | \( 30.00T \) |
| \( 0G_{Al}^{l} \) | \( 0 \) |
| \( 0G_{Si}^{l} \) | \( 0 \) |
| \( A^i \) | \( -10,695.4 - 1.823T \) |
| \( B^i \) | \( -4275.5 + 3.044T \) |
| \( C^i \) | \( 670.7 - 0.460T \) |
| \( A^{fcc} \) | \( -200 - 7.594T \) |
| \( A^{dia} \) | \( 89,138 - 31.445T \) |

The development of a complete description of the density of nucleation sites within an undercooled melt has been a continuing challenge. As mentioned above, the primary Si nuclei form on the surface of the particle. Therefore, it is assumed that the surface heterogeneous nucleation of Si could occur on the whole Al2O3 layer on the surface of the Al-20Si particle. Thus, the density of surface heterogeneous nucleation sites for primary Si can be expressed by [23]:

\[ n_S = \frac{N_A h}{V_m} \]  

(13)

where \( N_A \) is the Avogadro constant, \( h \) is the thickness of the Al2O3 layer (=1 nm) [36], and \( V_m \) is the molar volume of Al2O3 (=25.7 \times 10^{-6} g/mol) [36]. Thus, \( n_S \) equals \( 2.343 \times 10^{19} \) m^{-2}. For the interface nucleation of \( \alpha \)-Al, the density of interface heterogeneous nucleation sites in the melt is given by [15]:

\[ n_V = n_0^V \exp(c\Delta T) \]  

(14)
where \( n_0^0 \) is the base level of nucleation sites \((=10^{13} \text{ m}^{-3})\) and \( c \) is a positive constant \((=0.059)\) [15].

Based on this model (Equations (6)–(14)), the relationship between the nucleation rate and nucleation undercooling for the rapid solidification of single particles with a certain heterogeneity can be estimated. Employing the obtained parameters listed in Table 3, the data were fitted for Si and \( \alpha \)-Al phases. Here \( \theta_S, \theta_V, R_S, \) and \( R_V \) are fitting parameters. Levenberg-Marquardt nonlinear least-squares curve fitting was utilized with boundary conditions of \( 20^\circ < \theta_S, \theta_V \leq 180^\circ \) and \( 0.1 \text{ nm} < R_S, R_V < 10 \text{ nm} \). For the spherical cap model, the contact angle is larger than \( 20^\circ \) [37]. The fitted curves are indicated by dashed lines in Figure 5. The best fits to the data yield \( R_S = 1.29 \text{ nm} \) and \( \theta_S = 74.6^\circ \) for the surface nucleation of Si, and \( R_V = 2.61 \text{ nm} \) and \( \theta_V = 53.4^\circ \) for the interface nucleation of \( \alpha \)-Al. As shown in Figure 5, the modeling of the nucleation rates based on CNT with the inhomogeneous Poisson analysis of the thermal cycling behavior gives an accurate description of the 300 nucleation measurements for the single micro-sized Al-20Si particle that is rapidly solidified at 5000 K/s. Moreover, the nucleation mechanisms for the primary Si and secondary \( \alpha \)-Al are well described as surface and interface heterogeneous nucleation in a spherical cavity.

Table 3. Parameters used for nucleation rate analysis.

| Parameters                                      | Value                                      |
|------------------------------------------------|---------------------------------------------|
| Liquidus temperature (Si), \( T_{lSi} \)       | 684 °C [24]                                |
| Liquidus temperature (Al), \( T_{lAl} \)       | 597 °C [24]                                |
| Liquid diffusivity, \( D_l \)                  | \( 5 \times 10^{-9} \text{ m}^2/\text{s} \) [38] |
| Interfacial energy (Al in Al-Si system), \( \sigma_{slAl} \) | 0.169 J/m\(^2\) [39]                     |
| Interfacial energy (Si in Al-Si system), \( \sigma_{slSi} \) | 0.352 J/m\(^2\) [39]                     |
| Atomic spacing (Al), \( a_0^{Al} \)           | \( 2.6 \times 10^{-10} \text{ m (pure Al)} \) [27] |
| Atomic spacing (Si), \( a_0^{Si} \)           | \( 3.0 \times 10^{-10} \text{ m (Si atomic diameter)} \) [40] |

4. Conclusions

In-situ DFSC was successfully used to investigate the rapid solidification process of a single micro-sized Al-20Si (mass\%) particle at a cooling rate of 5000 K/s. By DFSC treatments at the same cooling rate, nucleation undercooling of one single micro-sized particle was correlated with the microstructure analysis of another similarly particle (with the same size). The results show that the microstructure of the rapidly solidified particle consists of primary Si, secondary \( \alpha \)-Al, and (\( \alpha \)-Al + Si) eutectics.

The nucleation rates of the primary Si and \( \alpha \)-Al dendrites were obtained by a statistical analysis of 300 solidification events (300 heating–cooling cycles of one single particle by DFSC). According to a modified classical heterogeneous nucleation theory, two different nucleation mechanisms of the primary Si and secondary \( \alpha \)-Al were proposed, i.e., surface heterogeneous nucleation for Si and interface heterogeneous nucleation for \( \alpha \)-Al.

By controlled rapid cooling and known nucleation undercooling via in-situ DFSC with microstructure characterization, it is possible to investigate the rapidly solidified structures and the nucleation mechanisms of single metallic particles. Therefore, this is a practical approach for a detailed investigation of the rapid solidification processes of metal particles and to discriminate surface and interface nucleation.

Author Contributions: Conceptualization, B.Y. and O.K.; methodology, Q.P. and B.Y.; software, B.Y.; validation, B.M., M.R., C.S. and O.K.; formal analysis, Q.P., B.Y. and D.L.; investigation, Q.P. and B.Y.; resources, B.M., A.S. and O.K.; data curation, Q.P. and B.Y.; writing—original draft preparation, B.Y.; writing—review and editing, Q.P., B.M., D.L., M.R. and C.S.; visualization, Q.P., B.Y. and A.S.; supervision, O.K.; project administration, B.Y., M.R. and O.K.; funding acquisition, B.Y., M.R. and O.K. All authors have read and agreed to the published version of the manuscript.
Funding: This research was founded by the German Research Foundation (DFG), grant numbers YA417/4-1 and KE616/29-1, and the German Research Foundation (DFG) priority program SPP2122, grant numbers RE1261/23-1 and KE616/27-1.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

Acknowledgments: B.Y. acknowledges support from the German Research Foundation (Project YA417/4-1). M.R. acknowledges support from the German Research Foundation (Project RE1261/23-1, SPP 2122). C.S. acknowledges financial support from the Ministry of Education and Science of the Russian Federation grant 14.Y26.31.0019. O.K. acknowledges support from the German Research Foundation (Project KE616/29-1 and KE616/27-1, SPP2122). The authors also thank Bilal Gökce from the University of Duisburg-Essen for XRF analysis.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Konrad, C.; Zhang, Y.; Xiao, B. Analysis of melting and resolidification in a two-component metal powder bed subjected to temporal Gaussian heat flux. *Int. J. Heat Mass Transf.* 2005, 48, 3932–3944. [CrossRef]
2. Konrad, C.; Zhang, Y.; Shi, Y. Melting and resolidification of a subcooled metal powder particle subjected to nanosecond laser heating. *Int. J. Heat Mass Transf.* 2007, 50, 2236–2245. [CrossRef]
3. Gusarov, A.V.; Yadroitsev, I.; Bertrand, P.; Smurov, I. Heat transfer modelling and stability analysis of selective laser melting. *Appl. Surf. Sci.* 2007, 254, 975–979. [CrossRef]
4. Gu, D.D.; Meiners, W.; Wissenbach, K.; Poprawe, R. Laser additive manufacturing of metallic components: Materials, processes and mechanisms. *Int. Mater. Res.* 2012, 57, 133–164. [CrossRef]
5. Li, Y.; Gu, D. Parametric analysis of thermal behavior during selective laser melting additive manufacturing of aluminum alloy powder. *Mater. Des.* 2014, 63, 856–867. [CrossRef]
6. Delahaye, J.; Tchuindjang, J.T.; Lecomte-Beckers, J.; Rigo, O.; Habraken, A.M.; Mertens, A. Influence of Si precipitates on fracture mechanisms of AlSi10Mg parts processed by Selective Laser Melting. *Acta Mater.* 2019, 175, 160–170. [CrossRef]
7. Kleiner, S.; Zürcher, J.; Bauer, O.; Margraf, P. Heat treatment response of selectively laser melted AlSi10Mg. *HTM* 2020, 75, 327–341. [CrossRef]
8. Prashanth, K.G.; Scudino, S.; Klauss, H.J.; Sreeddi, K.B.; Löber, L.; Wang, Z.; Chaubey, A.K.; Kühn, U.; Eckert, J. Microstructure and mechanical properties of Al–12Si produced by selective laser melting: Effect of heat treatment. *Mater. Sci. Eng. A* 2014, 590, 153–160. [CrossRef]
9. Aboulkhair, N.T.; Simonelli, M.; Parry, L.; Ashcroft, I.; Tuck, C.; Hague, R. 3D printing of Aluminium alloys: Additive Manufacturing of Aluminium alloys using selective laser melting. *Prog. Mater. Sci.* 2019, 106, 100578. [CrossRef]
10. Doubenskaia, M.A.; Zhirnov, I.V.; Telezheshkiy, V.I.; Bertrand, P.; Smurov, I.Y. Determination of true temperature in selective laser melting of metal powder using infrared camera. *Mater. Sci. Forum.* 2015, 834, 93–102. [CrossRef]
11. Li, X.P.; Wang, X.J.; Saunders, M.; Suvorova, A.; Zhang, L.C.; Liu, Y.J.; Fang, M.H.; Huang, Z.H.; Sercombe, T.B. A selective laser melting and solution heat treatment refined Al–12Si alloy with a controllable ultrafine eutectic microstructure and 25% tensile ductility. *Acta Mater.* 2015, 74, 71–82. [CrossRef]
12. Farshidianfar, M.H.; Khajepour, A.; Gerlich, A.P. Effect of real-time cooling rate on microstructure in Laser Additive Manufacturing. *J. Mater. Process Technol.* 2016, 231, 468–478. [CrossRef]
13. Hooper, P.A. Melt pool temperature and cooling rates in laser powder bed fusion. *Addit. Manuf.* 2018, 22, 548–559. [CrossRef]
14. Trivedi, R. The role of heterogeneous nucleation on microstructure evolution in peritectic systems. *Scr. Mater.* 2005, 53, 47–52. [CrossRef]
15. Gremaud, M.; Allen, D.R.; Rappaz, M.; Perepeko, J.H. The development of nucleation controlled microstructures during laser treatment of Al-Si alloys. *Acta Mater.* 1996, 44, 2669–2681. [CrossRef]
16. Wilde, G.; Sebright, J.L.; Perepeko, J.H. Bulk liquid undercooling and nucleation in gold. *Acta Mater.* 2006, 54, 4759–4769. [CrossRef]
17. Gandin, C.-A.; Mosbah, S.; Volkmann, T.; Herlach, D.M. Experimental and numerical modeling of equiaxed solidification in metallic alloys. *Acta Mater.* 2008, 56, 3023–3035. [CrossRef]
18. Trivedi, R.; Jin, F.; Anderson, I.E. Dynamical evolution of microstructure in finely atomized droplets of Al-Si alloys. *Acta Mater.* 2003, 51, 289–300. [CrossRef]
19. Das, S.K.; Perepeko, J.H.; Wu, R.I.; Wilde, G. Undercooling and glass formation in Al-based alloys. *Mater. Sci. Eng. A* 2001, 304–306, 159–165. [CrossRef]
20. Perepezko, J.H.; Sebright, J.L.; Höckel, P.G.; Wilde, G. Undercooling and solidification of atomized liquid droplets. *Mater. Sci. Eng. A* **2002**, *326*, 144–153. [CrossRef]

21. Perepezko, J.H.; LeBeau, S.E.; Mueller, B.A.; Hildeman, G.J. Rapid solidification of highly undercooled aluminum powders. In *Rapidly Solidified Powder Aluminum Alloys*; Fine, M.E., Starke, E.A., Eds.; ASTM International: West Conshohocken, PA, USA, 1986; pp. 118–136. ISBN 978-0-8031-4961-8.

22. Zhuravlev, E.; Schick, C. Fast scanning power compensated differential scanning nano-calorimeter: 1. The device. *Thermochim Acta* **2010**, *50*, 1–13. [CrossRef]

23. Yang, B.; Peng, Q.; Milkeret, B.; Springer, A.; Liu, D.; Rettenmayr, M.; Schick, C.; Keßler, O. Nucleation behaviour and microstructure of single Al-Si12 powder particles rapidly solidified in a fast scanning calorimeter. *J. Mater. Sci.* **2021**, *56*, 12881–12897. [CrossRef]

24. Murray, J.L.; McAlister, A.J. The Al-Si (Aluminum-Silicon) system. *Bull. Alloy. Phase Diagr.* **1984**, *5*, 74–84. [CrossRef]

25. Weck, E.; Leistner, E. *Metallographic Instructions for Colour Etching by Immersion, Part III: Non-Ferrous Metals, Cemented Carbides and Ferrous Metals, Nickel-Base and Cobalt-Base Alloys*; Deutscher Verlag für Schweiftechnik: Düsseldorf, Germany, 1986.

26. Milkeret, B.; Meißner, Y.; Ladewig, C.; Osten, J.; Yang, B.; Springer, A.; Keßler, O. Metallographische Präparation einzelner Pulver-Partikel. *Prakt. Metallogr.* **2021**, *58*, 129–139. [CrossRef]

27. Uttormark, M.J.; Zanter, J.W.; Perepezko, J.H. Repeated nucleation in an undercooled aluminum droplet. *J. Cryst. Growth* **1997**, *177*, 258–264. [CrossRef]

28. Yang, B.; Gao, Y.; Zou, C.; Zhai, Q.; Zhuravlev, E.; Schick, C. Repeated nucleation in an undercooled tin droplet by fast scanning calorimetry. *Mater. Lett.* **2009**, *63*, 2476–2478. [CrossRef]

29. Bokeloh, J.; Rozas, R.E.; Horbach, J.; Wilde, G. Nucleation barriers for the liquid-to-crystal transition in Ni: Experiment and simulation. *Phys. Rev. Lett.* **2011**, *107*, 145701. [CrossRef] [PubMed]

30. Simon, C.; Gao, J.; Ma, Y.; Wilde, G. Fast scanning calorimetric study of nucleation rates and nucleation transitions of Au-Sn alloys. *Scr. Mater.* **2017**, *139*, 13–16. [CrossRef]

31. NIST. *Chemistry WebBook: NIST Standard Reference Database Number 69*; Linstrom, P.J., Mallard, W.G., Eds.; National Institute of Standards and Technology: Gaithersburg, MD, USA, 2021.

32. Turnbull, D. Formation of crystal nuclei in liquid metals. *J. Appl. Phys.* **1950**, *21*, 1022–1028. [CrossRef]

33. Gutzow, I.; Schmelzer, J.W.P. *The Vitreous State: Thermodynamics, Structure, Rheology, and Crystallization*; Springer: Berlin, Germany, 1995.

34. Iwamatsu, M. Line-tension effects on heterogeneous nucleation on a spherical substrate and in a spherical cavity. *Langmuir* **2015**, *31*, 3861–3868. [CrossRef]

35. Thompson, C.V.; Spaepen, F. Homogeneous crystal nucleation in binary metallic melts. *Acta Metall.* **1983**, *31*, 2021–2027. [CrossRef]

36. Korb, L.J.; Olson, D.L. *ASM Handbook: Corrosion*; ASM International: Materials Park, OH, USA, 1992.

37. Cantor, B. Heterogeneous nucleation and adsorption. *Philos. Trans. R. Soc. A* **2003**, *361*, 409–417. [CrossRef]

38. Kalay, Y.E.; Chumbley, L.S.; Anderson, I.E.; Napolitano, R.E. Characterization of hypereutectic Al-Si powders solidified under far-from equilibrium conditions. *Metall. Mater. Trans. A* **2007**, *38*, 1452–1457. [CrossRef]

39. Gündüz, M.; Hunt, J.D. The measurement of solid-liquid surface energies in the Al-Cu, Al-Si and Pb-Sn systems. *Acta Metall.* **1985**, *33*, 1651–1672. [CrossRef]

40. Faraji, M. The Effect of Solidification Variables on the Microstructure of Hypereutectic Al-Si Alloys. Ph.D. Thesis, The University of Sheffield, Sheffield, UK, 2007.