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FIM-ATOM PROBE STUDY OF THE PRECIPITATION IN A Ti$_{46.3}$ - Ni$_{46.7}$ ALLOY

J.J. SWENS, P.F. WILLEMSE$^*$ and J. BEYER$^*$

SGM, Foundation for Advanced Metals Science, PO Box 8039, NL-7550 KA Hengelo (O), The Netherlands
$^*$University of Twente, Faculty of Mechanical Engineering, Laboratory of Materials Science, PO Box 217, NL-7500 AE Enschede, The Netherlands

Abstract - The precipitation process in a B2-TiNi-alloy has been studied during aging between 4373-973 K for various aging times. Earlier reported results obtained with TEM/EDX-analyses showed the development of three different precipitates. The compositions, based on crystallographic calculations were found to be between 55-65 at % Ni. In the present study FIM/AP-analyses have been used to verify these compositional variations. At the early stages of precipitation only Ti$_3$Ni$_4$ is formed which could be analysed both by EDX as well as FIM/AP-analysis. The results of AP-analysis seem to be in better agreement with the calculated value of 57.1 at % Ni than the EDX results.

The FIM images of the alloys show no crystallographic poles which is ascribed to the heavily twinned martensite transformation of the ordered B2-phase during cooling in the FIM. The Ti$_3$Ni$_4$-phase could be identified in the FIM images by the brightness and clear pole formation. The Ti$_5$Ni$_3$- and the Ti$_5$Ni$_9$-phase showed very brittle behaviour in the FIM, and AP-analyses showed better agreement with the calculated values than EDX, but can not be treated as conclusive yet.

1 - INTRODUCTION

TiNi alloys with near equiatomic composition (B2-phase) belong to the practically used shape memory alloys. The remarkable influence of the presence of precipitates on the shape memory effect (SME) like the occurrence of the all-round SME, has initiated a growing interest in the aging behaviour of these alloys. The precipitation process in alloys with excess Ni concentrations has been described as $\beta_1 + \beta_2 + Ti_3Ni_4 + \beta_3 + Ti_5Ni_3 + \beta_4 + TiNi_3$ [1]. The suggested compositions were based on crystallographic analysis of electron diffraction patterns. Chemical analysis by EDX however shows a systematical lower composition for all precipitates. Recently we reported a third metastable precipitate in a 53.7 at % Ni alloy with an equiaxed morphology having a cubic lattice with a lattice parameter close to $a = 0.88$ nm [2]. Field ion microscopy of TiNi-alloy has been reported only in diluted alloys of Ti in Ni [3,4]. The early stages of phase transformation (spinodal decomposition) has been reported in these references. He/He-gas was used for imaging these alloys. A reliable AP-analysis was found for pulse fractions of 20% or higher. For near equiatomic TiNi alloys no information is available as yet. In this paper we will describe the use of FIM/AP-analyses to verify the consistancy of different techniques in obtaining the compositions of the different phases.

2 - EXPERIMENTAL

A binary TiNi alloy with 53.7 at % Ni was prepared from 50.3 at % Ni alloy (Timet Corporation) and Ni of 99.95% purity by arc melting in argon atmosphere. The buttons were remelted seven times for solution treatment. Final homogenization was performed by annealing at 1323 K for 350 ks. in a quartz tube in high pressure argon and quenched by breaking the tube under water. Slices of 0.25 mm thickness were cut using a boron nitride cutting wheel and than aged between 873-973 K for different periods. Specimens were ground from the aged slices and electrolytically polished using a three stage method in 6% perchloric acid in acetic acid and 6% perchloric acid in 2-butoxy ethanol. The original alloy with 50.3 at % Ni was chosen as a reference for FIM-imaging and AP-analysis. Analytical electron microscopy (Jeol 200 CX) was used for crystallographic and morphological study whereas chemical analyses of thin foil specimens were performed with a Link EDX-analyser. A VG-FIM 100 was used for FIM- and AP-analysis. Tip temperatures varying from 30 to 1700 K were used.
80 K and imaging gas mixtures varying from pure Ne to 50/50 NE-HE were investigated to determine the best imaging conditions. The imaging gas pressure was $10^{-1}$ Pa. For AP-analyses high voltage pulse ratios of 15 and 20% were used to prevent preferential evaporation of one component. A fixed tip to screen distance of 55 mm was used. This implies a projected probe hole on the specimen varying from 2 to 4 mm.

3 - RESULTS AND DISCUSSION

Fig. 1 shows a low magnification micrograph of a specimen overaged at 973 K for 7.2 ks. The different morphologies of the metastable phases are indicated whereas the stable TiNi$_3$ intermetallic phase is not yet found. The thin lenticular plates (indicated as 1) are identified as the coherent rhombohedral phase with $a = 0.672$ nm, $\beta = 113.9^{\circ}$ and composition Ti$_3$Ni$_4$. Large plate precipitates (marked 2) can be seen on both sides of the grain boundary. This phase has been identified as Ti$_2$Ni$_3$ with a monoclinic crystal structure with $a = 0.414$ nm, $b = 0.828$ nm, $c = 1.352$ nm and $\gamma = 89.3^{\circ}$. The third identified phase (marked 3) is a cubic phase with $a = 0.88$ nm which is coherent with the B2 phase.

![Electron micrograph of an overaged 53.7 at % Ni specimen with different precipitated phases; 1: Ti$_3$Ni$_4$; 2: Ti$_2$Ni$_3$; 3: Ti$_5$Ni$_9$.](image)

The chemical compositions of these phases have been determined by EDX-analysis on thin foils. The results are given in Table 1. There are some difficulties in interpreting EDX-results in general and the present results in particular which has to be born in mind. The results are obtained with the standardless method using thin foil approximation. Care has to be taken that the analysis is performed on phases which are large enough to maintain through the whole thickness of the foil and that phases are not in diffracting conditions. If not so misleading results can be obtained. Our measured values can be seen from Table 1.

<table>
<thead>
<tr>
<th>Phase</th>
<th>$C_{\text{calc}}$, at % Ni</th>
<th>$C_{\text{EDX}}$, at % Ni</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 (Ti$_3$Ni$_4$)</td>
<td>57.1</td>
<td>55.7</td>
</tr>
<tr>
<td>2 (Ti$_2$Ni$_3$)</td>
<td>60</td>
<td>57.8</td>
</tr>
<tr>
<td>3 (Ti$_5$Ni$_9$)</td>
<td>64.1</td>
<td>59.5</td>
</tr>
</tbody>
</table>
The results show compositions which are systematically lower than those given by the crystallographic calculations. Although these phases are ordered, some deviation from stoichiometry could be explained by the presence of substitutional vacancies due to the aging treatment. Furthermore, the accuracy of the standardless EDX-technique is not better than a few percent which means that small differences in composition like in the present material are difficult to determine.

In order to obtain microchemical information on the different phases firstly the imaging characteristics for FIM had to be determined. Sinclair et al. [3] investigated the local order parameter in dilute alloys of Ti in Ni by FIM. Up till now however little work is available on the imaging of equiatomic TiNi alloys. Primarily we investigated a specimen taken from the reference alloy with 50.3 at % Ni. Fig. 2 shows the micrograph and the overall composition obtained by random area AP-analysis. No indication was found for the development of poles in the image. After variation of the image gas from pure Ne to Ne-He mixtures with 10, 20, 50% He best imaging was found with the 20% He mixture.

Imaging of the material was expected to be difficult due to the following arguments. While cooling the FIM-tip down to 30 K the ordered B2 high temperature phase transforms to monoclinic martensite with a very high density of twins and transformation dislocations. These martensite boundaries are visible as dark bands in the FIM-image (Fig. 2). Additionally, the high electric field may plastically strain the martensite due to its very low elastic limit. This has frequently led to catastrophic failure of the FIM-tips. The difference in ionization fields of Ne for Ti and Ni may cause different contrast of the respective species and it is probable as indicated elsewhere [3] that Ni will be the imaging species. The AP-spectrum also shows a large amount of H\(^+\) originating from the polishing procedure. The compositions of 49.5 at % Ni, obtained by AP-analysis is in satisfactory agreement with the chemical analysis giving 50.3 at % Ni.

Specimens with 53.7 at % Ni, homogenized at 1323 K were subsequently imaged in the FIM. Fig. 3 shows the FIM- and TEM-images. An early stage of homogeneous precipitation of the TiNi\(_3\) phase is visible in the TEM-image. This was confirmed by electron diffraction. The precipitate size was less than 10 nm. The FIM-image shows a small bright area with a clear pole. Although this precipitate which is taken to be TiNi\(_3\), is coherent with the B2 phase no continuity of the observed poles was visible in the matrix. Despite the relatively high Ni-concentration and thus a very low Ms temperature the transformation to martensite may still have occurred giving rise to strains which will mask the poles. The compositions obtained by AP-analysis from the areas indicated as precipitates, were in acceptable agreement with the calculated values. The concentration profile showed fluctuations in the composition which indicate the presence of precipitates. The size of the fluctuations is of the same order as measured from TEM-images.

Fig. 4 shows the TEM-image of the tip for a specimen aged at 873 K for 3.6 ks. TEM-analysis indicated the presence of both TiNi\(_3\) and TiNi\(_4\) precipitates, where TiNi\(_4\) precipitates were found to be relatively large. AP-analyses for the matrix and both precipitates are given in Fig. 5a-
c. The composition profile of the matrix shows fluctuations within a few percent due to the depletion of Ni into the precipitate. The overall matrix composition is 50.9 at % Ni. The composition for the Ti$_3$Ni$_4$ precipitates were found to vary between 56.6 and 57.4 at % Ni and for Ti$_2$Ni$_3$ precipitates between 59.6 and 60.6 at % Ni, all again in acceptable agreement with the calculated values. In specimens aged at 973 K for 7.2 ks, the same composition for the Ti$_2$Ni$_3$ was found and occasionally a precipitate composition of 65.2 at % Ni was measured. Although these results are not conclusive they strongly indicate that using FIM/AP-analysis accurate measurements of compositional variations due to precipitation in TiNi-SME alloys are possible.

Fig. 3a - FIM-micrograph of the homogenized 53.7 at % Ni alloy

Fig. 3b - Electron-micrograph and diffraction pattern of the homogenized 53.7 at % Ni alloy tip

Fig. 4 - Electron-micrograph of the 53.7 at % Ni alloy tip aged at 873 K for 3.6 ks
Fig. 5 - AP composition profiles of the 53.7 at % Ni alloy aged at 873 K during 3.6 ks for a) matrix b) Ti₃Ni₄ precipitates c) Ti₂Ni₃ precipitates. T = 55 K
4 - CONCLUSIONS
- FIM-imaging near equiatomic TiNi alloys has been shown to be possible with a Ne-He mixture at 30 K.
- Ti$_3$Ni$_4$ precipitates show clear poles. Specimens with fine homogeneously nucleated precipitates show to be more stable with respect to failure than those with coarse precipitates.
- Formation of martensite during cooling to imaging temperatures can cause early failure of the tips.
- Compositions of the matrix, Ti$_3$Ni$_4$ and Ti$_2$Ni$_3$ precipitates obtained by AF-analysis are in better agreement with those calculated from crystallographic data than the compositions obtained with EDX-analysis.

5 - REFERENCES