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Abstract

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Keywords

tomography, martensite, during, heat, treatment, steels, atom, observations, probe, decomposition

Disciplines

Engineering | Science and Technology Studies

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Observations of decomposition of martensite during heat treatment of steels using atom probe tomography

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Abstract. The decomposition of martensite during tempering or ageing is an important phenomenon as it leads to changes in the mechanical properties. These changes could take place during both steel manufacturing or in-service. Thus, their understanding is required to predict the material performance. Recent advances in the development of local electrode atom probes has led to a significant increase in the analysed volume of material (up to 100 millions of atoms) and at the same time reduced the acquisition times. This allows improvement in data statistics when investigating fine nanoscale features, such as solute segregation, clustering and ultrafine precipitation. Selected results of atom probe studies on the decomposition of martensite from bake hardening of a pre-strained Transformation Induced Plasticity (TRIP) steel and ageing of FeNiTiMnAl maraging steel are presented.

1. Introduction

Recent developments of local electrode atom probes [1-3] open new opportunities for analysis of martensite decomposition at the atomic level. These modern instruments allow the collection of data in a shorter period of time from a larger volume of material compared to the previous generation atom probe field ion microscopes. As a result, a more accurate and statistically better analysis of chemical composition, shape and distribution of fine microstructural features present in the material, such as nanoscale precipitates and clusters, is achieved.

For decades, the decomposition of martensite in various steel grades has attracted the attention from both fundamental and applied perspectives. This process leads to modification of the mechanical properties, which could have both beneficial and negative effects depending on the stage of martensite decomposition and the desirable in-service characteristics of the material. Whereas, there is a general agreement on the stages of decomposition taking place during tempering of carbon-containing ferrous martensite [4,5], some discrepancies still exist on the nature of the initial carbides formed. However, the atom probe tomography not only provides insight into the early stages of clustering in martensite, but also highlight the details of the precipitation sequence at a later stage.

Transformation-induced plasticity (TRIP) steels are promising candidates for applications in the automotive industry [6-9]. After forming, the automotive body is subjected to a paint baking process (typically for 20 minutes at 175-180°C), which results in an additional strength increase [7-9]. The microstructure of the TRIP steels consists of polygonal ferrite, carbide-free bainite morphologies, retained austenite and martensite. As previous studies using electron microscopy have shown [9-13], during pre-straining and bake hardening various microstructural changes take place: increase in dislocation density, especially in polygonal ferrite in the vicinity of hard martensite or retained austenite crystals, strain-induced transformation of austenite to martensite and precipitation of fine carbides in the bainitic ferrite.

FeNiTi (Mn,Al) maraging steels exhibit an unusual rapid hardening phenomenon after just 5 s ageing at 550°C [14-17]. In this condition it achieves approximately 50% of the maximum increase in strength without significant sacrifices in ductility. Based on the transmission electron microscopy data, there is no indication of precipitate formation during ageing times below 60 s. Thus, atom probe tomography is the most suitable technique to elucidate the decomposition of martensite in this steel.

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In this paper we present selected atom probe studies on decomposition of martensite during bake hardening of the TRIP steel, subjected initially to thermomechanical processing, and on the ageing behaviour of Fe-Ni-Mn-Ti-Al maraging steel.

2. Experimental

2.1 TRIP steel

The compositions and selected processing parameters for the studied TRIP steel are given in Table 1. It was subjected to thermomechanical processing simulation using a laboratory rolling mill at Deakin University. The details of the processing schedule are described elsewhere [13]. After the simulation of coiling in a fluidized bed furnace, the samples were quenched to room temperature. Tensile specimens were machined from the strip and subjected to 4% pre-straining (PS) before bake hardening (BH) at 175 °C for 60 min.

Table I. TRIP steel compositions and thermomechanical processing parameters

Steel		Element								Parameters		
		C	Si	Mn	Mo	Al	Cu	Nb	P	T _{AC} , °C	T _{IH} , °C	t _{IH} , s
AR	wt%	0.21	1.18	1.52	0.29	0.57	0.03	0.036	0.027	780	450	1200
	at%	0.95	2.3	1.51	0.16	1.15	0.026	0.02	0.05			
M-BH	at%	1.5 ±0.07	3.7 ±0.1	1.11 ±0.06	0	1.64 ±0.07	0.07 ±0.02	0.06 ±0.01	0.03 ±0.01			

Fe is balance. AR-as-received, M-BH - martensite after bake hardening, T_{AC}- accelerated cooling start temperature; T_{IH}- isothermal hold temperature, t_{IH}-holding time

Atom probe tomography (APT) studies were carried out using the Oxford nanoScience 3DAP at the Monash Centre for Electron Microscopy. The pulse repetition rate was 20kHz and the pulse fraction was 0.2. The sample temperature was 60K.

2.2 Maraging steel

An experimental steel with the following composition: Fe-20.1Ni-1.8Mn-1.6Ti-0.59Al-0.04Si-0.01C with P and S < 0.001 (wt%) was solution heat treated at 1100 °C for 12 h. Thin slices of ~0.4mm thickness were then cut and solution treated for 1h at 1050 °C. To complete the martensite transformation after water quenching the slices were immersed in liquid nitrogen for 10 minutes. Ageing was carried out in salt bath out at 550 °C. Detailed description of processing is given elsewhere [17]. Selected samples, which corresponded to the conditions of the best combination of strength and ductility (5 s ageing) and to the peak strength (600 s), were chosen for atom probe analysis. Data was acquired from a needle-shape sample inserted into a local electrode atom probe at the University of Sydney operating at a pulse repetition rate of 200 kHz, a pulse fraction of 0.2, and a sample temperature of 80K.

2.3 Atom probe data analysis

A detailed analysis of clusters and fine precipitates was carried out using the maximum separation envelope method [18] with a maximum separation distance between atoms of interest of 0.5 nm and a grid spacing of 0.1 nm. In order to minimise random solute fluctuations in the matrix, a minimum of 20 atoms was used to define the clusters. Their atomic composition was calculated from the number of atoms of each type forming a cluster or a particle as determined by the maximum separation envelope method. However, it should be noted that the maximum separation method aggressively removes the solvent atoms from particles and clusters, especially from the surface regions. Thus, using this method the solute content of the finest clusters (≤ 1 nm), in which there is a high proportion of surface to interior atoms, may be overestimated by ~25%. Using the data from the maximum separation method, the observed clusters and fine particles were grouped together based on the selection of non-overlapping ranges for their solute content. As output of the program, the size of the feature of interest is given by the radius of gyration (l_g), which is slightly smaller than its actual physical extent. The real size might be better represented by the Guinier radius (r_G) given by the following equation [18]:

$$r_G = \sqrt{\frac{5}{3}} \times l_g \quad (1)$$

The compositions of relatively large precipitates and of the matrix volumes free of visible clusters or precipitates were determined from the selected volumes with background noise subtraction based on the number of atoms. Concentration profiles were taken perpendicular to the interface of the feature of interest. Isoconcentration surfaces were also used to visualize the microstructural features.

3. Results and Discussion

3.1 Effect of bake hardening on martensite in TRIP steel

After TMP the multi-phase microstructure of TRIP steel consists of $15 \pm 3\%$ of polygonal ferrite, $16.5 \pm 3\%$ of retained austenite, $\sim 65 \pm 3\%$ of carbide-free bainite and the remainder martensite. As previous transmission electron microscopy studies have shown [10,13], the martensite was present in the form of thin layers between the bainitic ferrite laths or in a blocky form as part of the martensite/austenite constituent. However, it should be noted, that at the test temperature of atom probe studies all retained austenite was deemed to be also transformed to martensite. During cooling of strip to room temperature, some of the retained austenite crystals present in the microstructure, which are either chemically or mechanically unstable, transform to martensite. As previous research has shown, this martensite auto-temperers with the formation of fine (20-50 atoms) C-rich clusters [12,19]. Additionally, strain-induced martensite forms during pre-straining [9,13]. All of these types of martensite crystals undergo tempering during bake hardening.

The non-uniform segregation of C atoms is clearly visible in the atom map shown in Fig. 1a. The clusters

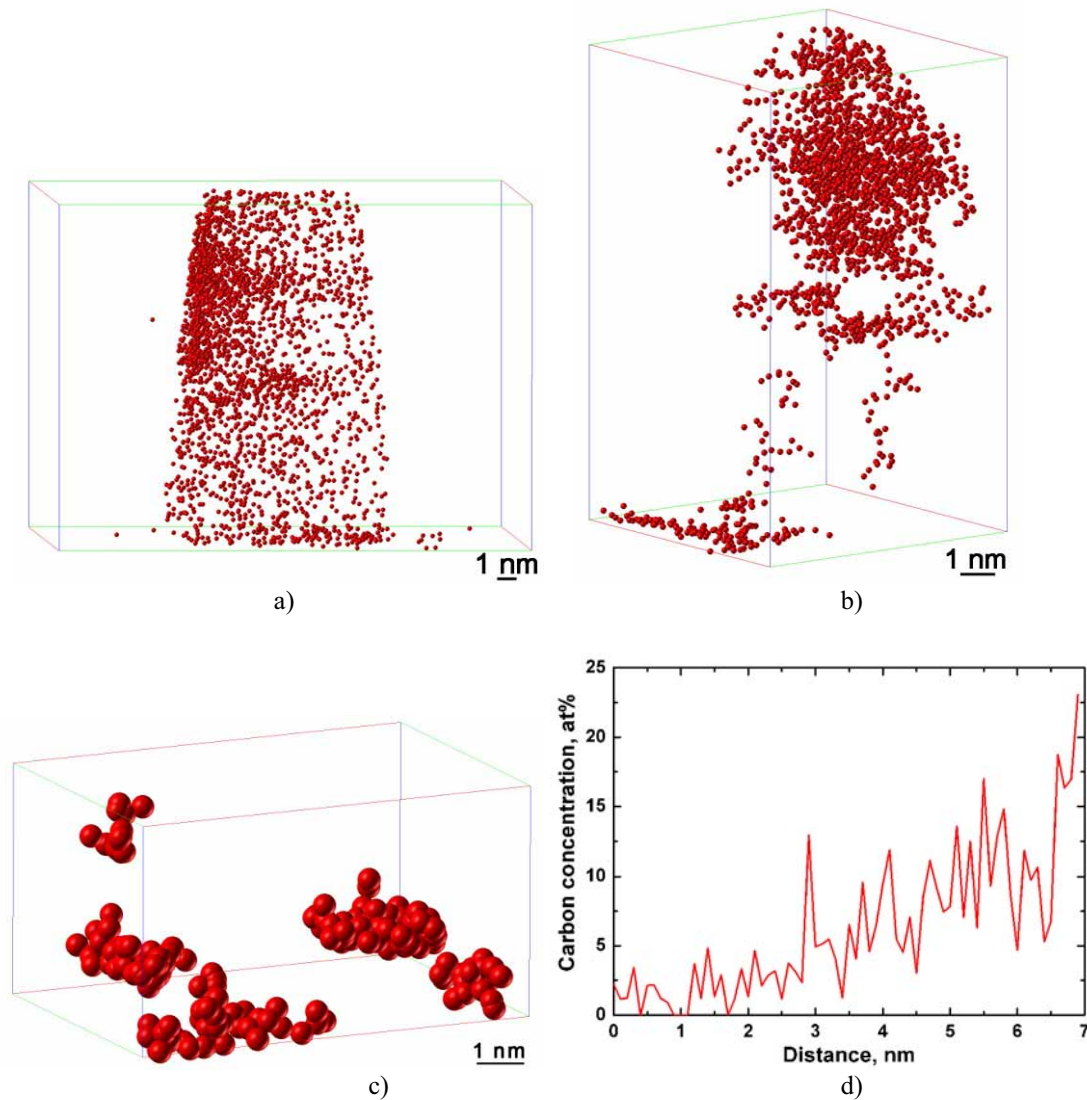


Fig. 1. C atom map (a), clusters (b and c) and representative C concentration profile across matrix/carbide interface (d) in the martensite of TRIP steel after bake hardening. All other atoms are removed in (b) and (c) by the maximum separation envelope method.

determined by the maximum distance separation method are shown in Fig. 1b from different points of view in order to give a better insight into the morphology of the clusters/fine carbides. An enlarged selected area with clusters is shown in Fig. 1c. The concentration profile indicates that the carbon concentration within the coarse carbide reaches approximately 20at%. The summary of the clusters composition and sizes are given in Table 2.

It is clear that the fine clusters are predominantly C-rich, as expected and in agreement with the results obtained for the early stages of martensite decomposition in the TMP condition [12]. They do not have the composition of Fe_4C , as suggested by Sherman *et al.* [20], but are close to the composition of FeC_8 . The density of clusters and fine precipitates is an order of magnitude higher than in the martensite after TMP [19,21]. With an increase of cluster/precipitate size, the C content decreases and the coarsest precipitates reach the composition of Fe_4C . It is surprising that it is not the equilibrium composition of Fe_3C , as was previously observed for several bake-hardened TRIP steels. It could be assumed that due to the limited volume of material analysed in this case the Fe_3C was not detected. However, the coarse precipitates, visible under TEM, were of that composition [13,21]. It is also worth noting, that the matrix composition of martensite (Table I) still shows a significant supersaturation in carbon, indicating that decomposition has not yet been completed. This is in agreement with other studies on the tempering of martensite using APT [21]. Based on the combined body of knowledge to-date, it can be concluded that a continuous set of iron carbide compositions from C-rich clusters to Fe_3C carbides is present in the tempered martensite.

Table II. Characterisation of clusters and carbides in the martensite of TRIP after bake hardening

Composition, at% (AP)					
C	Mo	Nb	Fe	r_G , nm	No. ions per feature
89.4±5.3	-	-	8.1±4.8	1.1±0.1	28±6
51.5±2.9	3.0±0.6	1.9±0.7	26.8±2.4	3.1±0.5	302±42
19.7±0.3	0.3±0.1	0.3±0.1	74.0±0.4*	4.8±0.5	11255± 230

*Traces of Mn and other elements

3.2 Decomposition of martensite in maraging steel

Analysis of the sample after 5 s ageing has clearly shown the solute segregation in the martensite crystals. Al, Ti and Mn atom maps (Fig. 2) depict the cluster formation. A more detailed analysis using the maximum separation envelope method has shown the preferential formation of Ti+Al and Mn+Fe clusters (Table III).

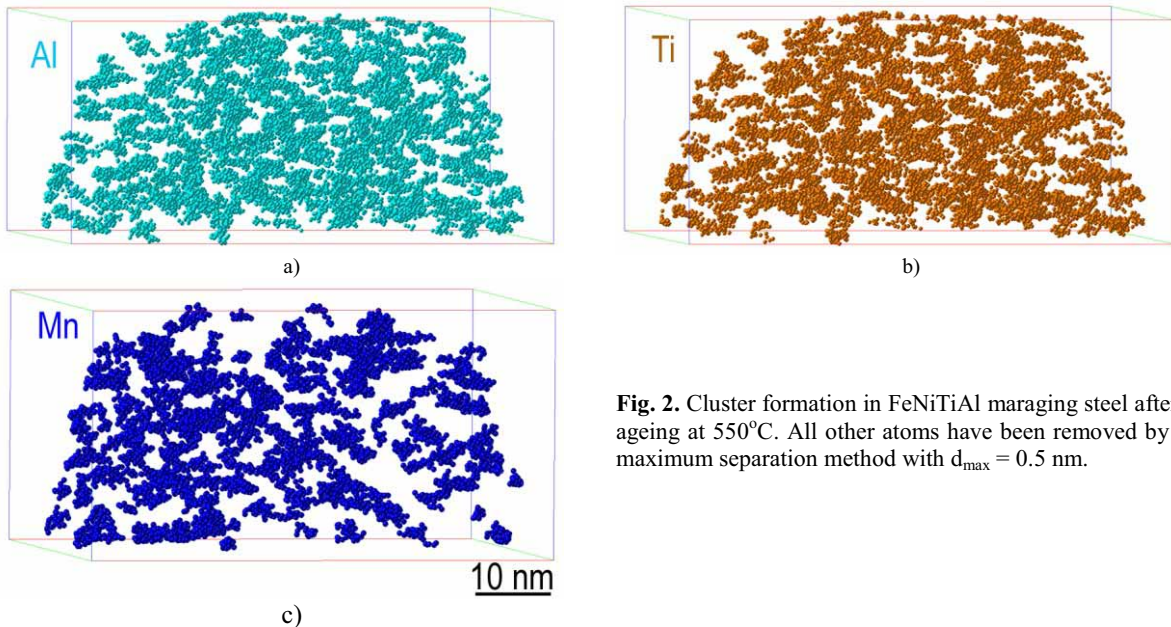


Fig. 2. Cluster formation in FeNiTiAl maraging steel after 5s ageing at 550°C. All other atoms have been removed by the maximum separation method with $d_{\max} = 0.5$ nm.

Whereas the composition of Mn-Fe clusters varies, their size remains constant. Contrarily, with an increase in size, Ti+Al clusters become richer in Fe and Ni, which could be explained by the deficiency of the maximum separation method, as described in section 2.3. However, there is no doubt in affinity between Mn-Fe and Ti-Al atoms, which results in the formation of these clusters. This is in agreement with the previous Mössbauer spectroscopy data on preferential formation of Fe-Mn bonds and displacement of Ti by Mn from solid solution,

which leads to the accelerated precipitation of Ni_3Ti [15]. Mn also has a strong affinity with Al [22] and with the progress of ageing the formation of nearly spherical $(\text{Fe,Ni})_3(\text{Mn,Al})$ particles was observed (Fig. 3). It could be suggested that not only dislocations, but Fe-Mn clusters, served as heterogeneous nucleation sites for these precipitates.

Table III. Characterisation of clusters and precipitates after 5 and 600s ageing, respectively.

Clusters / precipitates	Composition, at. %						r_G , nm/ xyz precipitates	No. ions per cluster/morphology
	Fe	Ni	Mn	Ti	Al			
Mn+Fe	0-18	0-12.5	65 -100	-	-		1.0±0.2	20-146
Ti+Al-rich	25-36	3 -18	-	16 -39	17 -47		1.4±0.3	28-441
	2-24	0-22	-	16-75	14 -72		1.1±0.2	22-177
	0	0-11	-	18-80	21-74		0.8±0.1	20-62
(Ni,Fe)₃Ti	9 ± 3	60 ± 3	1.7 ± 0.6	22 ± 2	3.1 ± 0.6		~9x3x16 ~9x8x14	plate-like rod-like
(Ni,Fe)₃(Al,Mn)	27 ± 5	40 ± 6	10 ± 4	2 ± 1	17 ± 3		~4x4x5	spheroidal

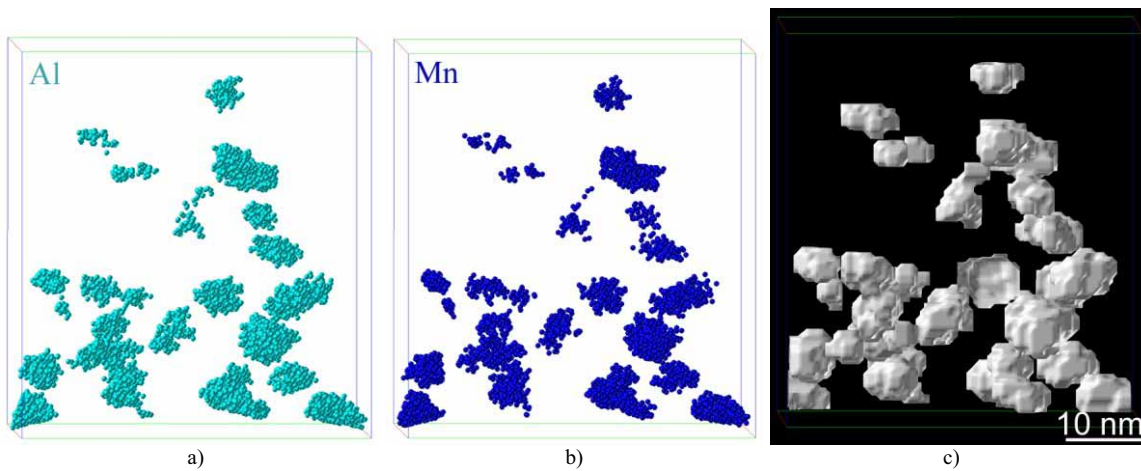


Fig. 3. Al (a) and Mn (b) atom maps with corresponding 30%(Al+Mn) iso-concentration surfaces showing formation of $(\text{Fe,Ni})_3(\text{Mn,Al})$ particles after 600 s ageing at 550°C.

After 600s ageing in addition to spherical precipitates, complex arrangements of plate-shaped and rod-like particles were clearly visible (Fig. 4). The selected volume analysis of these precipitates has shown that their composition is close to $(\text{Ni,Fe})_3\text{Ti}$ (Table III). It could be suggested that Ti+Al-rich clusters together with substructure were preferential nucleation sites for this precipitation. Based on TEM data, there was a long standing debate regarding the exact shape of Ni_3Ti precipitates in maraging steels being plate-like [16], needle-like [23-25] or rod-like [24,26]. As we now could conclude, both plate and rod morphologies of these precipitates co-exist at certain stages of decomposition. However, our atom probe tomography (APT) data also indicates that with further coarsening both type of precipitates, e.g. $(\text{Ni,Fe})_3\text{Ti}$ and $(\text{Ni,Fe})_3(\text{Al,Mn})$, tend to exhibit a predominantly rod-like morphology.

4. Conclusions

It has been shown that atom probe tomography is a powerful technique which allows complete chemical and 3D-spatial information on nanoscale features in steels to be obtained. The evolution from clustering to precipitation could be followed using APT, as was highlighted using TRIP and maraging steels as examples. However, some deficiencies of APT are noted and care needs to be taken when interpreting results for the features with the sizes about $r_G=1\text{nm}$.

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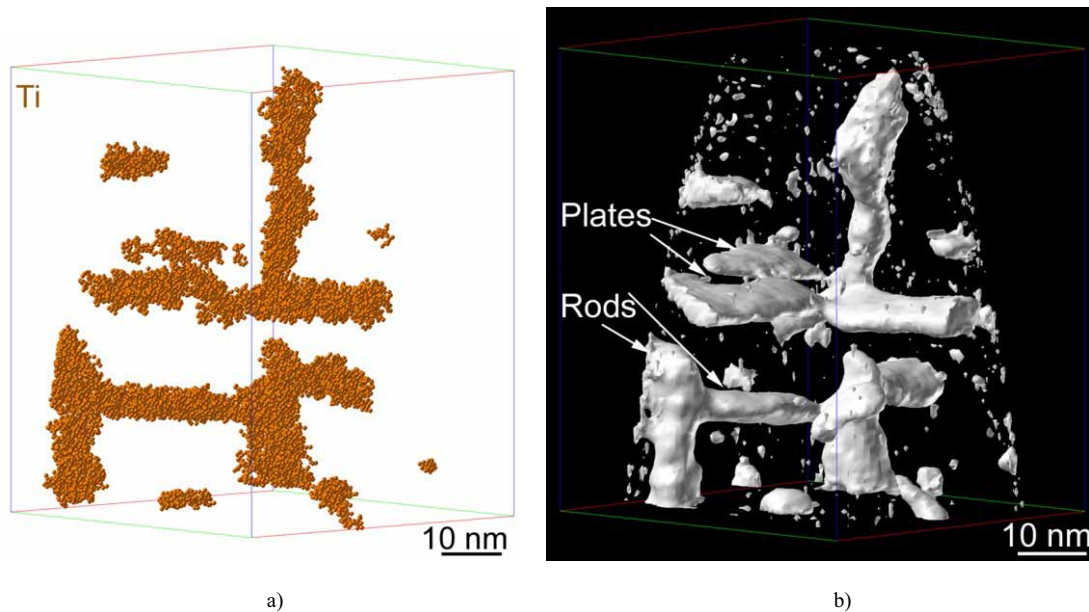


Fig. 4. Ti map (a) and 40% (Ni+Ti) iso-concentration surfaces (b) showing formation of predominantly $(\text{Fe,Ni})_3\text{Ti}$ precipitates in martensite after 600 s ageing at 550°C. All other atoms removed by maximum separation method in a).

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