Effect of CaO Addition on Microstructure and Damping Capacity of AM50 Magnesium Alloy

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The effects of a small amount of CaO addition (0.3 mass%) on the microstructure and damping properties of AM50 casting alloys have been investigated. The added CaO contributes to the formation of an Al$_2$Ca phase but reduces the total amount of compound particles by decreasing the $\beta$ phase content. The AM50–CaO alloy shows a slightly higher damping level within the strain-amplitude dependent region than does the alloy without CaO. The lower number density of compound particles acting as strong pinning points for dislocations is thought to be responsible for the improvement in the damping capacity.

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1. Introduction

Because of the inherent advantages such as low density, high specific strength, good castability and damping properties, Mg alloys have recently gained increasing attention as a versatile material with distinct impact on the society.\(^1\,\text{,}\,\text{2}\) Among the specific properties of Mg alloys, their low weight is an essential factor for consideration in designing transport vehicles and portable IT products: however, in view of safety, crashworthiness and comfort, the high damping characteristics of Mg alloys, which suppresses unwanted noise, irregular vibrations and resulting structural instabilities are also extremely important.\(^3\,\text{,}\,\text{4}\) Recently, it has been reported that an addition of CaO into pure Mg or Mg–Al based casting alloys enables the formation of dense films consisting of CaO and MgO on the molten surface without casting procedures involving SF$\text{\textsubscript{6}}$.\(^5\,\text{,}\,\text{6}\) Moreover, alloyed elemental Ca reduced from CaO results in improved mechanical properties at room and elevated temperatures by microstructural refinement, owing to CaO addition, is closely related to the alloying of elemental Ca reduced from CaO.\(^7\) There is little information available on the damping capacity of commercial Mg–Al based alloys with CaO. Accordingly, in the present study, we comparatively examine the damping capacities of AM50 casting alloy with and without the addition of CaO, and discuss the results on the basis of the microstructural evolution.

2. Experimental

Two AM50 based alloys having 0 and 0.3% (mass%) CaO, were prepared by remelting commercial billets supplied from the manufacturer and then casting them into a metallic mold. From the ingots, various specimens were prepared by machining for optical and electron microscopy, as well as for damping tests. The chemical compositions of the experimental alloys, as determined by an inductively coupled plasma (ICP), are listed in Table 1.

![Table 1 Chemical compositions of experimental alloys (mass%).](image)

<table>
<thead>
<tr>
<th>Composition</th>
<th>Al</th>
<th>Mn</th>
<th>Ca</th>
<th>Mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>AM50</td>
<td>5.09</td>
<td>0.31</td>
<td></td>
<td></td>
</tr>
<tr>
<td>AM50–0.3%CaO</td>
<td>4.94</td>
<td>0.21</td>
<td>0.28</td>
<td>bal.</td>
</tr>
</tbody>
</table>

The damping capacity was measured at room temperature using a dynamic mechanical analyzer (DMA, TA Q-800) in the strain amplitude range $1 \times 10^{-5}$ to $3 \times 10^{-3}$ vibrating in a single cantilever mode (frequency: 1 Hz). The dimension of the damping test specimen was $35 \times 12 \times 1 \text{mm}^3$. The damping capacity was evaluated in terms of the loss tangent (tan$\phi$), where $\phi$ is the phase lag angle between the applied strain and the responding stress. Microstructural evaluations were carried out by optical microscopy (OM, Nikon Epiphot-200BD), scanning electron microscopy (SEM, FEI QUANTA 200F) combined with energy dispersive X-ray spectroscopy (EDS, AMETEK PV72-60030F) and scanning transmission electron microscopy (STEM, JEOL JEM-ARM20F) with EDS (BRUKER QUANTAX 400).

3. Results and Discussion

3.1 Microstructures

Figure 1 shows the optical microstructures of AM50 and AM50–0.3%CaO alloys in as-cast state. The microstructures consist of primary $\alpha$(Mg) grains and intermetallic particles distributed within the grains and along the grain boundaries. It is seen that the CaO-containing alloy has a slightly finer-grained microstructure and a lower amount of intermetallic particles. The average grain size and the volume fraction of the intermetallic phase, measured by means of an image analysis, are $48 \mu\text{m}$ and $\sim$16% in the AM50 and $40 \mu\text{m}$ and $\sim$12% in the AM50–0.3%CaO alloys, respectively. This microstructural evolution, owing to CaO addition, is closely related to the alloying of elemental Ca reduced from CaO and is very similar to that of Mg–Al–Ca alloys.\(^10\,\text{–}\,\text{12}\) In a previous work, Amberger et al.\(^10\) reported that an addition of 1% Ca into AZ91D results in a decrease in the volume fraction of the $\beta$(Mg$_{17}$Al$_{12}$) phase by the formation of Al$_2$Ca (the total volume fraction of intermetallic particles may well be reduced because the Al$_2$Ca phase needs a higher amount

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of alloying elements than does the $\beta$ phase, which consumes Al and Mg simultaneously).

A TEM image and the corresponding EDS mapping on an intermetallic particle of the AM50–0.3%CaO alloy are presented in Fig. 2. Microanalysis performed on these particles indicates that the Al$_2$Ca phase is formed in the matrix. Figure 3 shows SEM images and EDS compositional analyses on the $\beta$(Mg$_{17}$Al$_{12}$) phase in the AM50 alloy with 0.3%CaO. It is noteworthy in Fig. 3 that lamellar and bulky types of the $\beta$ phase are seen in the microstructure and that Ca atoms dissolve in the $\beta$ phase. Inclusion of Ca in the $\beta$ phase was demonstrated theoretically by Min et al.\textsuperscript{13} They calculated the valence electron structure (VES) of the $\beta$ phase by using empirical electron theory. Their results suggested that the addition of Ca results in a more uniform distribution of valence electrons in the dominant bonds and within the whole unit cell, causing an increase in the melting point of the $\beta$ phase. Recent results by Kondori et al.\textsuperscript{14} confirmed that Ca atoms dissolve and distribute unevenly in the $\beta$ particles of the AM60–0.5%Ca alloy.

### 3.2 Damping capacity

Figure 4 shows the change in damping capacity ($\tan \phi$) with strain amplitude for the AM50 and AM50–0.3%CaO alloys. As seen in Fig. 4, the damping curves can be divided into two regions. At low strains, the loss tangent is only weakly dependent on strain amplitude, whereas a distinct dependency on strain amplitude evolves at higher strains. It is noticeable that both the CaO-containing and the CaO-free alloys exhibit a similar damping capacity at low strain amplitudes, but slightly higher damping levels are observed for the CaO-containing alloy at high strain amplitudes, even though the increment is not remarkable. This behavior is inconsistent with the results observed for Mg–Ca binary alloys, where the damping capacity decreases continuously upon increasing Ca content.\textsuperscript{15} After considering the previous reports that the addition of Mn up to 1% does not affect to the damping capacity in pure Mg\textsuperscript{16} and rather slightly improves the damping capacity in Mg–Ni alloys,\textsuperscript{17} the small difference in Mn content (0.1%) in the experimental alloys is not likely to contribute to the damping capacity. This confirms that a better damping capacity is due to the microstructural evolution in response to CaO addition.

It is well known that the total damping capacity ($\delta$) can be divided into two parts, namely, a strain-amplitude independent damping ($\delta_0$) and a strain-amplitude dependent damping ($\delta_{\text{II}}$). $\delta_0$ is related to the loss caused by forced vibrations of dislocation segments pinned by weak pinning points such as solute atoms, while $\delta_{\text{II}}$ results from the loss caused by dislocation segments pinned by strong pinning points such as precipitates and aggregates. According to the Granato and Lücke (G–L) theory,\textsuperscript{18,19} $\delta$ can be described as follows:

$$
\delta = \delta_0 + \delta_{\text{II}}(\varepsilon) \quad (1)
$$

$$
\delta_0 \sim \rho L_a^4 \quad (2)
$$

$$
\delta_{\text{II}}(\varepsilon) = \frac{C_1}{\varepsilon} \exp\left(-\frac{C_2}{\varepsilon}\right) \quad (3)
$$

Figure 1 Optical microstructures of (a) AM50 and (b) AM50–0.3%CaO alloys in as-cast state.

Fig. 2 TEM image showing an Al$_2$Ca particle (left) and EDS maps of the elements Mg, Al, Mn and Ca (right) in AM50–0.3%CaO alloy.

Fig. 3 (a) SEM image and (b) EDS analysis of the lamellar $\beta$ phase (A) and (c) SEM image and (d) EDS analysis of the bulky $\beta$ phase (B) in AM50–0.3%CaO alloy.
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4. Conclusion

The microstructures and damping properties of AM50 alloys with and without addition of CaO were comparatively investigated. The small amount (~0.3 mass%) of additional CaO causes the formation of the Al2Ca phase in the microstructure; however, the addition reduces the total amount of compound particles by decreasing the β phase content. The CaO addition leads to a better damping capacity at high strain amplitudes. The reason could be related to the lower number density of compound particles acting as strong pinning points for dislocations.

REFERENCES


\[
C_1 = \frac{\rho F_0 L_N}{6bEL_d} \quad (4)
\]

\[
C_2 = \frac{Kb \eta}{L_d} \quad (5)
\]

where \( \rho \) and \( L_d \) are the dislocation density and the average distance between weak pinning points, \( F_0, E, L_N, b, K \) and \( \eta \) are the binding force between dislocations and weak pinning points, the elastic modulus, the average distance between strong pinning points, the Burger’s vector, a constant and the size ratio of solvent and solute atoms, respectively. On the basis of the G–L theory, a slightly higher damping level in the strain-amplitude dependent region is plotted against \( 1/\varepsilon \) in Fig. 5 using the relation \( \delta = \pi \tan \phi \) (\( \tan \phi < 0.06 \)) relation. In this figure, \( \ln(C_1) \) and \( C_2 \) correspond to the Y-axis intercept and the slope, respectively. The obtained \( C_1 \) and \( C_2 \) values are \( 8.8 \times 10^{-4} \) and \( 3.96 \times 10^{-3} \) for the AM50 and \( 12.1 \times 10^{-4} \) and \( 4.05 \times 10^{-3} \) for the AM50–0.3%CaO alloys. The CaO-added alloy has a higher value of \( C_1 \), but an almost identical value of \( C_2 \), which supports well the experimentally observed enhanced damping capacity by CaO addition at high strain amplitudes.

![Fig. 4 Change in tan\( \phi \) with strain amplitude in AM50 and AM50–0.3%CaO alloys.](image)

![Fig. 5 G–L plots of AM50 and AM50–0.3%CaO alloys.](image)