Effect of MgO Content on Mechanical Properties of Directionally Solidified Pure Magnesium

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The pure magnesium was fabricated by directional solidification, and the effect of the distribution characteristics of magnesium oxide (MgO) on the mechanical properties of pure magnesium was investigated. The statistical results showed that the area fraction, number and size of MgO were decreased gradually from the top region ingot to the bottom region ingot, and these reflected the advantages of directional solidification technology in the controllability of MgO distribution characteristics. The top parts of magnesium ingots have the highest tensile strength (44 MPa), which is mainly due to the presence of a large amount of the coarse MgO. Though the coarse MgO increases the strength obviously, it has harm for the ductility of magnesium. The top parts of magnesium ingots have higher ultimate tensile strength, but lower failure strain (13% and -21% respectively) than the ingot at the center. These results indicate that if the suitable size and amount of MgO existed in magnesium matrix, it could avoid the disadvantages of MgO and provide positive effect for both the strength and ductility of magnesium alloys.

Keywords: Magnesium oxide, Directional solidification, Tensile properties, Diffusion mechanism

1. Introduction

Magnesium (Mg) is the lightest structural metal with rich resources on Earth, and it is the promising engineering material to improve energy efficiency and system performance in the automotive and electronic industry\textsuperscript{1,4}. Magnesium alloys consumption has increased to the third most used structural metal only behind iron and aluminum-based alloys\textsuperscript{7}. However, since Mg has low strength and poor plasticity, its applications are greatly limited. In the present, how to improve the ductility and strength of Mg at room temperature plays a key role in extending the application of Mg alloy\textsuperscript{4}. Nowadays, the various methods are used to strengthen Mg alloys, such as alloying, precipitation strengthening and fine-grained strengthening\textsuperscript{6-11}. Among them, the precipitation strengthening is one of the most effective methods to improve the mechanical properties of Mg alloys, because most of the intermetallic compounds (such as MgZn\textsubscript{12}, MgSn and Mg\textsubscript{13}(ZnY\textsubscript{11})) can enhance the mechanical properties of Mg alloys by pinning up dislocations and hindering the basal slip\textsuperscript{13,14}. As an active metal, Mg could easily form the eutectic compounds (such as Mg\textsubscript{17}Al\textsubscript{12} and Mg\textsubscript{2}Si\textsuperscript{19}) with adding elements. Unfortunately, these compounds always distribute in the grain boundary with net-shape morphology, mismatch with the Mg matrix, and it was harmful to improving the mechanical strength of Mg alloys\textsuperscript{17,19}. Conversely, magnesium oxides are perfect match with Mg matrix, and it also has uniform distribution characteristics and good thermal stability (melting point 2800 °C)\textsuperscript{20}, especially MgO is the most common phase in pure Mg and Mg alloys\textsuperscript{7,21,22}. However, the effect of MgO on the mechanical properties of Mg and Mg alloys are less reported, it can be attributed to the following reasons (i) it’s difficult to control the distribution of MgO (gradient distribution), and the researchers are more interested in grain boundary, grain size and grain orientation\textsuperscript{23,24}, and so on; (ii) many researches have been focused on the disadvantage of MgO on the surface properties of the Mg alloys\textsuperscript{25}. Additionally, it is generally accepted that the oxide (MgO) on the surface has a Pilling-Bedworth ratio < 1 leading to the porous oxide film of magnesium, which is not as protective as the passive films on Al\textsuperscript{21}. Furthermore, MgO could easily react with H\textsubscript{2}O in atmospheric conditions forming magnesium hydroxide (Mg(OH)\textsubscript{2}), and it is soluble in environments below pH=11, and many studies focus on the influence of MgO on the corrosion properties of magnesium alloys\textsuperscript{4,26,27}.

Directional solidification (DS) is a metallurgical technology that establishes a temperature gradient in a specific direction in the solidified area and the unsolidified area during the solidification of the alloy, which effectively changes the grain orientation and morphology in the solidification structure\textsuperscript{28,29}. Therefore, DS can be used to control the distribution of MgO. And there is no relevant research about the MgO structure and how it can affect the mechanical properties of Mg alloy. Nevertheless, the study of the distribution characteristic of MgO and its effect on the mechanical properties of magnesium alloy can help to understand the internal relationship between the distribution characteristic of MgO and the mechanical properties of Mg alloys. Most importantly, it’s also beneficial to develop the new technology about the recycling and reusing of Mg.
In order to investigate the effects of the distribution characteristic of MgO on the mechanical properties of magnesium alloys, this work used the direction solidification method to control the distribution characteristic of MgO in pure magnesium, and established a relationship between the distribution of MgO and the mechanical properties of Mg. Among them, pure magnesium is selected as the base alloy, which can greatly avoid the influence of other intermetallic compounds, and the directionally solidified magnesium ingot is coarse columnar grains that grow along the direction of heat flow, so the grain size and grain orientation of different regions are consistent, thus ensuring that magnesium oxide is the most important influencing factor.

2. Experimental Procedure

2.1. Preparation of casting specimens

Samples used in this work were prepared from commercially pure magnesium (Suzhou Haichuan Rare Metal Products Co. Ltd., China) by the directional solidification. The chemical compositions of the raw Mg were listed in Table 1. The pure Mg melted in a self-manufactured furnace, which structure was shown in Figure 1, and it was also described in our previous works. The directional solidification process was performed in an inert argon atmosphere to avoid the oxidation of the pure Mg. First, the samples were put into stainless steel crucibles with a 28/30 mm diameter (inside/outside diameter) and a length of 165 mm. Secondly, the specimens were heated to 1003 K over 1h and thermally stabilized for 2 h. When the magnesium is completely melted, turned off the bottom set of resistive wires labeled ‘②’ in Figure 1, cooled the bottom of crucibles by a water-cooled copper base, and then controlled the top set of resistive wires labeled ‘①’ in Figure 1, to keep the samples cool with 10 °C/min. At the end of the experiment, when the temperature went down to 673 K, turned off all the power and remained the sample in the furnace until the temperature went down to room temperature. As shown in Figure 1, the length of the sample was about 130 mm and the grain size was about 10mm.

2.2. Microstructural characterization

The casting samples were mechanically polished and etched with a 4% nital, an ethanol solution of picric acid and glacial acetic acid (2.0 g picric acid, 5 ml glacial acetic acid, 5 ml water and 25 ml ethanol). Microstructures of the alloys were observed by optical microscope (OM), scanning electron microscope (SEM) and using a JEOL JSM-6400 microscope equipped with Oxford Link Energy Dispersive X-ray (EDS) microanalysis spectrometer. A microstructural area of 4 mm$^2$ (a quarter of the cross-section) was grabbed from OM pictures at 200×. The quantitative distribution characteristic of MgO was statistics by image processing. The details of the quantification procedure of phase were documented elsewhere.

2.3. Calculation details and mechanical property test

Our calculations were performed using the Cambridge Serial Total Energy Package Code (CASTEP) based on density functional theory (DFT). The Generalized Gradient Approximation (GGA) was employed to evaluate exchange-correlation energy. Ultra-soft pseudo-potentials were used for electron-ion interactions and the electron wave function was expanded using plane waves. 2p$^6$3s$^2$ of Mg and 2s$^2$2p$^4$ of O are treated as valence electrons were involved in the calculations. The cut-off energy is 380 eV and the K-point of MgO is 12 × 12 × 12 and that of Mg is 18× 18 × 12 to

![Figure 1](image.png)

**Figure 1.** Schematic of the experimental set-up and the position and dimension of tensile specimens: ①- the top set of resistive wires, ②- the bottom set of resistive wires, ③- magnesium melt, ④- water-cooled copper base, ⑤- cooling water, ⑥- stainless steel crucible

| Alloy  | Al  | Mn  | Cu  | Fe   | Ni   | Si   | Mg   |
|-------|-----|-----|-----|------|------|------|------|
| Pure Mg | 0.025 | 0.016 | 0.004 | 0.0032 | 0.0005 | 0.017 | Remain |
ensure the convergence of the system energy and configuration at the plane wave group level. The Vickers hardness (HV) of the samples was measured with a micro-hardness tester (AHVD-1000XY, Jvjing) at room temperature using a load of 100 g for 15 s. Each sample continuously selected 10 hardness test points along the horizontal direction of the sample, the distance between each two test points is 0.6 mm, and the average hardness values were adopted as the results. In order to obtain the effect of position on mechanical properties of Mg, the tensile specimens of a cross section of 2 mm ×5 mm and 15 mm gauge length were machined from the bottom, center and top of the ingot, as shown in Figure 1. Uniaxial tensile test was performed on the bottom, center and top specimens at a constant strain rate of $10^{-4}/s$ in a computer controlling servo-hydraulic test machine at room temperature. At the end of the test, the fracture surface was observed by the SEM.

3. Results and Discussion

3.1. Distribution characters of MgO

Figure 2 showed the microstructures of the top, the center and the bottom regions of the pure magnesium ingot respectively, and it mainly consisted of well-development primary α-Mg and MgO. It should be noticed that the differences in the size, the amount and the morphology of MgO were observed among the three regions of the ingot. The size difference among MgO in the top region of ingot was bigger than the MgO in the other two regions, and the coarser MgO was mainly occurred at the top of ingot. As shown in Figure 2(d-f), there are not only MgO particles in the matrix, but also other impurity phases, and the element composition is shown in Table 2. In addition, a variety of elements were found in the impurity phase in the bottom region, among which the presence of Fe element had a passive impact on the mechanical properties of the magnesium matrix. In order to investigate the size difference and the distribution characteristic of MgO in casting, we counted the number density and the area fraction of MgO in the top, the center and the bottom region of casting, as shown in Figure 3. In the top region of ingot, when the size of MgO is less than 100 μm, the area fraction of MgO is maintained at a low value. But the area fraction of MgO increased sharply to 0.62% with the size of MgO larger than 100 μm. The area fraction is three times than other sizes of MgO. Additionally, the number density of MgO reduced sharply with the size of MgO increasing, high proportion of the number density of MgO (around 82%) belongs to 0-25 μm. This tendency was observed in both the center and bottom regions of ingot. In the center region of ingot, the area fraction of MgO displayed the same tendency as the top region, and the area fraction of MgO in the center region of ingot was larger than the top ingot (its size was in 0-25 μm, and 25-50 μm). The area fraction of MgO with the size larger than 100 μm was 0.58%, and it’s less than the area fraction of MgO in the top ingot. These data implied that the total area fraction of MgO and amount of coarser MgO reduced from top to center of ingot gradually. In the bottom of ingot, the area fraction of MgO

| Point | Mg    | O     | Si    | Al    | Fe    |
|-------|-------|-------|-------|-------|-------|
| Point 1 | 95.15 | 4.85  | —     | —     | —     |
| Point 2 | 64.99 | 3.55  | 31.46 | —     | —     |
| Point 3 | 72.55 | 27.45 | —     | —     | —     |
| Point 4 | 57.35 | 3.48  | 33.78 | 4.07  | 0.94  |

Figure 2. The morphology of the different regions of the casting: (a) (d) top; (b) (e) center; (c) (f) bottom
The area fraction of MgO from the top to the bottom ingot was 1.51%, 1.45% and 1.01%, respectively. These data indicated that the area fraction, number and size of MgO reduced from the top to the bottom of ingot. The coarser MgO mainly appeared in the top region of ingot and the bottom region’s impurity phase size was relatively small and uniform.

The reason for the distribution of MgO was considered to be related to the diffusion behavior of oxygen in the magnesium melt. During the melting process, the oxygen diffused into the magnesium melt surface through the protective gas (Argon) layer. The diffusion coefficient of oxygen in the protective gas layer could be obtained according to the Fuller-Schettler and Giddings correlation, as shown in Equation 1.

$$D = \frac{4.357 \times 10^{-2} T^{3/2}}{P \left( \frac{1}{V_A} + \frac{1}{V_B} \right)^{1/2}}$$

(1)

Where T is the temperature of the gas; $M_A$ and $M_B$ are the molar mass of A and B, $M_{O_2} = 0.032 \text{ kg/mol}$, $M_{Ar} = 0.04 \text{ kg/mol}$; P is the pressure of gas, $P = 1 \times 10^5 \text{ Pa}$; $V_A$ and $V_B$ represent the liquid molecular volume in the normal boiling point of gas A and B, $V_{O_2} = 2.56 \times 10^{-7} \text{ m}^3/\text{mol}$, $V_{Ar} = 2.86 \times 10^{-7} \text{ m}^3/\text{mol}$. When $T = 1003 \text{ K}$, we could solve $D = 620 \text{ m}^2/\text{s}$. According to the Fick’s first law, when D is constant, the diffusion speed is determined by the absorption speed of oxygen of magnesium melt. There were two forms of oxygen existing in magnesium melt, solid solution and compound. Because the magnesium was prone to react with oxygen, and MgO was the main phase of oxygen in the magnesium melt. The Gibbs free energy of the reaction between magnesium and oxygen could be determined by Equation 2.

$$\Delta G_{mol} = -RT \ln K = -RT \ln \frac{MgO}{[Mg][O]} = \Delta G_{mol} + \Delta G_f$$

(4)

$$\ln \left[ \frac{[O]}{[O]} \right] = 2.353 - 8856/T$$

(5)

Where [Mg] and [O] is the density of magnesium and oxygen atomic, the critical value of the oxygen density of reaction can be obtained from equation (5). When $K = 1003 \text{ K}$, [O] was equal to $3.34 \times 10^{-7}$. Since the critical value of the oxygen density was very low during the oxygen and magnesium reaction, the oxygen mainly existed as MgO and there were only very low levels of residual free oxygen existing in the magnesium melt. During the melting of the pure magnesium, the residuary air in the furnace diffused into the magnesium matrix and form a compact layer on the surface, as shown in Figure 4(a). However, as shown in Figure 4(b), with the holding time increasing, since the volume difference between compact layer and metallic matrix and the transport of Mg vapor, the internal stress would arise and lead the compact layer break\cite{20,21}. Many cracks occurred in the compact layer at this time. The cracks could provide easy paths for oxygen diffusion to the magnesium melt. And during the breaking process, many different sizes of MgO were escaped from the loose compact layer and appeared below the compact layer. In these MgO, the coarser MgO would float on the surface of magnesium melt, and the other fine dense MgO slowly moved toward the bottom of the crucible. And these fine MgO would grow up by swallowing other free MgO in the magnesium melt during the subsided processing. Therefore, the coarser MgO was mainly occurred at the top region of ingot. The area fraction, number and size of MgO in the top region of the ingot were larger than both the center and bottom ingot. This is why the MgO has a gradient distribution characteristic in the ingot.

### 3.2. Effect of MgO on mechanical properties

The mechanical property of MgO plays an important role in the reposed behaviors of magnesium and magnesium alloy during deformation process. Therefore, it was essential to investigate the elastic stiffness of MgO. The quotient between shear modulus (G) and bulk modulus (K) G/K can
be considered as an indication of the extent of fraction range in metals. A low value and a high value of G/K are associated with ductility and brittleness, respectively.

The Young’s modulus (E), Poisson’s ratio (ν) and anisotropy parameter (A_G) of Mg and MgO can be calculated and obtained by relevant formulas in the Wei39, as shown in Table 3. The G/K values were 0.442 and 0.775 for the Mg and MgO phase, respectively. As we know, the material was regarded as brittle if the value of G/K was more than 0.57. This suggested that MgO has brittleness. According to calculations, the bulk modulus of Mg (40.5 GPa) was much smaller than that of MgO (265.3 GPa), and MgO had larger shear modulus (111.4 GPa). These results implied that the shape of MgO was difficult to change or broken to small pieces during the deformation process and it had a strengthening effect on the magnesium matrix. Additionally, it was known that the elastic anisotropy could be measured using dimensionless quantity A_G. The results demonstrated that the Mg was more anisotropic than MgO, and it also agreed with the following calculation results of anisotropy. To investigate the elastic modulus anisotropy of MgO and Mg, a three-dimensional surface representation of elastic anisotropy was employed to show the variation of the elastic modulus with the crystallographic direction, and these results were shown in Figure 5. Since the shape of the three-dimensional surface would indicate the degree of anisotropy of phases, for example the spherical shape indicated the isotropic phase. From the three-dimensional surface images, it could be found that the Mg was much more anisotropic than MgO, since the shape of the three-dimensional surface image of MgO tends to spherical shape and the shape of Mg was fusiform shape.

Figure 6 showed the microhardness and stress-strain curves of the different regions of the casting. The mechanical properties were provided in Table 4. It can be seen from the microhardness of the three areas of the ingot that the microhardness of the top ingot is the largest (58.8±3.5 HV), which was related to the content and size of the MgO particles. However, the content of MgO particles in the bottom ingot was the least, and the microhardness was greater than that in the center ingot. This was mainly due to the large amount of impurity phases in the bottom ingot, and these impurity phases include MgO and mixtures (including Al, Fe and Si)40. Figure 6b was a typical strain strengthening curve. The stress increased gradually to the peak and then decreased drastically. It should be mentioned that in the center ingot, the elongation of sample (61%) was much higher than the bottom ingot (30%), although the area fraction and the size of MgO in bottom was smaller than that of the center. The other interesting finding was that the top region with a high area fraction of MgO showed high yield strength (9 MPa) and tensile strength (44 MPa). Combined with the above statistical results of MgO, it suggested that the magnesium matrix with finer MgO (<50 μm) had better plasticity than those with coarser MgO (>100 μm). But the coarser MgO increased strength much more significantly than those with finer MgO.

Figure 7 showed the fracture surfaces of tensile samples in the different regions of the casting alloy. All samples exhibited a fracture surface with cleavage facets and deep tearing ridges. These features showed evidence that the cross slipping and twinning played an important role.

### Table 3. Voigt (index V), Reuss (index R), and averaged macroscopic modulus for the Mg and MgO phase; all in Gpa except for ν (dimensionless)

| Phase | K_V | K_R | K | G_V | G_R | G | E    | ν     | G/K | A_G |
|-------|-----|-----|---|-----|-----|---|------|------|-----|-----|
| Mg    | 35.1| 35.1| 35.1| 16.3| 14.7| 15.5| 40.5 | 0.308| 0.442| 5.2%|
| MgO   | 143.7| 143.7| 143.7| 114.0| 108.7| 111.4| 265.3| 0.192| 0.775| 2.3%|

![Figure 4. Schematic of the stages for MgO growth on the surface of magnesium melt: (a) the initial state of magnesium melt; (b) distribution of different size of MgO in magnesium melt](image-url)
in the process of pure magnesium deformation at room temperature. In the top ingot, as shown in Figure 7a, the fracture surface contained substantial tearing ridges and a few cleavage facets. In comparison, as shown in Figure 7b, the amount of tearing ridges was much more than the top region, and the area of the cleavage facets on the fracture surface tend to decrease compared to the top region. As shown in Figure 7d, the small porosities were found on the tearing ridges. The formation of the porosity should be related to the finer MgO, which hindered the propagation of tearing ridges. And the tearing ridges were much shorter and more uniform than the top region. In contrast, as shown in Figure 7c, the bottom sample exhibited large cleavage facets, which was much flatter and broader than the other regions. This might be attributed to the brittle element, such as Fe, Si, and Al, which was segregated to the bottom of ingot during the directional solidification process. These impurity elements would change the interatomic force and alter the slipping mode.

From what had been discussed above, we might finally draw the conclusion that MgO had some positive effects on the mechanical behavior of magnesium alloys, and the finer MgO could enhance the strength of the magnesium with good plasticity as well. The MgO was not only smaller mismatch with the magnesium matrix, but also uniformly distributed in the whole grain, rather than segregation at the grain boundary which was often occurring in the other magnesium alloys. If we could avoid the harmful characteristics of MgO, it could become an important strengthening phase in magnesium and magnesium alloys.
4. Conclusions

In this work, the effect of microstructures (MgO particle distribution) and mechanical properties of the different areas of the directionally solidified pure magnesium ingot were studied. The main conclusions are as follows.

1. The distribution characteristics of MgO in the ingot were related to the diffusion behavior of oxygen closely. Many different sizes of MgO were formed because the residuary air in the furnace diffuses into the magnesium melt. The finer and denser MgO slowly moved toward the bottom of the crucible and the coarser MgO would float on the surface of magnesium melt. Therefore, the area fraction, amount and size of MgO in the top ingot were larger than the middle and bottom ingot.

2. The coarser MgO particles (>100μm) could enhance the microhardness and tensile strength of the magnesium matrix, and the finer MgO particles (<50μm) could improve its plasticity, comparing the mechanical properties of the magnesium ingot at the top and the center.

3. The poor mechanical properties of the bottom ingot are mainly due to the small size and amount of MgO in the bottom ingot, and the presence of mixed impurities (including Al, Fe and other elements).

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6. References

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Figure 7. SEM micrographs of fractured tensile samples in the different regions of the casting: (a) top; (b) center; (c) bottom; (d) zoom in of image (b).
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