High-Strength Behavior of the Al$_{0.3}$CoCrFeNi High-Entropy Alloy Single Crystals

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Abstract: The main disadvantage of fcc (face-centred cubic lattice) high-entropy alloys is the low stress level at the yield point ($\sigma_{0.1}$) at a test temperature above room temperature. This restricts their practical application at high test temperatures from 773 K to 973 K. In this study, we found that a high stress level was reached at the yield point $\sigma_{0.1} \approx G/100–G/160$ (G is the shear modulus) of the [001]- and [144]-oriented crystals of the Co$_{23.36}$Cr$_{23.29}$Fe$_{23.80}$Ni$_{21.88}$Al$_{7.67}$ (Al$_{0.3}$CoCrFeNi) high-entropy alloy (HEA) within a wide temperature range of 77–973 K under tension, due to the occurrence, of nanotwins, multipoles, dislocations under plastic deformation at 77 K and the subsequent precipitation of ordered L1$_2$ and B2 particles. It was shown that grain boundaries are not formed and the samples remain in a single-crystal state after low-temperature deformation and subsequent ageing at 893 K for 50 h. Achieving a high-strength state in the Al$_{0.3}$CoCrFeNi HEA single crystals induces the orientation dependence of the critical resolved shear stresses ($\tau_{cr}$) at $T \geq 200$ K ($\tau_{cr>[144]} > \tau_{cr[001]}$), which is absent in the initial single-phase crystals, weakens the temperature dependence of $\sigma_{0.1}$ above 573 K, and reduces plasticity to 5–13% in the [144] orientation and 15–20% in the [001] orientation.

Keywords: Al$_{0.3}$CoCrFeNi high-entropy alloy; single crystals; high-strength state; L1$_2$ and B2 particles; twinning

1. Introduction

High-entropy alloys (HEAs) are a new class of alloys that have a combination of unique properties, namely, high strain hardening, good plasticity and good ductile fracture strength [1–6]. At low temperatures, single-phase fcc (face-centred cubic lattice) HEAs are only comparable to high-strength materials ($\sigma_{0.1} \sim G/140–G/220$; here G is the HEA shear modulus, equal to 85, 81 and 67 GPa, respectively, at 77, 296 and 773 K [7]), which include materials with a yield point $\sigma_{0.1} \sim G/100$, and they are low-strength ($\sigma_{0.1} > G/360$) at $T \geq 296$ K, due to the strong temperature dependence of the yield point $\sigma_{0.1}(T)$ [2–6]. Commonly, a high level of $\sigma_{0.1}$ in single-phase polycrystals is achieved by a decrease in grain size. However, in fine-grained HEA polycrystals, within the high-temperature range at $T > 673$ K, the $\sigma_{0.1}$ sharply decreases as a result of grain-boundary sliding (GBS) [2].

The fcc Al$_{0.3}$CoCrFeNi alloy is one representative of the HEAs, which is characterized by medium stacking fault energy $\gamma_0 = 0.051$ J/m$^2$ and demonstrates a unique combination of properties, such as high strength, work hardening, good plasticity in tension at room and cryogenic temperatures, corrosion resistance, high oxidation resistance and excellent fatigue resistance [5,8–14].

The high level of strength properties at the yield point, achieved in the Al$_{0.3}$CoCrFeNi HEA due to Al alloying in the single-phase state, led to the development of twinning at $T < 296$ K. In poly- and single crystals of the Al$_{0.3}$CoCrFeNi HEA, twinning when interacting with slip provides a strong strain...
hardening, and at the same time retains plasticity up to 30–70% [5,10]. In addition, the Al0.3CoCrFeNi HEA is a precipitation-hardening alloy, and ordered L12 and B2 particles precipitate in it, which leads to an increase in the strength properties of this HEA [15–18]. Thus, with the precipitation of B2 particles, the tensile strength can be increased from 1200 MPa at room temperature to 1600 MPa at 77 K, while maintaining ductility up to 17% [10]. Despite this combination of unique properties at T < 300 K, the Al0.3CoCrFeNi HEA in the high temperature range at T > 300 K, like other fcc HEAs, remains low-strength [1–5]. This impedes its practical application at high test temperatures from 773 K to 973 K.

In the present paper, in relation to Al0.3CoCrFeNi HEA single crystals, an attempt has been made for the first time to achieve high stresses at the yield point 0.1, within the wide temperature range of 77–973 K, under tension, due to the occurrence of nanotwins, multipoles and dislocations under plastic deformation at 77 K, and the subsequent precipitation of ordered L12 and B2 particles. For this purpose, two orientations, [144] and [001], were chosen for the following reasons. Firstly, in the [144]-oriented crystals, the maximum Schmid factor for twinning, m tw = 0.5, is greater than the Schmid factor for slip, m sl = 0.4, and these crystals are favorable for obtaining a dislocation structure with twins [19,20]. To obtain a dislocation structure without twins, [001]-oriented crystals were chosen in which, conversely, the Schmid factor for slip, m sl = 0.41, is larger than the Schmid factor for twinning, m tw = 0.23 [5]. Secondly, in the chosen orientations, the m sl during deformation remains constant, due to the shear multiplicity in [001]-oriented crystals, which do not change during the precession in the [144]-oriented crystals [21]. This will allow the critical resolved shear stresses (CRSS) τcr of deformed crystals to be determined, provided that the single-crystal matrix is preserved after thermomechanical processing, due to plastic deformation at 77 K and subsequent ageing at 893 K for 50 h.

2. Materials and Methods

Single crystals of the Co23.36Cr23.29Fe23.80Ni21.88Al0.67 (Al0.3CoCrFeNi) (at.%) HEA were grown via the Bridgman method in a helium atmosphere using ingots cast in a resistance furnace (InterSELT, St.-Petersburg, Russia). To achieve a homogeneous distribution of the elements in the bulk of the ingots, they were remelted three times. The dog-bone-shaped tension samples had a gauge length of 12 mm and a cross section of 2 × 1.5 mm2. The samples were cut using wire electrical discharge machining. The damaged surface layer was ground off mechanically, and then electrically polished in 200 mL of an H3PO4 + 50 g CrO3 (phosphoric acid with chromium trioxide) electrolyte at room temperature. The crystals were homogenized in a helium atmosphere at 1473 K for 48 h, and then quenched in water. Homogenization leads to a uniform distribution of elements over the crystal volume, which, as will be shown below, is confirmed by the precipitation of L12 and B2 particles in the defect structure of the samples. To determine the orientation of the crystals, the diffractometric method was used by means of a DRON-3M X-ray diffractometer (Bourevestnik, St.-Petersburg, Russia) with monochromatic Fe Kα radiation, the technique for which was presented in [22]. The high-strength state ρ0.1 ≈ G/100–G/120 was achieved in two steps. The first step consisted of the low-temperature plastic deformation at 77 K of the [001]- and [144]-oriented crystals, within which a stress level of 700 MPa ≈ G/120 was achieved in both orientations. This stress level was reached at a strain of 50% in the [144]-oriented crystals and at strain of 20% in the [001]-oriented crystals. The second step consisted of the ageing at 893 K for 50 h in a helium atmosphere of the deformed crystals, followed by cooling in water. During the selected ageing process, firstly, ordered L12 and B2 particles were precipitated, which contributed to an increase in stresses of ρ0.1 [15–17]. Secondly, the temperature of ageing, 893 K, was not sufficient for the recrystallization of the dislocation structure [2,15,17], which allows the samples to be preserved following deformation in a single-crystalline state. The mechanical properties within a temperature range of 77 to 973 K were determined using an Instron 5969 universal testing machine (Instron, Nowood, MA, USA) at a strain rate of 4 × 10−4 s−1. For the test at 77 K, the sample and grips were immersed in a special vessel with liquid nitrogen, in which both were held for 10 min before the start of the test, and then the sample was deformed in liquid nitrogen. Tests in the temperature range of 200 to 973 K were carried out in a special heat chamber, which is included in the equipment of the Instron 5969 universal
testing machine. The heating/cooling rate of the chamber was 2 K/min. The sample was inserted in grips, heated/cooled together with the chamber, kept at each temperature for 30 min before testing, and then deformed. The CRSS values for slip were calculated using the expression $\tau_{cr} = \sigma_{0.1} \times m_{sl}$ (where $\sigma_{0.1}$ is the uniaxial stress at the 0.1% offset strain yield point). Transmission electron microscopy (TEM) studies were performed using a JEOL-2010 electron microscope (JEOL, Tokyo, Japan) at an accelerating voltage of 200 kV. The thin foils were prepared using double-jet electropolishing (TenuPol-5, “Struers”, Ballerup, Denmark) with an electrolyte containing 20% sulphuric acid in methyl alcohol at room temperature, with 12.5 V applied voltage. The chemical composition of the single crystals after quenching was determined using the X-ray fluorescence method, by means of a wave-dispersive X-ray fluorescence XRF-1800 spectrometer (SHIMADZU, Kyoto, Japan), giving the atomistic percentages Co = 23.36%, Cr = 23.29%, Fe = 23.80%, Ni = 21.88% and Al = 7.67% (at.%). After quenching, the single crystals formed an fcc-based substitution solid solution, and did not contain dispersed particles of the second phase.

3. Results and Discussion

TEM investigations have shown that a planar dislocation structure, with dislocation pile ups, developed in the quenched Al$_{0.3}$CoCrFeNi HEA single crystals after undergoing strain at 77 K [5]. In the [001]-oriented crystals, after undergoing strain of 20%, a high density of dislocations and multipoles was observed, but there were no twins. Thin twins with a thickness of 15–25 nm were found in the [144]-oriented crystals after undergoing strain of 50% at 77 K, simultaneously with a high density of dislocations and multipoles. Figures 1 and 2 display the dislocation structures after ageing at 893 K for 50 h of the deformed [001]- and [144]-oriented crystals of the Al$_{0.3}$CoCrFeNi HEA, having undergone up to 20% and 50% strain, respectively, at 77 K. During the ageing of the deformed crystals at 893 K for 50 h, recrystallization did not occur, grain boundaries were not detected and, therefore, the single-crystalline matrix was retained, as in samples deformed at 77 K (Figures 1 and 2). In the deformed [144]- and [001]-oriented crystals, ageing at 893 K for 50 h led to the precipitation of coherent L1$_2$ and non-coherent B$_2$ particles [15,17]. In the diffraction pattern, clear super-structural reflections from L1$_2$ and B$_2$ particles were detected (Figures 1a and 2c). The particles differed in size $d$ (L1$_2$ with $d = 5–7$ nm [17] and B$_2$ with $d = 30–40$ nm (Figures 1c and 2b). The simultaneous precipitation of L1$_2$ and B$_2$ particles in the matrix apparently suppressed the formation of the nuclei of grain boundaries, and the recrystallization process did not occur during this ageing, which requires further research and is not discussed in this paper. In polycrystalline Al$_{0.3}$CoCrFeNi HEA, particles were precipitated in the grain body, L1$_2$, and at the grain boundaries, B$_2$ [15]. In the initial quenched Al$_{0.3}$CoCrFeNi HEA single crystals, when aged for 50 h at 893 K, only L1$_2$ particles with the same size were precipitated, while B$_2$ particles were not precipitated, due to the very high nucleation barrier, as a result of the need to generate large elastic and fcc/B$_2$ interphase energies [17,18]. In the deformed [144]- and [001]-oriented crystals during ageing at 893 K for 50 h, L1$_2$ particles were distributed uniformly in the matrix, as was shown earlier in [17]. The precipitation of B$_2$ particles occurred along the boundaries of multipoles and at dislocations, as well as near the twin boundaries in the deformed [144]-oriented crystals (Figures 1 and 2). Therefore, we can infer that not only grain boundaries in polycrystals, but also dislocations, multipoles and twins in deformed crystals facilitate the nucleation of B$_2$ particles in an imperfect structure.
Figure 4 displays the temperature dependence of the yield point and indicates the level of strain at which the stress of 700 MPa was achieved during deformation.

Figure 3. The stress–strain response in the quenched [144]- and [001]-oriented crystals of the Al0.3CoCrFeNi high-entropy alloy under tensile strain at 77 K; (a) [144] orientation; (b) [001] orientation. The dotted line on the curve shows at what level of tensile strain the stress level of 700 MPa was achieved in the studied orientations. The stress-strain response of the [001]-oriented crystals shown earlier in [3] (with permission from Elsevier, 2020) is given in this figure to demonstrate the orientation dependence of the mechanical behavior of the studied crystals under tensile strain.
Figure 3. The stress–strain response in the quenched [144]- and [001]-oriented crystals of the Al0.3CoCrFeNi high-entropy alloy under tensile strain at 77 K; (a) [144] orientation; (b) [001] orientation. The dotted line on the curve shows at what level of tensile strain the stress level of 700 MPa was reached in the studied orientations. The stress-strain response of the [001]-oriented crystals shown earlier in [5] is given in this figure to demonstrate the orientation dependence of the mechanical behavior of the studied crystals under tensile strain.

Figure 4. (a, b) Temperature dependence of yield point and (c) critical resolved shear stresses of the Al0.3CoCrFeNi HEA single crystals under tensile strain. The temperature dependences of the yield point and critical resolved shear stresses of the quenched [001]-oriented crystals shown earlier in [5] (with permission from Elsevier, 2020) are given in this figure to demonstrate the absence of the orientation dependence of the critical resolved shear stresses in the quenched Al0.3CoCrFeNi HEA single crystals under tensile strain.

It can be seen that after low-temperature deformation at 77 K, the $\sigma_{0.1}$ of the [144]- and [001]-oriented crystals increased relative to the initial crystals in the studied temperature range; by 320 MPa at 77 K and 200–330 MPa at $T \geq 296$ K. The precipitation of L12 and B2 particles additionally increased $\sigma_{0.1}$ by 130 and 200 MPa at 77 K, and by 80–130 MPa at $T \geq 296$ K, relative to the deformed [001]- and [144]-oriented crystals, respectively (Figure 4a,b). As a result, in the [144]- and [001]-oriented crystals, $\sigma_{0.1} \approx G/100$–$G/160$ within a temperature range of 77–973 K, due to joint hardening with a
dislocation structure and ordered particles (Figure 4a,b). Thus, Al_{0.3}CoCrFeNi HEA single crystals became high-strength at T < 296 K, and close to high-strength at T ≥ 296 K. In this state, as shown in Figure 5, Al_{0.3}CoCrFeNi HEA single crystals remained plastic.

![Figure 5](image-url)

**Figure 5.** Plasticity in the Al_{0.3}CoCrFeNi HEA single crystals: (a) [\{144\}] orientation; (b) [001] orientation. Plasticity of the quenched [001]-oriented crystals was taken from [5] (with permission from Elsevier, 2020).

In the initial quenched [\{144\}] and [001]-oriented crystals of the Al_{0.3}CoCrFeNi HEA, the τ\textsubscript{cr} values were found to be independent of the crystal orientation within the studied temperature range, and the Schmid law was satisfied (Figure 4c, curve 1) [5,17]. The temperature dependence of the τ\textsubscript{cr}(T) of both orientations consists of two sections, characteristic of the slip deformation of fcc substitution alloys [5,23]. At T < 373 K, there is a strong temperature dependence of τ\textsubscript{cr}(T) (the thermally activated τ\textsubscript{S} component of τ\textsubscript{cr}), associated with the thermally-activated interaction of dislocations with substitution atoms. At T > 373 K, τ\textsubscript{cr}(T) is weakly dependent on temperature as G(T), and the athermal τ\textsubscript{G} component of τ\textsubscript{cr} is observed. The ratio τ\textsubscript{cr}(77 K)/τ\textsubscript{cr}(300 K) = 2.1. Consequently, the behavior of the initial Al_{0.3}CoCrFeNi HEA single crystals is typical of low-strength fcc alloys [23].

Upon reaching a high-strength state at the yield point σ_0.1, Al_{0.3}CoCrFeNi HEA single crystals showed features of mechanical behavior that had not previously been manifested in the quenched HEA single crystals. Firstly, in the deformed [\{144\}] and [001]-oriented crystals of the Al_{0.3}CoCrFeNi HEA, the τ\textsubscript{cr} were found to be independent of the crystal orientation at 77 K only, since at the low-temperature deformation, the same stress level of 700 MPa was artificially given, and m\textsubscript{al}[\{144\}] ≈ m\textsubscript{al}[001] [19,20]. At T ≥ 200 K, τ\textsubscript{cr} became dependent on the crystal orientation, and τ\textsubscript{cr}[\{144\}] > τ\textsubscript{cr}[001] (Figure 4c).
According to the known models of hardening under the assumption of the additivity of the contributions to \( \tau_{cr} \) [16,24], the effect of hardening from a defect structure (dislocations, multipoles, twins), L1 and B2 particles can be determined in the [\( \bar{1}44 \)]-oriented crystals,

\[
\tau_{cr} = \tau_{cr}^Q + \alpha G b p^{1/2} + \delta n G b x^{-1} + \tau_{cr}^{L12} + \tau_{cr}^{B2}
\]

(1)

and in the [001]-oriented crystals,

\[
\tau_{cr} = \tau_{cr}^Q + \alpha G b p^{1/2} + \tau_{cr}^{L12} + \tau_{cr}^{B2}
\]

(2)

Here, \( \tau_{cr}^Q \) are the critical shear stresses of the quenched undeformed crystals, which do not depend on the crystal orientation [5,25]; \( \Delta \tau_{cr}^{DS} = \alpha G b p^{1/2} \) is the hardening caused by dislocations (\( \alpha \) is the constant [3], \( G = 80 \) GPa [7] is the shear modulus of HEA, \( b = 0.25 \) nm is the Burgers vector of the perfect dislocation, \( \rho \) is the dislocation density); \( \Delta \tau_{cr}^{Tw} = \delta n G b x^{-1} \) is the contribution to hardening from twins (\( \delta \) is a constant, \( n = 5-7 \) is the number of dislocations in a pile up at a tensile strain of 5\%, \( x \) is the average distance between twin boundaries [26]); and \( \tau_{cr}^{L12} \) and \( \tau_{cr}^{B2} \) are the contribution to hardening from L1 and B2 particles, respectively.

L1 particles are coherent [15–17], and their effect of hardening is determined by the contribution of the elastic stress fields due to the mismatch between the lattices of the particle and matrix E, and the long-range order in Ni3Al particles:

\[
\Delta \tau_{cr}^{L12} = \Delta \tau_1 + \Delta \tau_2
\]

(3)

where \( \Delta \tau_1 \) is the contribution of elastic stress fields from the L1 particles [24],

\[
\Delta \tau_1 = 3 G / E \left( \frac{f_r}{b} \right)^{1/2}
\]

(4)

and \( \Delta \tau_2 \) is the stress due to ordering in the L1 particles [16,24],

\[
\Delta \tau_2 = \frac{\mu}{k b} \left( \frac{4 \mu f r}{\pi B} \right)^{1/2} - f
\]

(5)

Here, \( E = \Delta a / a_m \) is the lattice mismatch parameter of the initial matrix, \( a_m \), and L1 particles, \( a_p (\Delta a = a_m - a_p) \), \( r \) is the particle radius in the slip plane, \( f \) is the volume fraction of particles estimated by TEM, \( b = 0.25 \) nm is the modulus of the Burgers vector of the a/2 <110> dislocation, \( \mu \) is the antiphase boundary energy, \( k \) is the number of dislocations in the complex and \( B = (G b^2) / 2 \) is the line tension. The contribution to hardening from L1 particles, estimated earlier for single crystals of the studied HEA according to Equations (3)–(5), was 18 MPa [17]. Moreover, the \( \Delta \tau_{cr}^{L12} \) turned out to be the same for crystals of all orientations, since after the precipitation of L1 particles, the orientation dependence of CRSS was not observed, and the Schmid law was fulfilled [17].

B2 particles are non-coherent [15,27], and their contribution to hardening, according to [24], can be estimated by the Orowan equation:

\[
\tau_{cr}^{B2} = G b / L
\]

(6)

Here, \( L = 250 \) nm is the average distance between B2 particles. The contribution to hardening from B2 particles, with an average distance between them of \( L = 250 \) nm, was 80 MPa according to Equation (6), and turned out to be close to the experimentally obtained data, whereby \( \Delta \tau_{cr}^{B2} = 60-80 \) MPa in crystals of both orientations at 77 K. In both orientations, the contribution to the hardening from B2 particles was the same, and remained within the temperature range when studying the temperature dependence of \( \tau_{cr}(T) \) (Figure 4c). This correlated with the close volume fraction of B2 particles in the crystals of the studied orientations, which was ~20\%. The volume fraction of particles was determined.
from dark-field images, and 10 pictures from different places on thin foils were used to estimate it. At 77 K in the studied orientations, after ageing, the \( \tau_{cr} \) values turned out to be close (Figure 4c). In addition, in the Al\(_{0.3}\)CoCrFeNi HEA single crystals after ageing at 973 K at 50 h, when only B2 particles were precipitated, the \( \tau_{cr} \) values were not dependent on the crystal orientation in a wide temperature range, as was shown earlier in [27]. All this qualitatively indicates a similar volume fraction of the precipitated B2 particles in the deformed [T44]- and [001]-oriented crystals. Consequently, in the deformed [T44]- and [001]-oriented crystals of the Al\(_{0.3}\)CoCrFeNi HEA, the contribution to the CRSS value from B2 particles, as well as from L1\(_2\) particles, does not depend on the crystal orientation.

Thus, the analysis of Equations (1) and (2) and the experimental data presented in Figure 4c shows that the \( \tau_{cr}^{Q}, \tau_{cr}^{12} \) and \( \tau_{cr}^{B2} \) in the deformed and aged Al\(_{0.3}\)CoCrFeNi HEA single crystals are independent of the crystal orientation. Consequently, the orientation dependence of \( \tau_{cr} \) in the deformed and aged Al\(_{0.3}\)CoCrFeNi HEA single crystals is due to the orientation dependence of the dislocation structure and, in this case, the dislocation structure with twins makes the largest contribution to the hardening at the yield point, as compared to that without twins. So, \( \tau_{cr}^{Q}([T44]) = \tau_{cr}^{Q}([001]) = 70 \text{ MPa at 296 K.} \)

At 296 K, after low-temperature deformation at 77 K in the [001]-oriented crystals, where there was no twinning, the contribution to hardening at the yield point from the defect structure relative to quenched crystals was \( \Delta \tau_{cr} = \Delta \tau_{cr}^{DS} - \Delta \tau_{cr}^{O} = 160 \text{ MPa} - 70 \text{ MPa} = 90 \text{ MPa} \), and in the [T44]-oriented crystals with twins in the dislocation structure, \( \Delta \tau_{cr} = \left( \Delta \tau_{cr}^{DS} + \Delta \tau_{cr}^{tw} \right) - \tau_{cr}^{Q} = 215 \text{ MPa} - 70 \text{ MPa} = 145 \text{ MPa.} \) In the [T44]-oriented crystals at 296 K, the contribution to hardening at the yield point from the defect structure was 55 MPa greater than in the [001]-oriented crystals (Figure 4c). If we assume that when the stress level of 700 MPa was reached during the low-temperature deformation at 77 K, the contribution of \( \Delta \tau_{cr}^{DS} = \alpha G \rho \gamma_{cr}^{1/2} \) to the [T44]- and [001]-oriented crystals was similar, then the difference in the stresses \( \Delta \tau_{cr} \) in the deformed crystals can be seen to be due to twinning at 296 K. The estimate of the contribution from twins \( \tau_{cr}^{tw} = \delta n G \rho x^{-1} \) to hardening at the yield point—at an average value of \( x = 500 \text{ nm, as determined from TEM data,} \delta = 0.5 \) and \( n = 5 \)—was 100 MPa, which correlates quite well with the experimental data.

Moreover, the orientation dependence of the \( \tau_{cr} \) value of high-strength crystals within a wide temperature range is associated with the different temperature dependence of \( \tau_{cr}(T) \). In the deformed [T44]-oriented crystals with twins in the dislocation structure, the \( \tau_{S} \) and the temperature dependence of \( \tau_{cr}(T) \) were weakened (\( \tau_{cr}(77 \text{ K})/\tau_{cr}(300 \text{ K}) = 1.3 \)), and \( \tau_{cr}(T) \) was determined by the temperature dependence of the shear modulus \( G(T) \) for the HEAs [7]. In the deformed [001]-oriented crystals without twins in the dislocation structure, as in the initial crystals, the strong temperature dependence of \( \tau_{cr}(T) \) was preserved (\( \tau_{cr}(77 \text{ K})/\tau_{cr}(300 \text{ K}) = 1.7 \)). If, when hardened by plastic deformation, the \( \sigma_{0.1} \) in the [T44]- and [001]-oriented crystals was governed by slip, then the temperature dependence of \( \tau_{cr}(T) \) should have the same dependence as the initial crystals, and \( \tau_{cr} \) should not depend on orientation. A weak temperature dependence of \( \tau_{cr}(T) \), like \( G(T) \), was characteristic of twinning from the yield point and its safekeeping over a wide temperature range [28]. A qualitative confirmation of the development of twinning in the deformed [T44]-oriented crystals from the onset of deformation at \( T > 200 \text{ K} \) is the fact that within a temperature range of 200 to 473 K, the \( \tau_{cr} \) is close in magnitude to the CRSS for twinning, \( \tau_{cr}^{tw} = 200 \text{ MPa,} \) obtained with the [011]-oriented crystals of the Al\(_{0.3}\)CoCrFeNi HEA, with a Schmid factor for twinning, \( m_{tw} \), close to that of [T44]-orientation (\( m_{tw} = 0.47 \) in [011]-crystals and \( m_{tw} = 0.5 \) in [T44]-orientation) [5].

Secondly, the precipitation of L1\(_2\) and B2 particles preserves the orientation dependence of \( \tau_{cr} \), induced by low-temperature deformation at 77 K. However, in the deformed [T44]-crystals after ageing, \( \tau_{S} \) and \( \tau_{cr}(T) \) were again increased. The ratio \( \tau_{cr}(77 \text{ K})/\tau_{cr}(300 \text{ K}) = 1.5 \), and this has been revealed to be close in magnitude to the deformed [001]-crystals, with particles in which the onset of plastic flow was associated with slip and \( \tau_{cr}(77 \text{ K})/\tau_{cr}(300 \text{ K}) = 1.7 \). TEM investigations of the dislocation structure, after strain of 5% at 296 K in the aged [T44]-crystals, showed that the onset of plastic flow in these crystals was also associated with slip, rather than twinning. Ordered coherent L1\(_2\) particles, as was shown earlier in [17], suppress the development of twinning in the Al\(_{0.3}\)CoCrFeNi HEA single crystals.
Thirdly, in the strain-hardened \([\bar{1}44]\)- and [001]-oriented crystals, \(\sigma_{0.1}\) and \(\tau_{cr}\) were found to change only slightly within a temperature range of 573 to 973 K (Figure 4). In polycrystalline CoCrFeMnNi HEA with a grain size of 4.4 \(\mu\)m, \(\sigma_{0.1}\) decreased by 1.4 times with an increase in temperature from 673 K to 973 K, due to the GBS, which usually occurs in polycrystals at \(T > T_{cr}\) (\(T_{cr} = 0.5T_m\) is the critical temperature above which GBS occurs in polycrystals; \(T_m\)—melting temperature) (Figure 4b). In strain-hardened \([\bar{1}44]\)- and [001]-oriented crystals, there were no grain boundaries, but there were twins and multipoles. Twins are athermal obstacles to moving dislocations, and there is no sliding on them. Multipoles, like twins, are also obstacles to moving dislocations [5]. Both twins and multipoles provide a high stress level at the yield point \(\sigma_{0.1}\), and its weak dependence at \(T \geq 573\) K. However, the contribution to \(\sigma_{0.1}\) from multipoles is smaller than that from twins at \(T \geq 573\) K (Figure 4). In the hardened \([\bar{1}44]\)- and [001]-oriented crystals with L12 and B2 particles, the \(\tau_{cr}\) values change slightly within a temperature range of 573 to 873 K. At \(T > 873\) K, \(\tau_{cr}\) decreases, which can be associated with an increase in the size of B2 particles, and with the partial dissolution of L12 particles during heating and testing, since the temperature of 873 K is higher than the ageing temperature [15–17], and this requires additional study.

Finally, achieving a high-strength state at the yield point \(\sigma_{0.1}\) in the \(Al_{0.3}CoCrFeNi\) HEA single crystals via deformation and subsequent ageing was accompanied by a decrease in plasticity (3.5–13 times in \([\bar{1}44]\)-crystals and 2.3 times in [001]-crystals) (Figure 5), but at the same time, the samples retained their ductile fracture qualities. With an increase in the stress level at the yield point \(\sigma_{0.1}\), achieved via the dislocation structure during low-temperature deformation, the uniform strain in the crystals decreased. In the \([\bar{1}44]\)-orientation with twins in the dislocation structure, \(\sigma_{0.1}\) increased 3–5.5-fold, and the plasticity decreased to 7–13\% compared to the initial crystals within a temperature range of 296 to 973 K. In the [001]-oriented crystals without twins in the dislocation structure, \(\sigma_{0.1}\) increased 2–4-fold compared to the initial crystals of this orientation, but the plasticity remained equal to 15–20\% (Figures 4 and 5).

It should be noted that in the high-strength \([\bar{1}44]\)- and [001]-oriented crystals of the \(Al_{0.3}CoCrFeNi\) HEA under tension at a test temperature of 773 K, a serrated flow was observed on the \(\sigma(\varepsilon)\) curves, as in the quenched crystals of this alloy under tension and compression [8,17]. Consequently, in high-strength single crystals of the \(Al_{0.3}CoCrFeNi\) HEA upon tensile strain at 773 K, dynamic strain ageing was observed, associated with the formation of Cottrell atmospheres by mobile Al atoms near a moving dislocation core [8]. Dynamic strain ageing qualitatively confirms that during the precipitation of particles, some of the Al atoms are in a solid solution and lead to the blocking of moving dislocations. The locking and rapid unlocking of dislocations from the atmosphere of Al atoms leads to a serrated plasticity of up to 7–13\% in the \([\bar{1}44]\)-oriented crystals, and up to 15–20\% in the [001]-oriented crystals.

4. Conclusions

It was shown that the achievement of a high-strength state at the yield point \(\sigma_{0.1}\) in the \(Al_{0.3}CoCrFeNi\) HEA single crystals via low-temperature straining at 77 K and ageing at 893 K for 50 h provides a high stress level of \(\sigma_{0.1}\) at \(T > 296\) K, and a weak temperature dependence \(\sigma_{0.1}(T)\) at high temperatures \(T \geq 573\) K, which is determined by the absence of GBS yet, while on the other hand retaining plasticity.

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