Phase Transitions During Continuous Heating of Martensitic CuAlMn Alloys

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Abstract: Phase transitions in CuAlMn alloys containing up to 12 at% Mn and 27 at% Al continuously heated up to 550°C were studied using differential scanning calorimetry and transmission electron microscopy. Alloys containing less than 4.8 at% Mn exhibit an exothermal heat effect in the range 280 - 380°C due to γ2 precipitation. The additional endothermal peak above 500°C is caused by DO3 disordering. Alloys of intermediate Mn and Al content show only a reversible martensitic and ordering transformations during repeated cycling in the range -50 to +550°C. Alloys of higher Mn (6 - 12 at%) and lower Al content exhibit a diffused exothermal peak in the range 320 - 450°C due to bainitic transformation and a broad endothermal effect above 500°C due to L21/DO3 disordering. Bainitic plates possess 9R structure. They contain more Cu and less Mn and Al when compared to initial alloy compositions. Precipitates of γ2 compositions lie close to its equilibrium phase limits.

1. INTRODUCTION

CuAlMn alloys have been shown to exhibit a shape memory effect comparable to that of CuAlZn and CuAlNi alloys [1-3]. Martensitic transformation temperatures are very sensitive upon an ageing treatment causing initial their increase, then decrease [4-6]. This was explained as due to a change of matrix chemical composition [4, 5]; however, increase of stresses and changes of ordering, particularly in the first ageing period seem to be also important [6, 7]. Recent studies of the ordering transitions in CuAlMn alloys indicate a decrease of DO3 → B2 transition temperature with increasing manganese content [8]. Nakanishi et al [9] observed also an increase of the martensitic transformation temperatures accompanying a change of Mn site occupancy during ageing.

In alloys containing less than 5 wt% Mn γ2 precipitation was observed, while with increasing Mn content formation of bainite (α') occurred during isothermal ageing causing significant changes of the characteristic transformation temperatures and degradation of martensitic transformation, at final stages of ageing. Depending on the manganese and aluminium content the kinetics of precipitation is altered [4, 6, 7]. It is therefore of interest to know what is the effect of continuous heating on the precipitation and ordering changes in alloys with various Mn and Al content.
2. EXPERIMENTAL PROCEDURE
The alloys were cast in an induction furnace under argon atmosphere. After homogenization treatment at 750°C they were hot rolled down to 2 and 0.1 mm. The characteristic transformation temperatures were measured using a differential scanning calorimeter (DSC) Perkin-Elmer DSC-4 at a heating/cooling rate of 20K/min. Composition of alloys and Ms temperatures are given in Table 1. The alloys were quenched from 800°C into room temperature (RT) water except for alloys 1, 4 and 5 which were quenched in water at 60°C. Transmission analytical electron microscope Hitachi H-600 equipped with scanning attachment and Link detector was used for microstructure investigations. Thin foils were obtained by jet electropolishing in CrO₂ saturated H₃PO₄ solution.

<table>
<thead>
<tr>
<th>Alloy</th>
<th>Cu at%</th>
<th>Mn at%</th>
<th>Al at%</th>
<th>Ms[°C]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>71.4</td>
<td>1.5</td>
<td>27.1</td>
<td>18</td>
</tr>
<tr>
<td>2</td>
<td>70.0</td>
<td>2.9</td>
<td>27.1</td>
<td>-59</td>
</tr>
<tr>
<td>1A</td>
<td>68.5</td>
<td>4.8</td>
<td>26.7</td>
<td>-136</td>
</tr>
<tr>
<td>4</td>
<td>70.9</td>
<td>5.9</td>
<td>23.2</td>
<td>50</td>
</tr>
<tr>
<td>1B</td>
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<td>6.9</td>
<td>23.4</td>
<td>-30</td>
</tr>
<tr>
<td>5</td>
<td>70.5</td>
<td>9.0</td>
<td>20.5</td>
<td>5</td>
</tr>
<tr>
<td>3B</td>
<td>69.6</td>
<td>12.1</td>
<td>18.3</td>
<td>-145</td>
</tr>
</tbody>
</table>

3. RESULTS AND DISCUSSION
Fig. 1 shows the DSC curves obtained during heating and cooling of alloys 1, 2 and 1A (a) with low manganese content where γ₂ precipitation was expected to occur [4, 6, 7] and of alloys 3B and 5 (b) of high manganese content where formation of bainite was observed. There are two endothermal peaks, one due to a reverse martensitic transformation and another one (visible also for alloy 1A) due to DO₃ disordering above 500°C. All alloys show also one exothermal peak in the range 280-390°C likely due to γ₂ precipitation. Ordering and martensitic transformation are reversible and as such, both show reverse thermal effects during cooling. The temperatures of the ordering transformation lie close to that determined by Prado et al [8] from resistivity measurements.

A similar character possess DSC curves obtained for alloys 3B and 5 (Fig. 1b) although the exothermal effect is broader and shifted to higher temperatures. It is likely caused by the bainite formation [6,7]. The peaks related to ordering and disordering near 500°C are much broader than in the case of Fig. 1a and even clearly splitted for alloy 3B with the highest Mn content. This may be caused by partial DO₃ and L₂₁ ordering at higher Mn content as suggested by Nakanishi et al based on ALCHEMI method [9]. Ordering occurs at lower temperatures when increasing Mn content in agreement with the data of Prado et al [8].
Fig. 1 DSC cooling/heating curves from alloys 1, 2 an 1A (a) and 3B and 5 (b)

DSC curves of alloys 1B and 4 with intermediate Mn and Al content possess a different character. Contrary to the results presented in Fig. 1 no exothermal peak near 350°C can be seen on the heating curves. Peaks due to reversible transformations i.e. martensitic and ordering appear while heating and cooling also during repeated cycling. It results most probably from a low precipitation kinetics at this
composition range as already suggested in [6]. Even heating/cooling at the rate 5K/min do not change this behaviour what suggest usefulness of these alloys for higher temperature applications.

Application of transmission analytical electron microscopy allowed to determine the character of composition changes after continuous heating up to the end of diffusional transformation. Precipitates of $\gamma_2$ show only a few percent composition difference with the matrix what explains a high density of precipitates after continuous heating up to the end of exothermal transformation and the complete degradation of martensitic transformation. Fig. 3 shows the microstructure of alloy 3B continuously heated up to 450°C i.e. to the end of bainitic transformation. Microanalysis across the needle (marked by contamination spots) show an increase of 2 wt% of copper content and decrease of about 1 wt% of Mn and Al content. For thicker plates this difference can be double as high.

Fig. 3 Alloy 3B continuously heated up to 450°C at the rate 20K/min:
(a) Transmission electron micrograph showig a bainitic needle with contamination spots.
(b) Corresponding change of Cu, Al and Mn content across the bainitic needle.

Fig. 4 composition difference between the bainite and the matrix as a function of a plate thickness.
Fig. 4 shows the relationship between the $\Delta x$ (composition difference between the center of a bainitic plate and the matrix) and the plate thickness for alloy 5 continuously heated or isothermally aged at 300°C. It can be seen that $\Delta x$ grows linearly with increasing plate thickness. Extrapolated value of $\Delta x$ to 0 plate thickness do not approach to 0 composition difference. It may indicate that nucleation of bainite occurs with a certain composition difference. However, since the earliest nucleation stages were not observed, the participation of a shear mechanism cannot be excluded.

Fig. 5 shows a fragment of an isothermal section of the ternary CuAlMn system at 850°C with roughly marked estimated $M_s$ temperatures (by dotted lines) and positions of alloys investigated. Arrowheads point compositions of precipitating bainite or $\gamma_2$ phases (depending on what kind of precipitates form in a given alloy). Solid line arrows point compositions after continuous heating, while dotted line arrows after isothermal ageing at 300°C. One can see that arrows coming out of alloy positions 1, 2 and 1A are directed toward the center of $\gamma_2$ field, while those of alloys 5, 1B, 2B and 3B are pointing to the copper rich corner. Compositions of bainite lie within the $\alpha + \beta$ phase field range; only in an extreme case of alloy 5 they are touching the $\alpha$-Cu phase field (not much shifted at lower temperatures [10]), but $\gamma_2$ compositions lie also outside the $\gamma_2$ phase range. Since this phase range shrinks at lower temperatures [10], it may be caused by an microanalysis error due to a small size of $\gamma_2$ precipitates what may influence the results by matrix radiation.

Fig. 5 Fragment of an isothermal section of the CuAlMn phase diagram at 850°C with marked investigated alloys and precipitating phases compositions.
4. CONCLUSIONS.
1. Continuous heating of the CuAlMn alloys containing less than 4.8 at% Mn causes $\gamma_2$ precipitation in the range 280 - 380°C. It is connected with an exothermal heat effect. The additional endothermal peak above 500°C is due to $\text{DO}_3$ disordering.
2. Continuous heating of alloys of higher Mn content (6-12 at%) and lower Al content causes formation of a diffused exothermal peak in the range 320 - 450°C and a broad endothermal heat effect due to $\text{DO}_3/\text{L}_2$ disordering. At highest Mn content the disordering peak shows a clear splitting due to the separation of $\text{DO}_3$ and $\text{L}_2$ effects.
3. Alloys of intermediate Mn (5-6 at%), and Al (23-24 at%) content show only reversible martensitic and ordering transformations during repeated cycling in the range -50$\leftrightarrow$550 °C at the rates 5$\rightarrow$20K/min.
4. The compositions of bainitic plates are shifted towards the copper rich corner in the ternary CuAlMn phase diagram, when compared to the initial alloy compositions being outside the equilibrium $\alpha$-Cu field. With increasing thickness of plates its composition difference with the matrix increases. Compositions of precipitating $\gamma_2$ phase lie close to its equilibrium phase limits.

5. REFERENCES

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