Influence of mass fraction of SiC and undercooling on grain density function during heterogeneous nucleation primary phase in the AZ91/SiC composite

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INTRODUCTION

Grain size is one of the most important parameter which determined mechanical properties. Knowing element properties the proper application regions for it can be chosen to achieve best mechanical properties and performance. Nowadays simulation software can be use to predict the element microstructure. Those programs base on micro-macro model of crystallization. The model consists of partial differential equations (PDEs) that described the nucleation rate, diffusion in the casting, casting cooling speed and every single grain growth rate. Often it is hard to find the theoretical value of the parameters that appear in those PDEs. It is possible to find them from experiment. The experimental data that after applying statistical methods let us find approximated values of the so-called "fitting parameters" in the mentioned models [1 - 4].

AZ91 alloy analyzed in this study is hypereutectic alloy. The magnesium primary α-Mg phase is dendritic. During crystallisation there appears eutectic reaction. In this study influence of eutectic is omitted because magnesium primary phase has most significant influence on mechanical properties of the casting.

EXPERIMENTAL PROCEDURE

Composite casting

The AZ91 alloy was selected as the matrix for the composites. The chemical composition is shown in Table 1. The reinforcement particles are silicon carbide with an average diameter of 45 μm. Composite specimen with 0, 1, 2, 3 and 4 wt.% of SiC particles were prepared using a liquid mixing and casting process.

Table 1. **Chemical composition of AZ91 alloy**

Chemical composition, wt. %							
Al	Zn	Mn	Fe	Be	Si	Cu	Ni
9.03	0.6	0.2	0.0026	0.0011	0.0023	0.0016	0.00062

Processing of the magnesium composites consisted of mixing pre-heated SiC particles to 450 ^{o}C with liquid magnesium melt stirring and mould casting. About 1.4 kg of composite melts was prepared in an electric resistance furnace using a steel crucible under a SF_6/CO_2 gas atmosphere. The molten AZ91 alloy was held at 700 ^{o}C for 1 h. After putting SiC particles composite was stirred for 2 min, and then cast at 700 ^{o}C into mould to produce four plates of 100 x 100 x 10 (plate no 1), 15 (plate no 2), 20 (plate no 3) and 30 mm (plate no 4). The mould was made with resin sand hardened with CO_2. An un-reinforced AZ91 alloy was also cast at the same temperature (700 ^{o}C).

Thermal analysis

For the thermal analysis of AZ91 alloy and composite samples, cooling curves during solidification were obtained using a data acquisition system (Agilent) at a sampling rate of 5 data per second. A chromel-alumel (K-type) thermocouple positioned 50 mm from the bottom of the plate center, was used to monitoring temperature as the melt solidified

Microstructural analysis and grain size determination

The as-cast plates were sectioned at a distance 3 mm from hot junction of a thermocouple and next polished and etched before microstructural analysis. In order to visualization of grain boundaries of magnesium primary phase, the metallographic specimens were etched for 80-95 s. The chemical composition of solution was: 50 ml Distilled Water, 150 ml Ethanol, 1 ml Acetic Acid [5-7].

The etched specimens were examined using a light optical microscope Carl Zeiss AXIO Imager.A1 with cross polarized light and λ filter. The grains density was counted on the surface of etched specimens using image analysis NIS-Elements 3.0 Software. The images on computer display reveal arms of different dendrite grains as areas with different colours, Fig. 1.

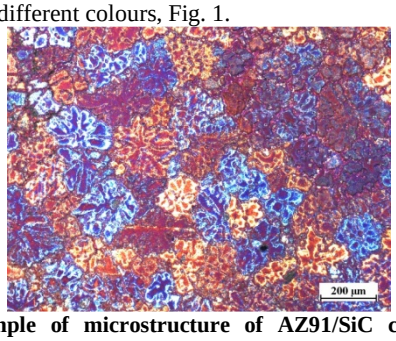

Fig. 1. Example of microstructure of AZ91/SiC composite for sample was cut from as-cast plate about thickness 10 mm with 2 wt.% of SiC

GRAIN DENSITY AND NUCLEATION MODEL

Data obtained from metallographic analysis can be used to calculate the grain's density per unit volume, N_V. To find this value Saltykov equation can be used [8]:

$$N_V = \frac{2}{\pi} \cdot N_a \cdot \left(\frac{1}{d}\right)_{mean}, \text{ m}^{-3} \quad (1)$$

where: N_V is mean the grain's density per unit volume, N_a is mean surface grain density, and $\left(\frac{1}{d}\right)_{mean}$ denotes average value of $\left(\frac{1}{d}\right)$ for all grains found on the polished section.

In this article the continuous nucleation model is taking into account. It is based on log-normal model described by Fras et. al in [4]:

$$N_V = \lambda \cdot \exp\left(-\frac{b}{\Delta T_{\max}}\right), \text{ m}^{-3} \quad (2)$$

where: λ [m^{-3}], b [K] – are model adjustment parameters, that should be find experimentally and ΔT_{max} denotes maximal undercooling.

It was shown by various authors that character of continuous nucleation [9, 10] is similar to model presented by Fras. According to this fact, as nucleation and crystallization simulation is performed, calculation of N_V step by step changes is connected with time by actual undercooling, denoted ΔT. During this calculation eq. (2) is used but for actual ΔT in the place of maximal undercooling ΔT_{max}.

Partial differential Fourier – Kirchhoff equation solved in parallel gives the actual $\Delta T(\tau)=T_N-T(\tau)$, where τ is time. The different maximal undercoolings, measured during the castings and connected volumetric grain densities gives us test values to calculate fitting parameters in equation (2). More complex model can be obtained if one use experimental data for different mass fraction of SiC particles, denoted mf_{SiC}. The test values can be than used to find

functions that describes λ and b parameters dependence on the mass fraction of SiC particles. The model that takes into account those functions can be expressed with formula:

$$N_V(\Delta T, mf_{SiC}) = \lambda(mf_{SiC}) \cdot \exp\left(-\frac{b(mf_{SiC})}{\Delta T}\right), \text{ m}^{-3} \quad (3)$$

The another parameter that is necessary for modeling of composite nucleation and crystallization is nucleation temperature, T_N. This value for composites depends on the mass fraction of SiC particles [2]. It also can be expressed with proper formula that can be finding statistically. Nucleation temperature can be obtained from thermoanalysis data, precisely from first derivative of cooling curve analysis, according to the procedure described by Kurz and Fisher [11].

RESULTS

Grain density of magnesium primary phase

The equilibrium phase for AZ91/SiC composite is the solid α-Mg solution, but during solidification a nonequilibrium eutectic (α-Mg - β-$Mg_{17}Al_{12}$) is also created and present in the un-reinforced AZ91 alloy and in the AZ91/SiC composite.

From thermoanalysis cooling curves the nucleation temperature, T_N, and maximal undercooling of primary phase, ΔT_{max}, can be calculated as a difference between nucleation and recalescence temperature. There are obtain from cooling curve and its first derivative. The nucleation temperature grows with increasing mass fraction of the SiC particles. The exponential dependence can be observed, it can be described with fallowing formula:

$$T_N(mf_{SiC}) = 606 - 5.8\exp(-90.4 mf_{SiC}), \text{ °C} \quad (4)$$

where: mf_{SiC} denotes dimensionless mass fraction of SiC particles in the composite.

The above formula was calculated, correlation coefficient for this fitting was R^2=0.991. The graph of statistically evaluated curve (4) is shown in Fig. 2.

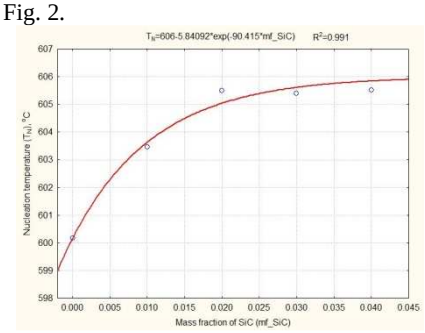

Fig. 2. The nucleation temperature dependence on mass fraction of SiC particles

Average grain diameter measurement data can be used also to calculate average volumetric grain density. This parameter is widely used to describe casting refinement of structure. It also is very useful during simulation, because it carries information about number of nuclei appearing in the unit of volume. The mean grain's density per unit volume, N_V, was calculated from Saltykov equation (1).

The grain's density N_V and corresponding maximal undercooling ΔT_{max} were used as the test values for approximate the adjustment parameters in Fras equation (2). For mass fraction of SiC particles the calculated equations have the following form:

- for 0 wt.% SiC, R^2=0.999, Fig. 3:

$$N_V = 1184 \cdot 10^9 \exp\left(-\frac{27.354}{\Delta T_{max}}\right), \text{ m}^{-3} \quad (5)$$

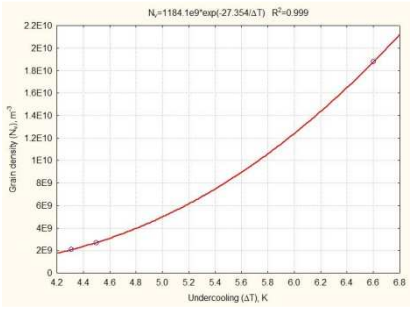

Fig. 3. The grain density dependence on undercooling for AZ91/0 wt.% SiC composite

- for 1 wt.% SiC, R^2=0.989, Fig. 4:

$$N_V = 5778 \cdot 10^9 \exp\left(-\frac{29.95}{\Delta T_{max}}\right), \text{ m}^{-3} \quad (6)$$

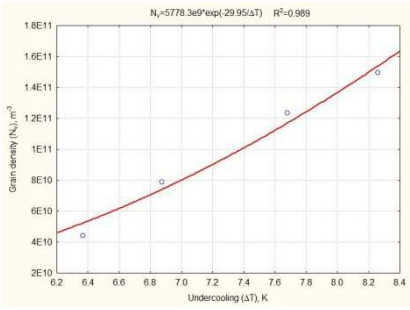

Fig. 4. The grain density dependence on undercooling for AZ91/1 wt.% SiC composite

- for 2 wt.% SiC, R^2=0.993, Fig. 5:

$$N_V = 5815 \cdot 10^{10} \exp\left(-\frac{85.9}{\Delta T_{max}}\right), \text{ m}^{-3} \quad (7)$$

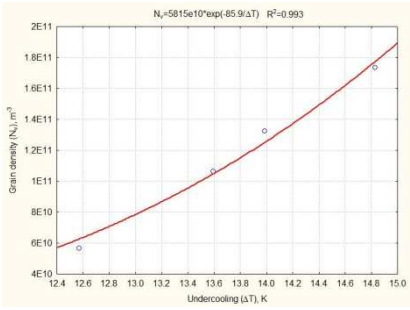

Fig. 5. The grain density dependence on undercooling for AZ91/2 wt.% SiC composite

- for 3 wt.% SiC, R^2=0.997, Fig. 6:

$$N_V = 1071 \cdot 10^{11} \exp\left(-\frac{102.26}{\Delta T_{max}}\right), \text{ m}^{-3} \quad (8)$$

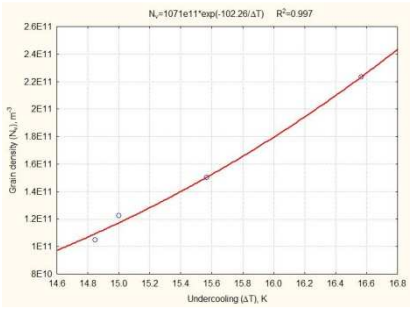

Fig. 6. The grain density dependence on undercooling for AZ91/3 wt.% SiC composite

- for 4 wt.% SiC, R^2=0.996, Fig. 7:

$$N_V = 1619 \cdot 10^{11} \exp\left(-\frac{104.85}{\Delta T_{max}}\right), \text{ m}^{-3}. \quad (9)$$

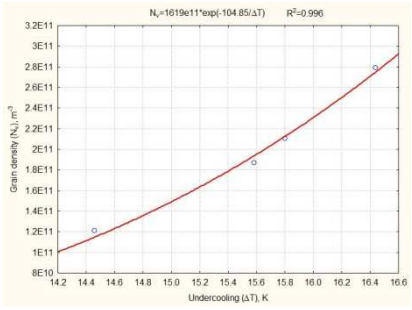

Fig. 7. The grain density dependence on undercooling for AZ91/4 wt.% SiC composite

The presented formulas can be used to describe the grain density function for specific mass fraction of SiC particles. Moreover if the maximal undercooling ΔT_{max} is replaced with actual under-cooling ΔT the presented equations can be use for continuous nucleation problem.

Further analysis of the experimental data leads to more general conclusion, that is, model for continuous nucleation that takes into account grain size. It can be described with the following expression:

$$N_V(\Delta T, mf_{SiC}) = 1.42 \cdot 10^{13} \exp\left(61.9 \cdot mf_{SiC} - \frac{36.25 \cdot \exp(29.3 \cdot mf_{SiC})}{\Delta T}\right), \text{ m}^{-3} \quad (10)$$

The correlation coefficient for this equation is $R^2 = 0.866$. Equation (10) can be used during nucleation simulation of the AZ91/SiC composites with different mass fraction of reinforcement particles.

The formula presented above gives possibility to calculate grain density continuously while melt temperature decreasing.

The nucleation model linked with the FK solving numerical scheme can be used to obtain very good approximation of the composite cooling speed, forming of the solid stage rate and to predict its microstructure.

Because of the lack of theoretical data, results of numerical analysis of composite nucleation phenomena (equations (5) – (10) presented above) can be very useful. Those equations linked with FK equation can give a lot of important information about AZ91/SiC composite crystallization phenomena.

CONCLUSIONS

The AZ91/SiC nucleation parameters as T_N and N_V can be described with mathematical formulas. Unknown adjustment parameters can be found using experimental data and statistical algorithms.

The mean volumetric grain density function shows grain density dependence on composite actual undercooling and mass fraction of SiC particles. This knowledge can be very useful for technologists during composite casting procedure preparation.

After setting the mass fraction of SiC particles and derivation the average volumetric grain density function gives information about nucleation rate. This is the key parameter for AZ91/SiC composite micro – macro model of crystallization.

ACKNOWLEDGEMENTS

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