Influence of Silver Addition on Structure, Martensite Transformations and Mechanical Properties of TiNi–Ag Alloy Wires for Biomedical Application

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Abstract: The microstructural and functional behavior of TiNi-based wires with a silver content of 0–1.5 at.% was evaluated. The concentration range for Ag doping determined for the TiNi wires with potential for the medical industry was 0–0.2 at.%. Microstructure analysis of TiNi wires with different silver contents at room temperature indicated a multiphase structural state. Various internal structures with tangled grain boundaries were formed by intense plastic deformation. The nanocrystalline structure and phase state of wire with the minimum silver content (0.1 at.% Ag) provide full shape recovery, the greatest reversible strain, and optimal strength and ductility. TiNi ingots with a high Ag content (0.5–1.5 at.%) cracked under minimum load due to excess silver that crystallized along the grain boundaries and broke cohesion bonds between the TiNi grains.

Keywords: shape memory alloy; silver; doping; mechanical properties; nanostructured material; wire

1. Introduction

The development and study of new TiNi-based materials is a promising direction in the field of medical materials science. The special properties of the alloys are shape-memory and superelasticity due to reversible austenite-martensite transformations caused by changes in external conditions such as load and temperature [1–4]. There are differences between thermal and stress-induced martensite transformations that affect the magnitude of the transformation strain [5,6].

TiNi-based wire is widely used in the manufacture of both temporary and long-functioning surgical implants because of bio-inertness and ability to undergo viscoelastic deformation with tissues, surviving millions of deformation cycles without failure [7,8]. In order to increase the survival rate of TiNi implants, an additional antibacterial effect is required. This effect can be achieved with silver doping [9,10]. At the same time, it is important to obtain a technological alloy that can be used to produce implants with high functional properties including fully reversible deformation, high strength, and ductility.

Nowadays, doping and coating of coarse-grained TiNi alloys with Ag with a concentration of 0.5 to 9 at.% without subsequent thermomechanical treatments is a common practice worldwide [11–15]. In terms of using TiNi–Ag materials in the medical industry, alloying TiNi with Ag provides the alloy with new attributes suitable for biomedical applications, improving its cytocompatibility and antibacterial capacity [11], while increasing yield and tensile strengths [16,17]. TiNi composite surface films doped with silver (4–10 at.%) enhance mechanical strength, biocompatibility, and corrosion.
resistance to implants [18–20]. However, as the silver concentration exceeds 5 at.%, the antibacterial properties of the alloy deteriorate. Therefore, doping with lower silver concentration is of particular interest. The shape memory effects of Ag-doped TiNi were studied at a concentration of 1.4 at.% silver [12], and martensite transformations as well as microhardness were mainly considered in the range of 0.6–1.9 at.% Ag [21–24]. However, the issue of how lower silver content (< 0.5 at.%) could affect the functional characteristics and deformability of TiNi–Ag wires has not yet been studied thoroughly.

The above-mentioned overview of the effect of high silver doping on the structure and properties of TiNi indicates the relevance of this issue for biomedical applications. During implant manufacturing, alloys are subjected to numerous thermo-deformation effects, so deformation properties are of high importance. At the first stage, the structure, shape memory, as well as mechanical and biocompatibility properties of TiNi-based alloys with 0–1.5 at.% Ag were studied before thermo-mechanical treatments [25,26]. Interpretation of the resulting dependencies between phase composition, microstructure, functional and mechanical properties served as the scientific basis for creating new implants from TiNi wires with enhanced performance.

This study is the second stage of research aimed at creating biocompatible TiNi alloys with high functional properties and an additional antibacterial effect. The purpose of the current study is to determine the range of silver content for the production of TiNi–Ag wires obtained by intense plastic deformation and to study their structure, phase composition, shape memory effect and mechanical properties.

2. Materials and Methods

TiNi–Ag alloys were melted in an induction furnace according to the following doping scheme: Ti50Ni50–XAg (X = 0, 0.1, 0.2, 0.5, 1, and 1.5 at.%). The combined method of intensive plastic deformation, consisting of multiple processes of cold rolling and intermediate annealing (T = 400–450 °C), was used to produce wire with a diameter of 1 mm. Three main production stages were used: i) rolling of ingots (8 cycles) up to a bar 80 mm thick; ii) rotary-forging of round bars up to a 3.5 mm wire (7 cycles); iii) hot wire drawing up to 1 mm (25 cycles). The following requirements of cold deformation processes were fulfilled: low degree of compression per cycle, and intermediate annealing to reduce work-hardening effects and increase metal ductility. Intermediate annealing temperatures were selected in such a way that the recrystallization of grains occurred, but there was no increase in grain size.

Ingots with a high Ag content (0.5–1.5 at.%) cracked under minimum compression using a rolling mill (Figure 1).

![Figure 1. Rolling and drawing of TiNi–Ag with different silver content.](image-url)
a screening effect of the surface oxide layer on the matrix. Therefore, a special method of sample preparation was developed for the examination of the matrix of 1 mm thick wire. In order to obtain a flat surface, 20 wire segments were connected with a thin TiNi thread into a dense “bundle” and filled with epoxy resin at the base. After that, the end surface of the “bundle” was mirror polished. The shape memory effect parameters were studied in tension at a constant load of 2 kg. Transmission electron microscopy (TEM) imaging of thin foils were performed using a JEM-2100 transmission electron microscope (JEOL, Tokyo, Japan) at an accelerated voltage of 200 kV. The mechanical properties were studied using an Instron testing machine under tensile strain conditions.

3. Results and Discussion

Ingots with a high Ag content (0.5–1.5 at.%) cracked under minimum compression, as depicted in Figure 1. The deformation process is significantly affected by the grain/grain boundary interface interaction, since grain boundaries have a chemical composition and characteristics that differ from those of the grains themselves [27,28]. The grain boundaries have a significant impact on the deformation conditions, since their chemical composition and properties differ from those of the grains. There are zones at the grain boundaries that prevent deformation. The movement of atoms in the grain boundaries required for intergranular deformation is complicated by the presence of insoluble Ag phases. Silver phases in TiNi alloys with an Ag concentration of more than 0.5. at.% were formed as a result of their limited solubility in TiNi [29]. The liquation of excess silver along the grain boundaries in the alloys broke cohesion bonds between TiNi grains. Thus, the presence of phases along the grain boundaries changes the deformation mechanism; as deformation is developing, cavities arise in the form of microcracks and micropores, which lead to cracks nucleating at specific points and propagating across the material.

XRD analysis showed that TiNi wires with different silver concentrations at room temperature are characterized by a multiphase structural state (Figure 2). The purity of the silver used as a dopant was 99.9%, as confirmed by a certificate. However, as is evident from the presented XRD patterns, no traces of Ag-related phases were revealed. The findings on elemental mapping reported in our previous work have also proved no aggregate of metallic Ag0 or oxides AgxOy [25]. The main structural component of the wires is intermetallic TiNi in B2 and R modifications. This is confirmed by the angular distribution and high intensity of the main reflections of these phases. In the angular interval, 2θ = 42–43°, a pronounced splitting of the main reflection of the B2 phase with a partial overlap of {110}B2 and {303}R peaks was found (inset in Figure 2a). These results correlate with the electrical resistivity ρ(T) data in the same series of samples [25].

The sizes of the coherent scattering regions (CSR) of the B2 phase in the studied wires do not exceed 30 nm, which indicates a fine-crystalline structural state after multiple mechanical and thermal effects (Figure 2d). Qualitative XRD analysis revealed the presence of structural reflections from TiNi phases with a stoichiometry of Ti2Ni, Ti3Ni4, and TiNi3 with a total volume fraction not exceeding 15 vol.%, which were distributed differently depending on the composition. The occurrence of Ni-rich phases is stemmed from the segregation of Ti to the alloy surface during drawing, with subsequent formation of an oxide layer through intermediate annealing [30,31]. No phases with Ag were detected in the XRD diffraction pattern of wire samples. A small amount of silver is not sufficient for the formation of individual intermetallic compounds; therefore, the silver completely dissolves in the matrix replacing titanium in its sublattice, as indicated by a decrease in the lattice parameter (Figure 2e).

Silver dissolves in the TiNiB2 phase to a limited extent, up to 0.26 at.% [11,12]. The addition of 0.1 at.% Ag causes the occurrence of reflections of the B19’ phase. An increase in silver content in combination with intense plastic deformation leads to an increase in the preferential orientation of the crystal lattice of B2-phase grains in the (200) direction from 17 to 35%.

An increase in the concentration of Ag in TiNi solid solution decreases the B2 lattice parameter, thereby changing interatomic interaction forces and increasing internal elastic stresses Δd/d from $1.2 \times 10^{-3}$ to $12.45 \times 10^{-3}$ (Figure 2f), provoking a $B2 \rightarrow R \rightarrow B19'$ phase transformation. The volume
fraction of the peritectic Ti$_2$Ni phase decreases from 7 to 4 vol.%, and the amount of the metastable phase Ti$_3$Ni$_4$ increases from 5 to 8 vol.%.

Figure 2. XRD patterns of TiNi wires with silver concentrations of 0 (a), 0.1 (b), and 0.2 at.% (c); concentration dependences in the parent phase on: coherent scattering regions (d), internal stresses (e), lattice parameter (f), phase volume fraction (g).

Qualitative XRD analysis of the diffraction pattern of the wire with 0.2 at.% Ag showed a high content of B19' martensite and Ni-rich fine phase Ti$_3$Ni$_4$, and structural lines from the secondary Ti$_2$Ni and Ti$_3$Ni$_4$ phases that were less intense than those of the wire with 0.1 at.% Ag. Assessment of the volume fractions of phases showed that when the silver concentration increases, the volume fraction of B2 austenite and secondary phase Ti$_2$Ni decreases, while that of Ti$_3$Ni$_4$+TiNi$_3$ increases (Figure 2g).

The average grain size and microstructural features evident from Figure 3 indicate that in alloys with different silver contents, the intense plastic deformation formed various internal structures where the average grain size changed nonlinearly. The small addition of Ag has an ambiguous effect on the processes of relaxation of internal stresses and dynamic recrystallization, variously inhibiting or inducing them. Figure 3 shows that the grain size and the number of deformation defects at the boundaries and in the grain volumes are different, and phases of different sizes are formed.

The TiNi wire without Ag is characterized by a fully crystalline ultrafine-grained structure with high dislocation density and an average size of recrystallized grains of 500 ± 50 nm (Figures 3 and 4). In Figure 4a–c inhomogeneous contrast of the extinction contour in the subgrains indicates the presence of a substructure in which the curvature of the crystal lattice, level of stresses, and density of dislocations is high. The lamella TEM image with the corresponding electron diffraction patterns (EDP) enabled to identify of the TiNi$_{B2}$ matrix, white elongated Ti$_2$Ni inclusions 50–150 nm in size along the grain boundaries, and fine Ni-rich Ti$_3$Ni$_4$ precipitates 20–60 nm in size in individual areas of the matrix.
In the TiNi wire with 0.1 at.% Ag, a more homogeneous nanocrystalline structure was formed without dislocation defects and with an average grain size of 100 ± 50 nm. No dislocation defects were detected in the grains because the small grain sizes prevent the development of dislocation slip. There are areas with inhomogeneous contrast of displacement bands types because of intergranular stresses near the boundaries, caused by the grain orientation difference on the boundary (Figure 4d). EDP identification from different areas of the matrix showed the presence of parent B2, martensite R and B19’ phases, Ti2Ni phase with sizes of 10–50 nm, and Ti3Ni4 and Ti2Ni3 phases (10–30 nm). Finely dispersed Ti3Ni4 precipitates are uniformly dispersed throughout the volume (Figure 4e). The total number of Ti2Ni particles along with the TiNi grain boundaries was half of the precursor TiNi wire, which is in good agreement with the XRD data.

In the TiNi wire doped with 0.2 at.% Ag, an ultrafine-grained structure with dislocation defects similar to that of the TiNi wire (Figure 4f–h) and average grain size of 250–300 nm (Figure 3c) was observed. Identification of diffraction patterns from individual wire areas confirms the presence of the parent B2 and R phases, and secondary Ti2Ni and Ti3Ni4 ones. No diffraction reflections from martensite B19’ were found in the investigated lamella, although the volume fraction of this phase in the XRD spectrum attained 18 vol.%. Figure 3e–g indicates the EDP for typical B2, Ti2Ni, Ti2Ni3, and Ti3Ni4 phases persist in the wires studied.

The formation of ultrafine-grained and nanocrystalline structures is a result of thermomechanical treatment by rolling and drawing with intermediate annealing [7,32] during which a number of excess equilibrium TiNi3 and metastable Ti2Ni3 and Ti3Ni4 phases are precipitated [33–35]. The structures formed have a positive effect on the resistance of the material to plastic deformation.

Figure 3. Transmission electron microscopic (TEM) images of TiNi wires with different Ag concentrations: 0 at.% (a), 0.1 at.% (b), 0.2 at.% (c); concentration dependence of the average grain size (d); TEM image of the parent and Ti2Ni phase, Ti3Ni4, Ti2Ni3 precipitations with corresponding electron diffraction patterns (EDP) presenting in the studied wires (e–g).
Ni-rich secondary phases, in which energy dissipation occurs during the movement of martensite, increase in the 0–0.2 at.% range. The thermal hysteresis amount of energy dissipation during the phase transformation, increases from 55 to 68 °C with a silver concentration increase in the TiNi solid solution and the presence of a high density of dislocations as a result of intense plastic deformation.

Figure 4 shows the features of the shape memory effect (SME) for all wire samples; and the measured SME parameters are given in Table 1. In the cooling-heating cycle under a constant tensile load, all wires are characterized by high reversible deformation values. However, in the wire with 0.2 at.% Ag, the defect structure of the B2 phase has suppressed the martensite transformation due to an increase in the Ag concentration in the TiNi solid solution and the presence of a high density of dislocations as a result of intense plastic deformation.

The onset temperature of direct martensite transformation (MT) insignificantly depends on the silver concentration and remains stable at 38 ± 2 °C. This means that the crystal lattice of the B2 phase is stable to martensite transformations at the macro level under load despite a change in silver concentration. It is known that doping of TiNi alloys with a third element leads to a sharp decrease in the MT temperature or complete suppression, especially under conditions of intense plastic deformation. However, noble elements (palladium, platinum etc.) at low concentrations form solid solutions based on the B2 structure and have contrary effects on the stability of B2 austenite [36], as in our case with silver [17]. The thermal hysteresis \( \Delta T \), which characterizes the amount of energy dissipation during the phase transformation, increases from 55 to 68 °C with a silver concentration increase in the 0–0.2 at.% range.

The main contribution to the width of thermal hysteresis is made by incoherent particles of the Ni-rich secondary phases, in which energy dissipation occurs during the movement of martensite boundaries in shape recovery.
In wires with 0.1 and 0.2 at.% Ag, during MT under loading with decreasing temperature, leads to the formation of dislocation defects during the movement of the austenite-martensite (A→M) interfaces. Then, upon heating during reverse MT, these defects impede the movement of the M→A interface. This phenomenon leads to a broadening of the temperature range of the reverse MT under load, as also reported in [37,38].

The characteristic temperatures $M'_f$, $A'_s$, and $A'_f$ decreased, as residual strain occurs in TiNi wires with 0.2 at.% Ag. The residual strain is caused both by the accumulation of the plastic strain component and by defects that arise during intense plastic deformation. The plastic strain component is the main mechanism for the dissipation of elastic energy. These phenomena significantly affect the movement of interfaces and lead to irreversible effects. The residual strain appears because of the irreversible plastic deformation caused by the high content of finely dispersed Ti$_3$Ni$_4$ precipitates, which do not undergo martensite transformations. The slope of the thermal hysteresis loop is related to the high heterogeneity of the structure and the density of defects in the initial TiNi wire and in the TiNi wire with 0.2 at.% Ag, and is in good agreement with TEM studies.

![Figure 5. Experimental curves in terms of $\varepsilon$-T of TiNi wires with different content of Ag: 0 (a), 0.1 (b), and 0.2 (c) at.%.](image-url)
Table 1. Comparative shape memory effect (SME) parameters in TiNi and TiNi-Ag wires.

|               | \(M'_\text{fs}, \degree C\) | \(M'_\text{fr}, \degree C\) | \(A'_\text{fs}, \degree C\) | \(A'_\text{fr}, \degree C\) | \(\Delta T, \degree\) | \(\varepsilon_\text{rev}, \%\) | \(\varepsilon_\text{res}, \%\) |
|---------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-------------------------|-------------------------|-------------------------|
| TiNi          | –36                         | –164                        | –22                         | 25                          | 55                      | 6.2                     | 0                       |
| TiNi49.9–Ag0.1| –40                         | –145                        | –14                         | 13                          | 60                      | 6.5                     | 0                       |
| TiNi49.8–Ag0.2| –38                         | –108                        | –27                         | 27                          | 68                      | 4.7                     | 0.2                     |

Figure 6 shows the mechanical behavior of the wires during tensile tests. Tensile tests performed at room temperature show that \(B2 \rightarrow R \rightarrow B19'\) martensite transformation caused by deformation occurs in all wires, as evidenced by the phase yield region \(\Delta \varepsilon_y\). The linear section depicted on the \(\sigma-\varepsilon\) curves corresponds to the strain accumulation of the \(B2\) phase. The presence of a stress plateau indicates the development of the stress-induced \(B2 \rightarrow R \rightarrow B19'\) martensite transformation and an increase in the volume fraction of the \(B19'\) phase [39]. A comparative analysis of stress–strain diagrams demonstrates the effect of silver content on the length of the phase yield section \(\Delta \varepsilon_y\), tensile strength \(\sigma_y\), and maximum strain before fracture \(\varepsilon_b\) (Table 2). The maximum values of the \(\Delta \varepsilon_y\), \(\sigma_y\), and \(\varepsilon_b\) in the stress–strain curves belong to the TiNi wire doped with 0.1 at.% Ag.

Figure 6. Stress-strain curves of TiNi–Ag wires with different Ag content: 0 (1), 0.1 (2), and 0.2 (3) at.%.

Table 2. Mechanical properties of the TiNi–Ag wires.

|               | \(\sigma_y, \text{MPa}\) | \(\sigma_y, \text{MPa}\) | \(\Delta \varepsilon_y, \%\) | \(\varepsilon_b, \%\) |
|---------------|--------------------------|--------------------------|----------------------------|-----------------------|
| TiNi          | 350 ± 20                 | 1370 ± 20                | 7 ± 0.5                    | 32 ± 0.5              |
| TiNi49.9–Ag0.1| 435 ± 20                 | 1450 ± 20                | 10 ± 0.5                   | 34 ± 0.5              |
| TiNi49.8–Ag0.2| 400 ± 20                 | 1330 ± 20                | 8 ± 0.5                    | 31 ± 0.5              |

The homogeneous nanocrystalline dislocation-free structure with a significant volume fraction of finely dispersed reinforcing \(\text{Ti}_3\text{Ni}_4\) precipitates is the rationale for the high strength and ductility of the TiNi wire doped with 0.1 at.% Ag. In addition, with increased density of grain boundaries, the uniformity of plastic strain in the material increases, and the internal stress level decreases. The lower mechanical properties of other wires are associated with a high density of dislocations, which are stress concentrators and can cause destruction [38,39]. The presence of a large volume fraction of incoherent \(\text{Ti}_2\text{Ni}\) particles in the TiNi wire and their interaction with the dislocation substructure also lead to a strong stress concentration and make the wires less durable.

4. Conclusions

Ultimately, the concentration range for Ag microalloying was found for TiNi wires (0–0.2 at.%) obtained by intense plastic deformation. The combined method of intense plastic deformation,
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consisting of multiple processes of cold rolling and intermediate annealing (T = 300–450 °C), was used
to produce wire with a diameter of 1 mm. The optimal balance between strength, ductility, and shape
memory parameters was achieved at a silver concentration of 0.1 at.%. The mechano-thermal treatment
promotes the formation of a submicrocrystalline structure and recrystallization of the finely dispersed
phases TiNi_3, Ti_2Ni_3 and Ti_3Ni_4, and reduces the size of Ti_2Ni particles. In the wires, the minimum
silver concentration (0.1 at.% Ag) provides full recovery of the shape (ε_{res} = 0%), the greatest reversible
strain (ε_{rev} = 6.5%), and optimal strength (σ_{b} = 1450 MPa) and ductility (ε_{b} = 34.4%) properties. This is
achieved by virtue of the dislocation-free homogeneous nanocrystalline structure with an average
grain size of 100 ± 50 nm and a high concentration of Ti_2Ni dispersion-strengthening precipitates
in the TiNi(Ag) matrix. TiNi wire 1 mm in diameter and doped with 0.1 at.% silver may be a
suitable alternative to implants of known titanium alloys for use as actively functioning implants in
biomedical applications.

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