Continuous Heating Dissolution and Continuous Cooling Precipitation Diagrams of a Nickel-Titanium Shape Memory Alloy

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Abstract Binary NiTi alloys are the most common shape memory alloys in medical applications, combining good mechanical properties and high biocompatibility. In NiTi alloys, the shape memory effect is caused by the transformation of an austenite phase to a martensite phase and the reverse process. Transformation temperatures are strongly influenced by the exact chemical composition of the NiTi phase and the presence of precipitates in the microstructure induced by thermo-mechanical treatment, especially solution annealing and ageing. Isothermal time-temperature precipitation diagrams can be found in the literature. Cooling is frequently not considered, as water quenching is typically assumed to be sufficient. To the best of our knowledge, continuous heating dissolution (CHD) and continuous cooling precipitation (CCP) diagrams do not exist. Differential scanning calorimetry (DSC) is a common method to analyse the austenite/martensite transformation in shape memory alloys, but it has not yet been used to analyse precipitation processes during continuous temperature changes. We have enabled DSC to analyse dissolution and precipitation processes in situ during heating as well as during cooling from the solution annealing temperature. Results are presented as CHD and CCP diagrams, including information from microstructure analysis and the associated changes in the austenite/martensite transformation temperatures.

Keywords NiTi • Differential scanning calorimetry (DSC) • Continuous heating dissolution diagram (CHD) • Dissolution • Continuous cooling precipitation diagram (CCP) • Precipitation

Introduction

NiTi alloys are the most common shape memory alloys in medical applications, combining good mechanical properties and high biocompatibility [1–5]. In binary NiTi alloys, the shape memory effect is characterised by the transformation of austenite (B2-structure) to martensite (B19\textsubscript{0}-structure) and the reverse process [6]. The low-temperature B19\textsuperscript{0} martensitic phase forms during cooling of the high-temperature B2 austenite phase [7]. In the solution-annealed and quenched condition, the transformation takes place in one step from B2 to B19\textsuperscript{0} during subsequent heating or cooling in the temperature range from −100 to +100 °C. The transformation temperatures are strongly influenced by the exact chemical composition of the NiTi phase and the presence of precipitates induced by thermo-mechanical treatments, especially solution annealing and ageing. The formation of Ni\textsubscript{4}Ti\textsubscript{3} precipitates during ageing or dislocations due to thermo-mechanical treatment hinders the formation of B19\textsuperscript{0}. Depending on the size and density as well as the nucleation site density, these precipitates disturb the direct transformation from B2 to B19\textsuperscript{0}. Beyond that, a second martensitic phase called the intermediate R-phase exists [3]. It has been observed that in the solution-annealed, quenched, and aged stage, the transformation...
takes place in two steps in the temperature range from -100 to +100 °C. The initial transformation is from B2 to R-phase, which transforms to B19' and the reverse [3, 7, 8]. The precipitation sequence during isothermal ageing treatment in NiTi alloys can be described as follows:

\[ \begin{align*}
  \alpha & \xrightarrow{\text{supersaturated B2 NiTi}} \alpha_1 + \text{Ni4Ti3} \\
  & \quad \xrightarrow{\text{}} \alpha_2 + \text{Ni3Ti2} \\
  & \quad \xrightarrow{\text{}} \alpha_3 + \text{Ni3Ti}
\end{align*} \]  

(1)

\( \alpha_1, \alpha_2, \) and \( \alpha_3 \) are matrices with different Ni concentrations [11]. Typically, ageing is done below 600 °C in the two-phase region NiTi and Ni3Ti according to the NiTi phase diagram [10, 12]. During ageing at lower temperatures and shorter ageing times, Ni4Ti3 appears in the microstructure, while at higher temperatures and longer ageing times, the equilibrium phase Ni3Ti will be precipitated. For intermediate temperatures and times, the metastable Ni3Ti2 forms. Besides, it is known that the precipitation process and resulting transformation temperatures are also dependent on the heating rate to the solution annealing/ageing temperature as well as the cooling rate from the solution annealing/ageing temperature [8, 11, 13–16]. However, currently, no continuous heating dissolution (CHD) and continuous cooling precipitation (CCP) diagrams for NiTi alloys exist, and only isothermal time–temperature transformation diagrams can be found in the literature [1, 11, 17].

Differential scanning calorimetry (DSC) is a common method to analyse the austenitic and martensitic transformation in shape memory alloys (see, for example, Refs. [1, 7, 9, 10]), but it has not yet been used to analyse the phase transformations according to the precipitation sequence of Ni3Ti (Eq. 1). Recently, DSC has successfully been used to establish CHD and CCP diagrams for several alloying systems, like Al, Fe, Mg, or Ni [18–20]. In this work and for the first time, the DSC method was applied to the analysis of dissolution and precipitation kinetics during continuous temperature changes in a binary NiTi alloy.

**Materials and Methods**

The investigated alloy was a Ni-rich binary NiTi shape memory alloy. In the delivered condition, the material was a Ø 6 mm cylindrical rod with an austenite finish temperature of -5 °C ± 10 K. The cylindrical rod was formed by means of round forging from a cast block that was manufactured using vacuum induction melting (VIM). The present alloy was delivered with a nominal composition of 50.9 at.% Ni and 49.1 at.% Ti. Before the DSC experiments, all samples were solution annealed (1000 °C for 30 min in the Setaram Labsys Evo DSC, 700 °C for 30 min in the Perkin Elmer Pyris Diamond DSC/8500 DSC) and cooled at a rate of 0.1 K/s to create a defined initial condition for subsequent experiments. The sample dimensions and resulting sample masses for the three used types of DSC devices are as follows:

- Setaram Labsys EVO DSC: Ø 4.9 mm x 11.6 mm, \( \approx 1420 \) mg.
- Setaram Sensys Evo DSC: Ø 5 mm x 14.4 mm, \( \approx 1850 \) mg.
- Perkin Elmer Pyris Diamond DSC/8500 DSC: Ø 6 mm x 1 mm, \( \approx 180 \) mg.

The applied scan rates were varied over a wide dynamic range, from 0.01 to 5 K/s for the heating experiments and from 0.01 to 1 K/s for the cooling experiments. Further cooling experiments with 0.001 to 0.03 K/s rates were performed for metallographic analysis. The DSC data evaluation for the heating experiments was performed according to Ref. [21] and for the cooling experiments according to Ref. [22]. The experimental setup to perform high-temperature DSC with solution annealing temperatures of 1000 °C in a heat flux DSC Setaram Labsys EVO is described in Ref. [20]. The technical ceramic Rescor960 with the same geometry as the sample was employed as an appropriate reference material for the baseline measurement. DSC samples were packed in commercial ceramic crucibles. Two sample measurements and one baseline measurement were performed for each parameter setup.

For DSC experiments with temperatures up to 700 °C, the heat flux DSC Setaram Sensys Evo and the power compensated Perkin Elmer Pyris Diamond DSC were used. According to the Ni-Ti phase diagram, a temperature of 700 °C is sufficiently high to anneal in the single phase field of NiTi [12]. Later on, the DSC experiments will prove this assumption. In this case, the stable \( \alpha \)-titanium alloy Ti5Al2.5Sn with the same geometry was used as a reference material for maximum temperatures of 700 °C.

To investigate the influence of quench-induced precipitation during cooling from the solution annealing temperature on the martensite transformation, the determination of transformation temperatures from austenite to martensite and the reverse was conducted. Therefore, the standard test method described in ASTM F2004-05 was adapted using a Perkin Elmer Pyris Diamond DSC with sample dimensions of Ø 6 mm x 1 mm. After solution annealing at 700 °C for 30 min and cooling with different rates, a common ageing treatment at 500 °C for 20 min was performed in a protective nitrogen atmosphere. Afterwards, the temperature range was set from -80 to +100 °C at controlled heating and cooling rates of 0.05 K/s. Fig. 1 shows the applied time/temperature profile for determination of the transformation temperatures. In the case of these experiments, one sample was scanned multiple times in a series of varied cooling rates after solution treatment. This experiment
series focussed on the determination of variations in the characteristic transformation temperatures. No baseline measurements were done in this special case. This is to allow programming of the series in one experiment-program, and as the raw heat flow was found to be sufficiently precise to allow evaluation of relative variations in the transformation temperatures.

To analyse the changes in the microstructure related to the reactions detected in DSC, optical microscopy (OM) as well as scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDS) were performed. The heat-treated samples were cold-embedded in epoxy resin and then mechanically ground and polished with water-free lubricants. The final polishing was done with a 0.05 μm oxide suspension. The polished samples were rinsed and etched as per following the etching agent: 62.5 ml water H2O, 18.75 g potassium metabisulfite K2S2O5, 12.5 ml hydrochloric acid HCl, and 1.5 g ammonium hydrogen fluoride NH4HF2. Microstructural images were acquired by optical microscopy using a LEICA DMI5000 M, Co. Leica Microsystems GmbH, Wetzlar, Germany.

The SEM samples were analysed by a field emission SEM (MERLIN®VP Compact, Co. Zeiss, Oberkochen, Germany) equipped with an EDS detector (XFlash 6/30, Co. Bruker, Berlin, Germany) and analysis software (Quantax400, Co. Bruker, Berlin, Germany). SEM-secondary electron (SEM-SE) images were obtained using a high efficiency Everhart–Thornley-type HE-SE detector at 5 kV acceleration voltage. Representative areas of the samples were analysed and mapped to determine the elemental distribution based on the EDS-spectra data by the QUANTAX ESPRIT Microanalysis software (version 2.0).

The embedded and polished samples were mounted on the SEM carrier with adhesive conductive carbon and aluminium tape (Co. PLANO, Wetzlar, Germany). The acceleration voltage for the EDS analysis was set to 20 kV.

Figure 2 shows metallographic images obtained by SEM (polished and unetched) in the initial condition of the NiTi. In the cross section in Fig. 2a and b, many small black particles can be observed in the grey matrix. In Fig. 2c and d, images are displayed in the longitudinal direction where aligned particles are evident. EDS proves that the inclusions show a high concentration of titanium and carbon. Furthermore, the nickel content is significantly lower than in the surrounding matrix. The large size (up to a few hundred micrometres) and the high density of those particles correlate with non-metallic inclusions like titanium carbides and oxides, which were also observed in Ref. [23–25]. According to the literature, these inclusions can be traced back to the primary shaping process.

Results and Discussion

Heating Experiments

DSC curves during continuous heating using a broad range of heating rates are shown in Fig. 3. For a better understanding, the curves have been offset downwards from the slowest rate on top to the fastest at the bottom. Horizontal dotted lines correspond to the zero level for the curves. Upward deviation of the DSC curve from the zero level indicates domination of endothermal reactions, whilst downward deviations of the DSC curve from the zero level indicate domination of exothermal reactions. One must be aware that the (partly opposing) reactions might superimpose and DSC is just measuring the sum of all heat effects at present. Three different DSC devices were used, as shown by the black, red, and blue curves. In general, there is a good match between the three types of DSC devices used to cover a wide range of heating rates as well as high temperatures.

The DSC-heating curves of 0.01 K/s show an exothermic precipitation reaction between 230 and 430 °C with a distinct peak at 300 °C (peak a). Additionally, an overlapping precipitation reaction appearing as a shoulder at 350 °C can be seen (peak b). This holds for rates up to 0.1 K/s, while the shoulder is no longer seen at higher heating rates. It is believed that in the context of these reactions, parts of the precipitation sequence of Ni3Ti (Eq. 1) occur. After crossing the zero level at 430 °C, a subsequent endothermal dissolution reaction with a peak at 580 °C can be observed (peak C). It can be assumed that previously precipitated particles will be dissolved during this endothermal reaction. It can be seen that the
dissolution reaction is finished at 600 °C and solution annealing above 600 °C is sufficient to dissolve the alloying elements. A further increase in the temperature results in a horizontal progress of the DSC curve at zero level, indicating a reaction free zone. With increasing heating rate, the precipitation and dissolution reactions and corresponding DSC peaks are significantly shifted to higher temperatures (indicated by the vertical grey spline). In the same order, they are increasingly suppressed. For the fastest investigated heating rate of 5 K/s, the peak of the exothermal precipitation reaction can be seen at 530 °C, which means a shift of about 230 K in comparison to heating at 0.01 K/s. The peak of the endothermal dissolution reaction shifts from 580 °C at 0.01 K/s to 620 °C at 5 K/s. The suppression of ongoing reactions during heating can firstly be seen by the disappearance of the shoulder during the exothermal reaction for heating rates higher than 0.1 K/s as well as by the reduction of areas under the exothermal and endothermal peaks.

Figure 4 shows the resulting continuous heating dissolution (CHD) diagram for the specific batch and initial condition of the NiTi alloy 50.9 at.% Ni and 49.1 at.% Ti are investigated. Therefore, start and end temperatures of characteristic precipitation and dissolution reactions were evaluated and plotted in a temperature/time diagram. From the CHD diagram, during heating to typical ageing temperatures, distinct precipitation and dissolution reactions occur. This means that depending on the considered temperature and heating rate, certain stages in the precipitation sequence of Ni₃Ti (Eq. 1) can be reached prior to the actual ageing treatment. This aspect must be considered when selecting ageing temperatures and times as premature
precipitation of nickel rich particles can significantly influence the transformation temperatures. In consequence different heating rates, especially slower heating rates, can lead to different transformation temperatures under otherwise identical ageing conditions. After running through the different areas of precipitation and dissolution, the heating rate specific solvus temperature is reached. Above this temperature, all major alloying elements are in a solid solution.

Cooling Experiments

DSC curves from continuous cooling using a broad range of cooling rates are shown in Fig. 5. For a better understanding, the curves have been offset downwards from the slowest rate on top to the fastest at the bottom. Horizontal dotted lines correspond to the zero level for the curves. Upward deviation of the DSC curve from the zero level indicates exothermal reactions. During cooling, only exothermic precipitation is expected; thus, the issue of superimposed reactions is less critical for curve interpretation. Three different DSC devices were used, which are shown by the black, red, and blue curves. Also, during cooling, there is a good match between the three types of DSC devices used to cover a wide range of cooling rates as well as high-temperatures. DSC curves after solution annealing at 700 and 1000 °C are nearly identical. This agrees with the CHD diagram in Fig. 4.

Considering a cooling rate of 0.01 K/s, one obvious exothermal precipitation reaction occurs between 420 and 310 °C. With increasing cooling rate, the precipitation reaction is shifted to lower temperatures and gets increasingly suppressed. The peak temperature for 0.01 K/s is 395 °C, while it is shifted to 275 °C at 1 K/s. However, though the DSC curve of 0.01 K/s is quite noisy, another weak, preceding precipitation peak might be present at 520 °C. This assumption is supported by the shoulder seen on the high-temperature side of the subsequent exothermic peak at 0.03 K/s and 0.05 K/s. It can be assumed that this reaction at higher temperatures would be even more pronounced at slower cooling as considered here.

Figure 6 shows the reduction of the total specific precipitation enthalpy with increasing cooling rate. The specific precipitation enthalpy can be calculated by integrating the peak area. In this case, the highest specific precipitation enthalpy is 3.6 J/g for the slowest cooling rate of 0.01 K/s. This value is nearly constant for the next two faster cooling rates. For 0.1 K/s, the specific precipitation enthalpy drops down to 1.7 J/g and decreases further with the increase in the cooling rate. For the highest investigated cooling rate of 1 K/s, a precipitation enthalpy of just 0.3 J/g remains, which is already close to the detection limit of DSC [18, 22]. From this, it can be concluded that the critical cooling rate to completely suppress quench-induced precipitation is just slightly higher than 1 K/s (considering a logarithmic cooling rate increment).

The start and end temperatures of the precipitation reaction during cooling from 1000 °C, respectively, to 700 °C for 30 min were evaluated. The corresponding
time–temperature data points are plotted in Fig. 7 showing the CCP diagram for the specific batch of the NiTi alloy 50.9 at.% Ni and 49.1 at.% Ti. The two fastest cooling rates were performed starting from a temperature of 700 °C; nevertheless, these two were integrated into the same diagram. The critical cooling rate is estimated to be 3 K/s, as the specific precipitation enthalpy for 1 K/s is almost zero (see Fig. 6). The dashed line proposes the extrapolated start and end temperature for quench-induced precipitation limited by the critical cooling rate at 3 K/s.

It can be assumed that quench-induced precipitation of phases like Ni₃Ti₃, Ni₃Ti₂, or Ni₃Ti from the precipitation sequence towards Ni₃Ti (Eq. 1) is observed here. As only one clear exothermal precipitation peak occurs at a cooling rate of 0.01 K/s, it is expected that slower cooling would result in multiple overlapping precipitation peaks regarding the precipitation sequence towards Ni₃Ti. This behaviour is well known from other alloying systems, for example, on the basis of Al, Fe, or Mg [18, 19].

Transformation Behaviour after Cooling with Different Rates

The influence of the applied cooling rate variation (0.01–5 K/s) from the solution annealing temperature on the transformation behaviour between −80 and +100 °C after additional ageing is shown in Fig. 8. The cooling step including the endothermal transformation from the high-temperature austenite (B2) phase to the low-temperature martensite (B19°) phase is shown in Fig. 8a. During cooling of the sample previously cooled at 0.01 K/s from the solution annealing temperature, at least three distinct peaks can be found. During those peaks, a sequence of transformations occurs depending on the specific ratio of Ni/Ti and the size, amount, and location of precipitates and dislocations [7–9]. It can be seen from the diagram that faster cooling results in a slight shift of the transformation peak 1 to higher temperatures and the disappearance of the transformation peak 2 for cooling rates higher than 0.3 K/s. In addition, the transformation peak 3 significantly shifts to lower temperatures with increasing cooling rate from −18 °C at 0.01 K/s to −38 °C at 3 K/s. Comparing the three fastest investigated cooling rates (1–5 K/s), no differences can be recognised for the transformation behaviour from austenite to martensite. This agrees with the finding that the critical cooling rate is just slightly higher than 1 K/s (Fig. 7).

In Fig. 8b, the subsequent heating from −80 to +100 °C is shown. During heating of the low-temperature martensite (B19°) phase, two distinct exothermal transformation peaks can be seen, resulting in the formation of the high-temperature austenite (B2) phase. Peak 4 dominates the transformation during heating. The transformation peak 5 only shows up as a shoulder at transformation peak 4. At the temperature at which the overall reaction expires and adopts the course of the imaginary baseline, the transformation is complete and the austenite finish temperature Af has been determined. Af has been evaluated and summarised in Fig. 6. For a prior cooling rate of 0.01 K/s from the solution annealing temperature and additional ageing, the Af temperature has been determined to be 55.1 °C. For the next faster cooling of 0.03 K/s, the Af temperature drops to 48.6 °C, staying nearly constant for further increases in the cooling rate. Although the Af temperature is nearly the same, the transformation behaviour differs. The shoulder from peak 5 gets increasingly suppressed and shifted to lower temperatures.
In the same manner, we have evaluated the martensite finish $M_f$ and austenite start $A_s$ temperatures (Fig. 8). Increasing transformation temperatures $M_f$, $A_s$, and $A_f$ with a lower cooling rate (Fig. 6) can be explained by the preceding precipitation from the solid solution during cooling. Transformation temperatures rise with decreasing Ni content of the NiTi phase [6, 26], and the Ni content itself is reduced by the precipitation sequence (Eq. 1). Thus, stronger precipitation during cooling causes increasing austenite/martensite transformation temperatures.

**Microstructures after Cooling with Different Rates**

The metallographic images from optical microscopy of the investigated NiTi alloy are shown in Fig. 9. These samples were continuously cooled from 700 °C for 30 min to room temperature.
temperature with varying cooling rates from 0.001 to 0.03 K/s to promote quench-induced precipitation. No ageing treatment was added. In this state, transformation from austenite to martensite starts significantly below room temperature, i.e. the analysed microstructure consists mainly of austenite. Etching reveals the austenite grains.

Beside the austenitic matrix, obviously a high density of bright polygonal low aspect ratio particles of a few micrometres in diameter exists randomly distributed on all of the images. The size of a few micrometres and the distribution of those particles correlate with non-metallic inclusions like titanium carbides and oxides, which have also been observed in Fig. 2 as well as by [23–25]. Hence, these particles appear independent of the cooling rate.

After the slowest investigated cooling rate of 0.001 K/s, at least two types of quench-induced precipitates are observed. One type appears as elongated, bright-grey precipitates without a specific shape and orientation and a length of about 20 μm. It seems that around these particles, precipitate-free zones can be seen. That is, a large proportion of the alloying elements have diffused into these large precipitates. At higher cooling rates, these precipitates are no longer visible, i.e. their formation during cooling is suppressed. As a second type, many blue coloured needle-shaped precipitates can be seen at 0.001 K/s, ranging in length from up to 20 μm down to a few μm. With a thickness of only a few hundreds of nm, they exhibit a high aspect ratio. These particles appear in distinct orientations. Since the material under study is a binary alloy of Ni and Ti, these precipitates are assumed to come from different stages of the precipitation sequence Ni₃Ti (Eq. 1). However, within the scope of this work, we need to keep the open question of which exact phases are precipitated. Certainly, with increasing cooling rate, these needles become smaller and less in number. After cooling at 0.03 K/s, no quench-induced precipitates can be seen using optical microscopy. As the DSC-signal clearly indicates an exothermal precipitation reaction during cooling at this rate (Fig. 5), further particles are probably formed at the nanoscale, which need further analysis by transmission electron microscopy (TEM) in the future.

Conclusion

In this work, the successful application of a sophisticated, systematic differential scanning calorimetry (DSC) in situ analysis of dissolution and precipitation processes during continuous heating and cooling of a Ni-rich binary NiTi shape memory alloy has been shown for the first time. From the systematic DSC analysis combined with optical metallography, we conclude the following:

- Quench-induced precipitation during cooling from the solution annealing temperature was detected by DSC and confirmed by microstructural analysis. The size and number of precipitates decrease with increasing cooling rate.
- A critical cooling rate for quenching from the solution annealing temperature of about 3 K/s has been determined.
- The impact of existing (quench-induced) precipitates on the transformation temperatures has systematically been analysed.
- Pre-existing precipitates, i.e. nickel rich particles, can significantly influence on the transformation temperatures. In consequence, different heating rates, especially slower ones, can lead to different transformation temperatures under otherwise identical ageing conditions.
- It can be further concluded that quench-induced precipitation (QIP) also strongly influences the transformation behaviour of austenite to martensite and the reverse. Specifically, with increasing suppression of QIP, the martensite to austenite transformation temperature decreases. E.g. after cooling at various rates from solution treatment and additional ageing (500 °C 20 min), the martensite finish temperature during cooling at 0.05 K/s from 100 °C shifts from —21 °C (preceding cooling at 0.01 K/s) to about —41 °C after a preceding cooling at 5 K/s. A continuous heating dissolution (CHD) and a continuous cooling precipitation (CCP) diagram for NiTi have been established. These diagram can be used for the selection of heating and cooling rates in processing NiTi components.
- Future work needs to analyse the nature of quench-induced precipitation on the micro and nanoscale.

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Declarations

Conflict of interest The authors declare that they have no conflict of interest.

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