

Shape Memory Response of Polycrystalline NiTi12.5Hf Alloy: Transformation at Small Scales

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Abstract The transformation behavior of NiTiHf alloys is intriguing. In NiTiHf alloys, the experimental transformation strains have been reported to be considerably lower than theoretical transformation strains. In this study, the transformation strain is established with very careful strain measurements at small scales in isobaric and isothermal experiments. Because of the heterogeneity of strain distributions, the results depend on the sub-region considered. The measured local transformation strain can be as high as 6.0 % in compression which is in very good agreement with theoretical calculations for NiTi12.5Hf. The comprehension of NiTi12.5Hf alloy was furthered upon extensive microstructural characterization including high-resolution electron microscopy, establishing the volume fractions of precipitates and twin type. The volume fraction of precipitates is similar to that of Ni-rich binary NiTi alloys. Meanwhile, the twinning modes in the martensite are compound and Type I twins which were used in the theoretical calculations of transformation strains. This material also generates a high work output and represents a foundation for understanding higher Hf compositions.

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H. Sehitoglu huseyin@illinois.edu **Keywords** Shape memory · Pseudoelasticity · Digital image correlation · Transformation strain · High-resolution electron microscopy

Introduction

NiTi alloys undergo reversible phase transformations with exceptional shape memory and pseudoelastic behaviors which allow their utilization for a wide range of biomedical and industrial applications [1, 2]. The ternary additions (such as Hf, Cu, Fe, Pd, and others) can improve the functionality of shape memory alloys [3, 4] via changing transformation temperatures. However, in most cases, the addition of ternary elements (Cu, Fe, Pd) has reduced the transformation strain magnitudes compared to binary NiTi. Experimental results on NiTi20 Hf [5-8] also exhibit lower transformation strain than binary NiTi alloys [9]. In view of the inferior NiTiHf transformation strains measured at macroscale compared to the theoretical values, the question remains whether better agreement can be achieved if experimental strain measurements were made at smaller length scales. In addition, high Hf compositions are often tested at high temperatures where slip-induced plasticity may curtail the transformation. Studying a lower Hf content such as 12.5 at.% Hf could provide a better assessment of transformation strain change with increasing Hf content. Moreover, precise characterization of the NiTiHf microstructure (precipitate volume fraction and twin modes) is needed. For complete study, the SMA experiments must consider both isobaric temperature cycling (shape memory), and isothermal pseudoelasticity. The strain levels for the two cases may differ because of the ease of plastic flow (or fracture) in pseudoelasticity experiments at higher strains. In this work, we undertake a

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thorough investigation following a multi-scale approach with the aim to provide a better understanding of the transformation strain in NiTi12.5Hf alloy.

The most recent publications in Hf alloyed NiTi have been focused primarily on 20 at.% Hf [8] and recently on 25 at.% Hf compositions [10]. The experimental transformation strains for 20 at.% Hf are less than 4 % in most cases compared to theoretical values at 8 %. To our knowledge, in the case of NiTi12.5Hf composition, such comparisons have not been reported. Experiments on 12.5 at.% Hf are undertaken in the present work, and theoretical strain values are determined. The theoretical transformation strains were established utilizing energy minimization and lattice deformation theories (LDT), and a straightforward comparison between theoretical and measured transformation strains is then presented.

Previous works provided macroscopic strain measurements in NiTiHf over a large domain (mm) measured with an extensometer. The measured transformation strain is thus averaged over a large specimen area containing a large number of grains. These grains are expected to show different transformation strains which are highly orientationdependent in NiTiHf-based alloys. Large strain heterogeneities are thus expected in the polycrystalline alloy, but such heterogeneous nature cannot be captured with classical strain measurement techniques. In this study, we measured transformation strain via digital image correlation (DIC) in order to determine its magnitude at small scales [10]. Local strain measurements via DIC [11] focus only on the grains (or a group of grains) in which transformation develops. The response in this region provides insight into the intrinsic transformation behavior as the surrounding domains remain largely untransformed. In addition to DIC, at smaller length scales, the characterization of the microstructure was achieved by utilizing SEM, TEM, and scanning transmission electron microscopy (STEM) including high-resolution observations to precisely determine the precipitate volume fraction and the type of twin planes.

Because of the importance of precipitation in NiTiHf alloys, we characterize the precipitate volume fraction with three-dimensional microscopy techniques. We also investigate the influence of different aging treatments on the transformation temperatures and transformation strains to achieve the most favorable transformation strain conditions. We focus most of our results on a particular aging treatment and aging temperature (600 °C/100 h) that results in austenite peak temperatures near 90 °C and precipitate size of 200 nm. The precipitate morphologies and twin interfaces were characterized through high-resolution transmission microscopy. Based on the twin-type observed microscopically (Type I-2 and compound twins), the theoretical transformation strains are computed. Importantly, with the addition of Hf, the elevation of transformation strains in compression compared to the stoichiometric NiTi is primarily due to the monoclinic angle increase from 97° for NiTi to 99° for the NiTi12.5Hf case [12]. The observed localized transformation strains in the present study (near 6 % in compression) confirm this finding. Therefore, it is important to use the NiTi12.5Hf as a foundation for studies of Hf-modified NiTi alloys.

Experiments

The Ni₅₁Ti_{36.5}Hf_{12.5} alloy investigated in this work was produced by plasma arc melting. The ingots were sectioned into $4 \text{ mm} \times 4 \text{ mm} \times 8 \text{ mm}$ compression samples by electrical-discharged machine. The samples were homogenized in the vacuum chamber for 4 h at 1000 °C, followed by solution treatment at 900 °C for 1 h and quenching in iced water. The samples were successively aged at various temperatures for different time periods. Thin NiTi12.5Hf foils, 40-50 mg, were cut by a low-speed diamond saw for differential scanning calorimetry (DSC) analysis. The surface of the compression specimens was polished by 600- and 1200-grit SiC paper. A fine speckle pattern was then airbrushed onto the surface of the compression specimens for DIC strain measurements. The isothermal and isobaric compression experiments were both conducted on a MTS servo hydraulic load frame. During isothermal experiments, the compression specimens were deformed in displacement control at a strain rate of approximately 10^{-4} /s. The load frame was controlled by customized LabView software and images were automatically captured during loading and unloading by an IMB-202FT CCD camera and Navitar optical lens providing a final image resolution of 2 µm/pixel. The temperature was controlled by Lepel induction generator through induction coils. The grips were cooled by means of copper wires connected with a liquid nitrogen tank allowing sub-zero cooling. A Raytek infrared sensor was used to monitor the temperature of the specimen. In order to explore the shape memory properties in the isothermal condition, the sample was heated above the austenite finish temperature (A_f) to ensure complete transformation into austenite state. Subsequently, the temperature was gradually decreased to the test temperature. In isobaric test, images were manually captured during heating and cooling with a temperature step of 5 °C. During the isobaric test for the long-aged heat treatment, the temperature was cycled between 0 and 140 °C. The axial strain fields obtained from these experiments through DIC were used to characterize stress-strain and strain-temperature behaviors of the material.

Specimens for TEM and SEM were electro-polished by the twin-jet method in an electrolyte solution of HNO₃ and methanol (1:5, v/v) at about 255 K. The SEM observations were conducted on a Carl Zeiss Ultra 55 system. The focused ion beam slicing and SEM observation were repeatedly performed on a FEI Scios DualBeam system. A FIB slicing interval was set to 5 nm. The successive tomographic images were analyzed with the software Amira (Maxnet, Inc.). Selected-area electron diffraction experiments and TEM observations were carried out on a JEM-2000EX system operated at 200 kV. High-angle annular dark-field (HAADF)-STEM observations were performed on a JEM-ARM200F system (Cs-corrected 200 kV STEM). The electron probe size and current were 0.10 nm and 20 pA, respectively. For the HAADF-STEM imaging, the convergence semi angle was fixed to 22 mrad. The collection semi angle was set between 90 and 170 mrad.



Fig. 1 a DSC curves showing the influence of the most significant heat treatments on the phase transformation temperatures, and **b** effect of aging time on austenite $(A_{\rm PT})$ and martensite peak temperatures $(M_{\rm PT})$

Experimental Results

The most important DSC results are shown in Fig. 1a. Compared to the homogenized case, the transformation temperatures decreased upon solution treatment and subsequent aging at 637 °C for 1 h. In addition, for the case of 605 °C/4 h and 665 °C/4 h heat treatments, the transformation temperatures are still similar to those of the homogenized case. Increment of the transformation temperature can be observed in the long-time aging cases, i.e., 637 °C/11 h and 637 °C/100 h. In order to investigate the shape memory behavior at the highest temperatures possible, long aging times were chosen (the gray curve in Fig. 1a). In Fig. 1b, the influence of the aging time on the $A_{\rm PT}$ and $M_{\rm PT}$ temperatures is presented for a fixed aging temperature 637 °C. Both $A_{\rm PT}$ and $M_{\rm PT}$ increase from 1 to 15 h of aging. $M_{\rm PT}$ increases faster than $A_{\rm PT}$ resulting in a big drop of hysteresis, $A_{\rm PT}$ - $M_{\rm PT}$, at the lower aging time range. The hysteresis, $A_{\rm PT}$ - $M_{\rm PT}$, shows a minimum at 10 h aging and increases slightly with further aging time.

In Fig. 2, precipitation in the material matrix for the case of 637 °C/100 h was captured through SEM. Based on the volume fraction of particle in grains 1 and 2, the overall volume fraction was calculated to be approximately 20 %. The cross-section shape of the precipitate is oblate spindle like as shown in (b) and (c), which is also similar to the one reported for Ti-rich NiTiHf after 150 h of aging [13]. However, the overall shape of the precipitate can be more complicated as shown in Fig. 3. The size of the precipitate is measured as approximately 200 nm. These facts are also confirmed by three-dimensional visualization of the

Particle Volume Fraction 0.20



Fig. 2 a Secondary electron image showing precipitation distribution in Ni₅₁Ti_{36.5}Hf_{12.5} aged at 637 °C/100 h. **b** and **c** Enlarged images captured from the *framed areas* in **a**

precipitate structure (movie is included as a supplement) in Fig. 3.

Isobaric experiments (Fig. 4) were conducted under different applied compressive loads. In Fig. 4, we also report the strain fields at the minimum temperature of each thermal cycle under the specific compressive stress. The strain fields were obtained using the reference image captured at the maximum temperature and zero stress. The



Fig. 3 Three-dimensional visualization of the precipitate structure (movie is included as a supplement)

solid line was determined averaging the axial strain fields over all DIC region (4 mm \times 3 mm). It is worth pointing out the heterogeneous nature of the strain fields where low and high strains are exhibited at small scales. The heterogeneity of the strain distribution is mainly due to the fact that finite volumes retain their austenitic structure. This observation is further confirmed with X-ray diffraction of the samples based on the B2 peaks. In order to quantify the maximum transformation strain in the straintemperature plots, we also include the curve representing the local temperature-strain behavior of the region that displays the largest strain localization (dashed line). The area which was used to collect localized data is shown in the strain contour plots by a black box. Besides, the red line and blue line represent heating and cooling, respectively. From 0 to 500 MPa applied stress, negligible irreversibility was measured either for the averaged strains over the DIC region or for the local. The maximum local transformation strain was obtained to be 5.5 % at 500 MPa. The hysteresis was found to be approximately 40 °C. In addition, the transformation temperatures will also tend to increase with increasing of applied stress. According to Fig. 4, when the applied stress is 0 MPa, M_f , M_s , A_s , and A_f are 20, 35, 50, and 70 °C, respectively. All the temperatures increase by 15 °C when a 500 MPa stress is applied. The trend matches with the



Fig. 4 Isobaric tests at various compressive stresses for the Ni_{51} -Ti_{36.5}Hf_{12.5} alloys aged at 637 °C/100 h. The corresponding DIC results at each stress level show the strain *contour* at the minimum

temperature of the thermal cycle. The *boxes* displayed in the DIC strain fields indicate the region used to calculate the local strains and the corresponding strain-temperature curves (*dashed line*)



Fig. 5 Stress and strain relationship at various temperatures in which a shape memory effect (SME) dominates and b pseudoelasticity (PE) and plasticity (PLS) dominate. The local in situ strain measurement (*blue curves*) was extracted from the strain fields delimitated by the *boxes* displayed in the DIC images



Fig. 6 Flow stress as a function of temperature when the Ni₅₁Ti_{36.5-} Hf_{12.5} is heat treated at 637 °C/100 h and 637 °C/92 h. The *inset* shows NiTi and NiTiCu₉ for comparison

other experimental results published in the literature [7]. However, the thermal hysteresis stays relatively stable at a level of 40 $^{\circ}$ C.





Fig. 7 Pseudoelastic behaviors of the $Ni_{51}Ti_{36.5}Hf_{12.5}$ alloy at various temperatures after the 637 °C/4 h aging treatment. The local in situ DIC strain measurement is shown in *blue curve*



Fig. 8 Pseudoelastic behavior for three different aging treatments for the Ni₅₁Ti_{36.5}Hf_{12.5} alloy. The local in situ DIC strain measurement is shown in *blue curve*. This figure is significant because it points the high work output in these alloys near 50 J/cm³



Fig. 9 The last load step at 55 °C for the $Ni_{51}Ti_{36.5}Hf_{12.5}$ alloys aged at 637 °C/4 h with DIC results showing strain *contours* at both maximum and minimum loads under compression. The local in situ DIC strain measurement is shown in *blue curve*

In Fig. 5, we report the stress-strain curves obtained during isothermal experiments in the temperature interval between pure martensite deformation and pure austenite



Fig. 10 a Three-dimensional illustration of the strain *contour* at the peak stress, $\sigma = -1200$ MPa, for 637 °C/4 h aging, and b strain as a function of position in X direction at the maximum and minimum load in A

deformation (slip). To gain more insight into strain heterogeneity, the black curve indicates the averaged strain over a large area on the sample (4 mm \times 3 mm), while the blue curve indicates the local stress-strain behavior. It is important to note that transformation is localized at small scales. The red lines underneath the stress-strain curves indicate the recoverability after the thermal cycle following heating and cooling to the test temperature at zero load. At the martensite start temperature, 40 °C, the yield stress displays a minimum. The corresponding residual local strain (3 %) was then fully recovered after the heating/cooling thermal cycle. At temperatures where stressinduced martensite can be obtained (85-100 °C), good pseudoelasticity is found and shown in Fig. 5b. Based on these results, the maximum work output can be estimated using a stress level of 1200 MPa and a strain level of 4.4 % as 50 J/cm³. This magnitude exceeds the work output levels observed for other NiTiHf alloys.

The relationship between 0.1 % offset flow stress and temperature is plotted in Fig. 6. A linear relationship with a slope of 14.5 MPa/°C between flow strength and temperature is shown. The stress corresponding to the M_d temperature is nearly 1200 MPa which is substantial. The transformation temperatures are marked in the figure. In the inset, similar relationships for NiTi [14] and NiTiCu₉ [15] are presented to provide an insight into the comparison of flow stress among the three materials.

The pseudoelastic behavior for shorter aging treatment times, 637 °C/4 h and 637 °C/2 h, was also investigated. The stress–strain characterizations obtained through isothermal tests are shown in Fig. 7. As shown, when a 4 h aging treatment is implemented, near-perfect pseudoelastic

behavior was obtained at relatively lower temperatures. Perfect pseudoelastic behavior is captured at both 40 and 55 °C for 637 °C/4 h heat treatment. At 70 °C, irrecoverable processes (slip of austenite and/or slip of transformed martensite) occur and small residual strains were measured upon unloading. The shape memory effect is obtained at room temperature for this heat treatment. It is also interesting to observe that pseudoelastic behavior is captured at 40 °C which is below austenite finish temperature, approximately 60 °C.

In Fig. 8, the pseudoelastic behaviors for different aging treatments at different load steps were reported and compared. The temperatures selected for the tests were all at $A_{\rm f}$ -5 °C. In all three cases, pseudoelasticity was captured at a relatively high stress level above 800 MPa. The overall strains recovered at a same stress level are similar for all three cases. When a compressive stress of 1200 MPa is applied, maximum total strain in the order of 5 and 5.5 % are recovered locally upon unloading for the 637 °C/2 h and 637 °C/4 h aging, respectively. For these aging treatments, the transformation strain is determined to be 3.5 and 4.2 % as shown in Fig. 8. No apparent irreversibility was observed as well. On the other hand, at the similar stress level, obvious plastic strain, 0.5 %, is accumulated locally for the 637 °C/100 h aging. The plastic strain accumulation due to long-time aging also verifies the fact that over-aging would deteriorate the pseudoelasticity.

The strain contour plots captured at the maximum and minimum load are shown in Fig. 9 for the 637 °C/4 h aging treatment. The amount of total strain in the order of 4.4 % is fully recovered, while 3.3 % of which is due to phase

Fig. 11 The stereographic triangle of the **a** CVP, **b** LDT strains, **c** bright field image, **d** corresponding electron diffraction pattern showing $(011)_{B19'}$ Type I twin and four variants of precipitate (*P*), **e** HAADF-STEM image, and **f** its inverse fast Fourier transformed image showing mirror symmetry of lattice *planes* with respect to $(011)_{B19'}$



transformation. In addition to the pseudoelastic behavior, the hysteresis is found to be the lowest for the 637 $^{\circ}C/4$ h case among all. In the DIC images that are shown in Figs. 5, 7, and 9, the transformation strain distribution in this material is not uniform. In order to appreciate such phenomenon

explicitly, a three-dimensional illustration of the axial strain contour is reconstructed into Fig. 10a. At the maximum and minimum load, an averaged strain over the peak area A is plotted as a function of positions in X direction in Fig. 10b. The local strains pointwise reach 6 % levels.



Fig. 13 The measured polycrystalline texture in NiTi12.5Hf. The measured texture favors (011) aligned grains in the loading direction (Z axis)

Calculations of Transformation Strains and High-Resolution Electron Microscopy

To gain further insight into the NiTi12.5Hf transformation behavior, the theoretical transformation strains were calculated using the following lattice constants. For austenite $a_0 = 0.3075$ nm, and for the monoclinic martensite a = 0.2970 nm, b = 0.4060 nm, c = 0.4770 nm, and monoclinic angle = 98.9° [12]. The correspondence variant pair (CVP) formation strain (for Type I-2 twinning) and the LDT strains were established using similar procedures as outlined in [9]. Figure 11a, b provides the transformation strains for CVP and LDT, respectively. The results show maximum transformation strains of 7 %. Figure 11c shows the bright field image and Fig. 11d illustrates its corresponding electron diffraction pattern in the Ni₅₁₋ Ti_{36.5}Hf_{12.5} aged at 637 °C for 100 h. The pattern consists of two sets of $[100]_{B19'}$ reflections with the $(011)_{B19'}$ Type I twin relation. Since the sharp line contrast between alternate platelets in (c) is parallel to the trace of $(011)_{B19'}$, the platelets are in $(011)_{B19'}$ Type I twin relation. This twin relation is clearly confirmed in HAADF-STEM and its inverse fast Fourier transformed images in (e) and (f), respectively. From the electron diffraction experiments, the structure of precipitate is the same as that reported by Santamarta et al. [16]. There are four sets of precipitate marked as 1, 2, 3, and 4 around $0\bar{1}0_M$ and $00\bar{1}_T$ spots in the enlarged pattern inserted in the upper right corner in (d) [16]. Figure 12a, b indicates that there are three sets of B19' martensite plate consisting of $(001)_{B19'}$ compound twins marked as A, B, and C. There are also $1/3 \ 110^*_{B2}$ type reflections derived from precipitate marked as P in (b). The interfaces between three martensite plates are not straight and sharp in comparison to those in Ni_{50.7}Ti_{29.3}Hf₂₀ alloy furnace cooled from 1000 °C.

The X-ray diffraction results are shown in Fig. 13 where the Z direction represents the loading axis of the specimen. The results show a preferential texture with a higher volume fraction of grains in $\langle 011 \rangle$ direction. The local transformation strain measured through experiment (~6% considering the uncertainty of the strain measurements, as reported in Table 1) is in a good agreement with theoretical Table 1 Comparison of experimental and theoretical transformation strains for NiTi12.5Hf

Material (polycrystalline)	Experimental compressive transformation strain	Theoretical compressive transformation strain
NiTi12.5Hf	5.5 ± 0.5 % (Local DIC area)	6.8 % (Local, $\langle 011 \rangle$ pole)

The results show that the local strain measurements and theory are close to each other while the entire DIC area gives a smaller average strain compared to theory (CVP strain)



Fig. 14 The variation of transformation strain in [011] direction as a function of Hf content in NiTiHf alloys. The result of the present material 12.5 at.%. Hf is shown with a *blue marker*. Note that lattice constants from several investigators are utilized to calculate the strains. The 6.25 and 12.5 at.% Hf lattice constants are obtained from DFT calculations

calculation of transformation strain for the $\langle 011 \rangle$ pole, which is 6.8 %. Theoretical values are still difficult to be matched as the DIC strain measurements provided in this study do not reach the nano-scales. Such resolution would be required in order to avoid the effect of the non-transforming precipitates. As stated earlier, the NiTiHf alloys attain higher transformation strains based on theoretical calculations compared to the binary NiTi case. This is illustrated in Fig. 14 where the transformation strains are shown as a function of increasing Hf content. The results for [011] compression are computed because this is the pole that produces higher strains in compression compared to others. In Fig. 14, the lattice constants (cubic austenite and monoclinic martensite crystals) are either obtained from X-ray diffraction in previous studies or from density functional theory [6-9, 12, 17]. We note that the 12.5 at.% Hf represents a median strain in the range of alloys in Fig. 14. Three transformation strains, CVP strain and CVP + detwin strain, LDT theory strains, have been computed. The results shown in Fig. 14 represent the transformation strain calculated from the LDT theory. The differences between these three strain measures are not high in compression [9]. The straight line in Fig. 14 is drawn to aid the eye.

Summary and Discussion of Results

The work explored the underpinnings of NiTi12.5Hf transformation strains with experiments and theory. Precise local measurements are needed during the experiments to gain a true measurement of the transformation strains. Because of large domains of untransformed material, the overall transformation strains could be lower than theory, while the local levels are in agreement with theory. The maximum local measured strain levels reached 7 % (Fig. 10) which exceeds the level in other NiTiHf studies. The results also point to the importance of characterization of the microstructure establishing the type of twinning of the martensite and the volume fraction of precipitates. This knowledge represents an important step to make comparisons between theory and experiment. Since the precipitates cannot transform, the measured strain levels can be lower than expected.

The shape memory and pseudoelasticity response of the new shape memory alloy $Ni_{51}Ti_{36.5}Hf_{12.5}$ were chosen because it permitted studies of shape memory and pseudoelasticity without premature fracture. Hence, the flow stress behavior (Fig. 4) near the M_d temperatures can be measured readily and compared to binary NiTi and other NiTiX alloys. As pointed earlier, the Hf provides a stronger strengthening element compared to Fe, Cu, and Pd [12]. The flow stress levels were near 1200 MPa for the NiTi12.5Hf alloy and using a macroscopic strain of 3.5 % results in a high work output (near 50 J/cm³) which is likely the highest among transforming alloys. While the NiTi alloys with high Ni contents [18] also exhibit high flow stresses, the transformation strains are less than 3.5 % in those cases.

During studies on shape memory alloys, it is important to seek treatments to achieve the optimum levels of transformation strains, transformation temperatures, and temperature hysteresis. We note that transformation temperatures of this material were investigated with various aging treatments (637 °C) ranging from 1 to 100 h. We observe that the transformation temperatures rapidly increase with aging time from 1 to 10 h, reaching saturation at longer aging times beyond 10 h (see Fig. 1a, b for details) with austenite peak temperatures near 100 °C.

For the heat treatment at 637 °C for 100 h, the shape memory property was investigated during isobaric experiments at different stress levels. The material possessed small residual strain up to -500 MPa. In Fig. 4, the maximum strain was in the order of 7 % locally and later fully recovered upon heating. This is comparable to the strain level predicted based on the LDT theory utilizing the lattice constants of austenite and martensite phases. It was clear from the DIC results that the grains which yield the highest transformation strain do not extend to the entire specimen. The shape memory effect and pseudoelasticity were also investigated via isothermal compression tests at different temperatures as shown in Fig. 5. Pseudoelasticity extended to 115 °C in this class of alloys. As the temperature increases, excellent pseudoelastic behavior was found at 85 and 100 °C. The plastic irreversible strains developed beyond 115 °C. Similarly, further efforts are required to capture sub-grain deformation which would enable strain measurements at higher resolution and potentially match with theoretical transformation strains levels in other NiTiHf alloys.

As shown in Fig. 10, the strain distribution is inhomogeneous with maximum localized strains exceeding 6 %. Higher transformation strains are favored near the [011] pole with much lower strain near [001] with [111] being the intermediate (Fig. 11a, b). Measurements with X-ray diffraction on our samples confirm the preference for grains in (011) direction. It is important to note that for the case of increasing hafnium, the monoclinic angle increases (from 96.8° for NiTi to 98.9° for NiTi12.5Hf) which in turn increases the transformation strains. In general, the high compressive strain values relative to the binary NiTi could be an important advantage of the high Hf-rich alloys in addition to the elevated temperature capability. We note, however, the results point to a large reduction in strains at the macroscale, i.e., polycrystalline experimental strains are near 3.5 %. This is attributed to the presence of untransforming domains.

Comparing NiTiHf alloys to other ternary shape memory alloys, the strengthening associated with Hf is significant. In fact, the flow stress rises with increasing Hf content and reaches a saturation at higher Hf contents [12] (see Fig. 7). Therefore, the flow stress and transformation strain combination appear to be very attractive for lower Hf contents such as 12.5 at.% Hf though the choice of Hf could depend on the application. The predicted transformation strains for NiTiHf alloys, are higher than NiTiCu [18] and NiTiFe alloys, and also with higher temperature capability as discussed above.

We provided a comparison of the transformation strain calculations for compression in Fig. 14 for the $\langle 011 \rangle$ direction. The results show unequivocally the elevation of the transformation strains with increasing Hf content. Such an increase can be exploited to design potentially NiTiHf compositions with higher transformation strain levels by modifications in crystallographic texture and Hf contents. One of the issues is clearly the heterogeneity in the transformation at the macroscale which needs to be further explored in other Hf alloys.

Overall, the work underscores the need for a multifaceted approach combining (i) precise determination of the values of lattice constants with diffraction methods or electronic structure calculations, (ii) high-resolution microcopy to identify twin types and volume fraction of precipitates, (iii) measurements of local strains in a heterogeneous environment, and (iv) phenomenological martensite strain calculations to gain a stronger understanding of the behavior of NiTiHf alloys. The results point to the need for a better understanding of transformation strain progression during transition from small to large scales in future studies and to ascertain the role of grain boundaries on the advance of transformation fronts in NiTiHf class of alloys.

Conclusions

The work demonstrated the following conditions:

- (1) In NiTi12.5Hf alloy, local strain measurements with DIC result in high transformation strains and provide an intrinsic measure of transformation since domains of transformed and untransformed material co-exist in NiTiHf alloys. At the local level, the strains can reach 6 % in experiments which agrees with transformation strain predictions near the $\langle 011 \rangle$ pole (~ 6.7 %). When larger volumes are considered, the measured transformation strains fall to 3.5 %.
- (2) In the NiTi12.5Hf alloy, the transformation stress levels approached 1200 MPa in compression, and with macroscopic strain levels of 3.5 %, the work output of such alloys is near 50 J/cm³. This level of work output exceeds the higher Hf alloys where the stresses are similar while the experimental strains are much smaller. Compared to binary NiTi alloys, the work output is also higher in compression.
- (3) A thorough microstructural characterization utilizing (SEM), (TEM), (STEM), and X-ray diffraction was undertaken to gain a better understanding of the NiTi12.5Hf. The X-ray diffraction studies pinpointed the strong preference for $\langle 011 \rangle$ grains supporting the high strain measurements locally. The microscopy observations provided quantitative insight into the precipitate volume fraction and the twinning type in this alloy.

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