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High-velocity deformation of $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy: Remarkable resistance to shear failure

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The mechanical behavior of a single phase (fcc) $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy (HEA) was studied in the low and high strain-rate regimes. The combination of multiple strengthening mechanisms such as solid solution hardening, forest dislocation hardening, as well as mechanical twinning leads to a high work hardening rate, which is significantly larger than that for Al and is retained in the dynamic regime. The resistance to shear localization was studied by dynamically-loading hat-shaped specimens to induce forced shear localization. However, no adiabatic shear band could be observed. It is therefore proposed that the excellent strain hardening ability gives rise to remarkable resistance to shear localization, which makes this material an excellent candidate for penetration protection applications such as armors.

Most commercial alloy systems are based on one principal element such as Fe, Al, Ni, with minor additions of other elements to improve properties. More recently, the high-entropy alloy design strategy has emerged, in which five or more elements are mixed in equiatomic or near equiatomic concentrations, such that the high configurational entropy $\Delta S_{\text{mixing}}^{\text{conf}}$ promotes solution formation^{1–3}. The configurational entropy is often ignored in the phase prediction of conventional alloys, because they are based on one major element and their phases would have small mixing entropies. A randomly-equimolar mixed solid solution will have a high configurational entropy, which needs to be considered for the prediction of the equilibrium phases⁴. High-entropy alloys represent a new field of alloy development that focuses attention away from the corners of phase diagrams toward their centers.

Many high-entropy alloys have shown exceptional properties, including remarkable fracture toughness at low temperature⁵, high magnetization⁶, and resistance to fatigue^{7,8}. More recently, Li *et al.*⁹ developed a non-equiatomic $\text{Fe}_{80-x}\text{Mn}_x\text{Co}_{10}\text{Cr}_{10}$ HEA system which exhibits both exceptional high strength and ductility at room temperature. The authors attribute the excellent performance to the versatile deformation mechanisms (including dislocation mediated plasticity, mechanical twinning, and stress induced phase transitions), producing a continuous and steady strain hardening.

The processes that occur when bodies are subjected to rapidly changing loads can differ significantly from those that occur under static or quasi-static situations¹⁰. When materials are subjected to large plastic deformation, the temperature increases because a large fraction of the plastic work is converted into heat. Localization of plastic flow can occur when the heating causes thermal softening that overcomes strain hardening and strain-rate hardening. Adiabatic shear bands (ASB) can act as precursors to brittle or ductile failure¹¹. Thus, it is of great importance to study the dynamic response and microstructure evolution of high-entropy alloys.

Results and Discussion

The alloy contains coarse grains with size $\sim 500\ \mu\text{m}$ and large annealing twins shown in Fig. 1(a). The single phase (fcc) $\text{Al}_{0.3}\text{CoCrFeNi}$ alloy, as demonstrated by neutron-diffraction pattern¹² in Fig. 1(b), contains two elements (Cr and Fe) that crystallize with the body-centered cubic (bcc) structure, two elements (Ni and Al) as face-centered cubic (fcc) and one (Co) as hexagonal close-packed (hcp). This single phase fcc crystal structure is consistent with the XRD analysis of $\text{Al}_x\text{CoCrFeNi}$ (x in molar ratio, $x = 0\text{--}2.0$) high-entropy alloy¹³. The elemental

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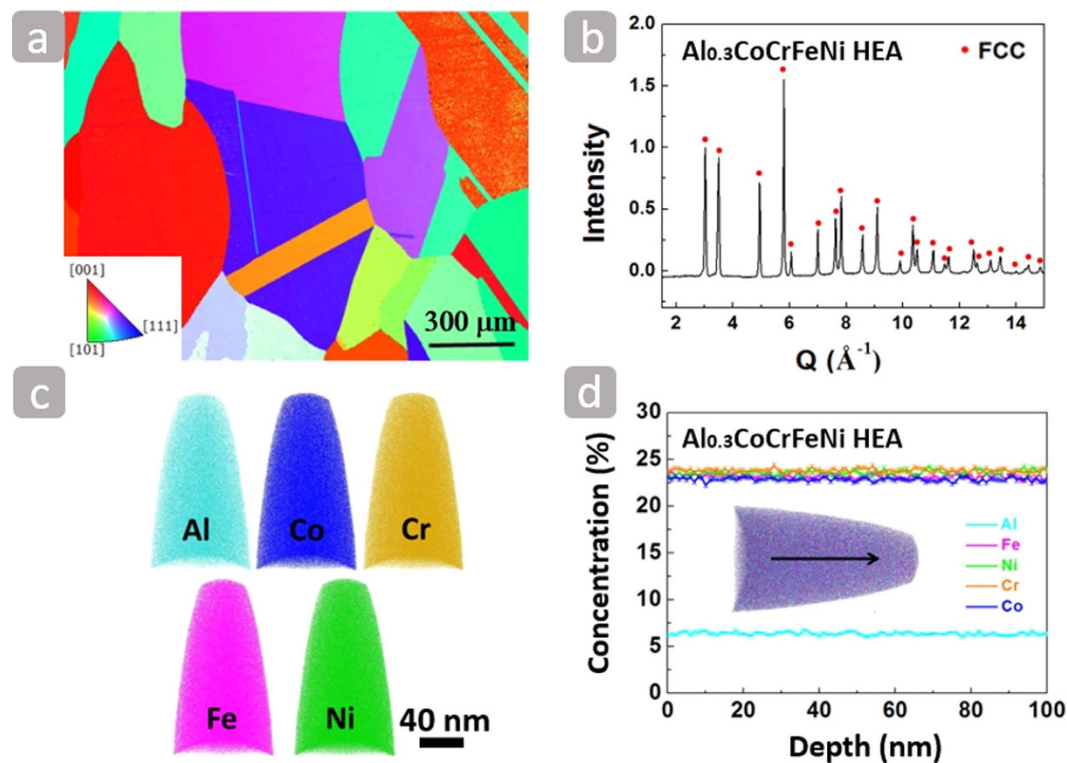


Figure 1. (a) Electron backscattered diffraction (EBSD)-inverse pole figure (IPF) showing the microstructure of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy with annealing twins. (b) Neutron-diffraction pattern¹² of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy. (c) APT analysis showing the homogeneous distribution of Al, Co, Cr, Fe, and Ni elements. (d) APT results of one dimensional element concentration taken along the black arrow.

maps of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy in Fig. 1(c) show that the distribution of Al, Co, Cr, Fe and Ni within the analyzed volume is homogeneous with no indication of segregation or clustering at the atomic scale. The atom probe tomography (APT) results of Fig. 1(d) show the concentrations of the different elements along with the black arrow, which represents the nominal alloy composition.

The compressive true stress-true strain curves at room temperature are plotted in Fig. 2(a). The $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy has a high strain hardening ability during plastic deformation both at quasi-static and high strain rates. The stress-strain curve of coarse-grained (CG) (grain size $\sim 75\ \mu\text{m}$) pure aluminum¹⁴ is also shown in Fig. 2(a) for comparison. Figure 2(b) shows changes of strain hardening rate θ (defined by $\frac{d\sigma}{d\epsilon}$) as a function of true strain for the alloy and CG pure Al at a strain rate of $10^{-2}\ \text{s}^{-1}$. The strain hardening rate of the alloy (above 1000 MPa) is significantly higher than that of CG pure Al. Figure 2(b) shows a negative-slope region followed by a positive-slope region. Such transition is due to a change in the deformation mechanism from dislocation slip-dominated plastic deformation to twinning-dominated plastic deformation. Work hardening suppresses both necking (in tension) and localization (in shear). Both phenomena are interconnected. Lu *et al.*¹⁵ showed that nanocrystalline copper exhibited much higher ductility when the grain boundaries are nanotwins. The cause of the high strain hardening in low-SFE fcc alloys, particularly high-Mn steels, has been widely discussed¹⁶. This high strain-hardening effect is due to the interstitial C atoms of C-Mn dipoles interacting strongly with dislocations or mechanical twins providing barriers to dislocation motion with increasing dislocation storage and thus decreasing their mean free path¹⁷. In the case of dynamic deformation, thermal softening plays an additional role. Xiong *et al.*¹⁸ reported a maximum temperature increase of 55 K for a twin-induced plastic TWIP steel deformed at a high rate of $2400\ \text{s}^{-1}$. This temperature rise was not sufficient to form an adiabatic shear band. In single-phase high-entropy alloys, the lattice is locally distorted because of the occupation of the same crystallographic sites by atoms with different sizes. This local distortion effect is an essential strengthening mechanism^{19–21}. The combination of several strengthening mechanisms such as solid solution hardening, forest dislocation hardening, as well as twinning can lead to an excellent work hardening ability. The yield stress σ_y of the alloy increases from 216 MPa to 503 MPa when the strain rate increases from $10^{-4}\ \text{s}^{-1}$ to $1.8 \times 10^2\ \text{s}^{-1}$. The corresponding strain-rate sensitivity m (defined by $\frac{\log \sigma_y}{\log \dot{\epsilon}}$) is about 0.053 as shown in Fig. 2(c). The strain-rate sensitivity of the pure Al is about 0.028, which is only half of that of the alloy²². In order to study the thermal softening, the dynamic response at the strain-rate $1800\ \text{s}^{-1}$ and different temperatures was examined. The thermal softening parameter $\frac{d\sigma_y}{dT}$ shown in Fig. 2(d) is $\sim -0.4\ \text{MPa/K}$.

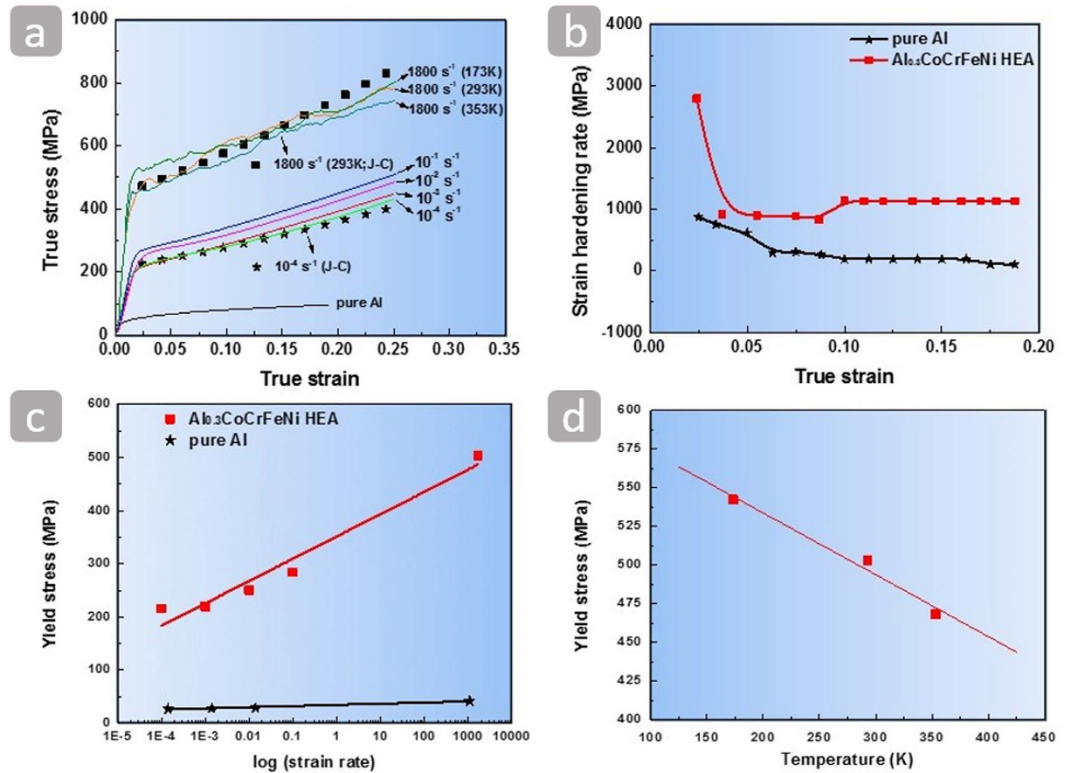


Figure 2. (a) True stress-true strain curves of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy at different strain rates (with Johnson-Cook model²³ fitting curves) and true stress-true strain curve of the CG pure aluminum at the quasi-static 10^{-2}s^{-1} . (b) Strain hardening rate as a function of true strain of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy and pure aluminum at the strain rate 10^{-2}s^{-1} . (c) Yield stress as a function of $\log(\dot{\epsilon})$. (d) Thermal softening of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy at 1800s^{-1} .

The propensity to shear localization can be quantified from the constitutive response. The Johnson-Cook model is the most common phenomenological constitutive equation used to describe the plastic behavior of materials¹⁰. A modified Johnson-Cook constitutive equation²³ is used here:

$$\sigma = (\sigma_0 + B\epsilon^n)(1 + C \log \frac{\dot{\epsilon}}{\dot{\epsilon}_0}) (\frac{T}{T_r})^\lambda \tag{1}$$

where σ_0 is the yield stress, $\dot{\epsilon}_0$ (10^{-4}s^{-1}) is a reference strain rate and T_r (293 K) is a reference temperature. A, B, n, C and λ are experimentally determined parameters: $\sigma_0 = 216\text{MPa}$, $B = 1000\text{MPa}$, $C = 0.145$, $n = 1.2$, $\lambda = -0.18$, $T_r = 293\text{K}$. The work of the deformation can be used to calculate an (adiabatic) temperature rise:

$$dT = \frac{\beta}{\rho C_p} \sigma d\epsilon \tag{2}$$

where ρ is the density and C_p is the specific heat capacity. The parameter β , which is the efficiency of the conversion of the strain energy into heat, is usually taken as 0.9. The density of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy was measured to be 7860kg/m^3 . The specific heat C_p is approximated to be $460\text{J/kg}\cdot\text{K}$ by using a weight averaging method $C_p = w_i \sum_{i=1}^5 C_{pi}$ (w_i is the weight percent and C_{pi} is the specific heat for each element of the alloy). A relationship between the temperature and plastic strain, at a fixed strain rate, is obtained by substituting Eq. (1) into Eq. (2):

$$T = \left[T_r^{-\lambda+1} + \frac{1 + C \log \frac{\dot{\epsilon}}{\dot{\epsilon}_0}}{\rho C_p T_r^\lambda} \times 0.9(1 - \lambda) \epsilon \left(\sigma_0 + \frac{B\epsilon^n}{n+1} \right) \right]^{\frac{1}{1-\lambda}} \tag{3}$$

The temperature at an extreme high strain rate of 10^4s^{-1} and true strain of 0.5 is only 402 K, representing a temperature rise of 109 K.

Joseph *et al.*²⁴ reported that both arc-melted and direct laser sintered $\text{Al}_{0.3}\text{CoCrFeNi}$ alloy samples (with FCC structure) exhibited a yield strength of 200 MPa, excellent strain hardening, and had tensile failure strain above 1. Li *et al.*²⁵ discovered that tensile strength and elongation of the FCC $\text{Al}_{0.3}\text{CoCrFeNi}$ alloy increased with decreasing temperatures and reached 1010 MPa and 68% at 77 K. The Charpy impact energy of the $\text{Al}_{0.3}\text{CoCrFeNi}$ alloy was approximately 413 J at room temperature and 328 J at 77 K. This superior ductility and ultrahigh Charpy

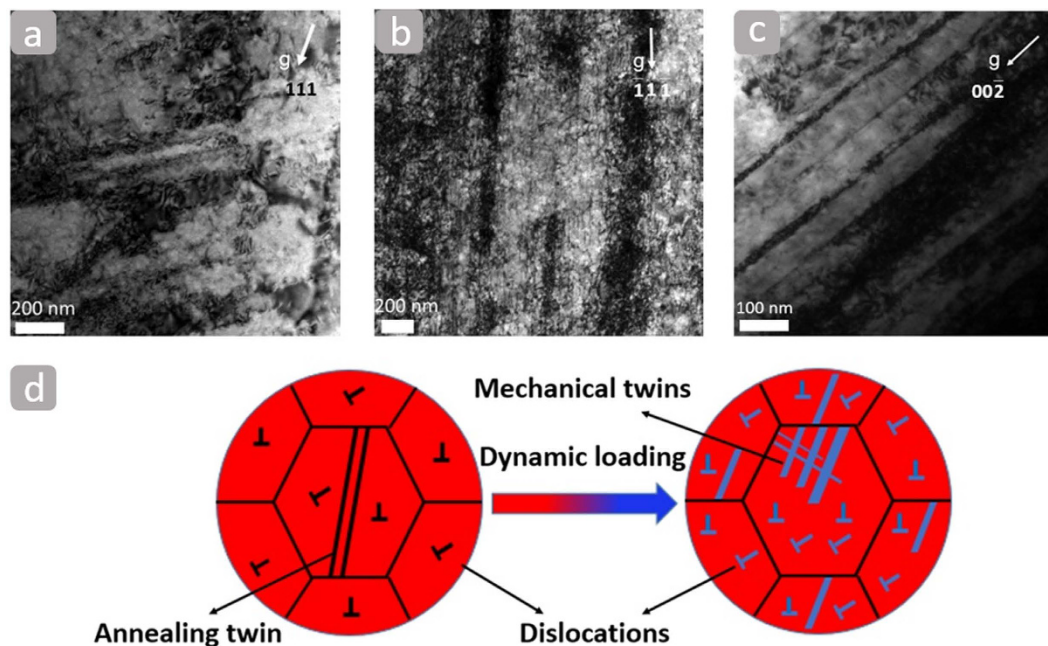


Figure 3. TEM bright-field images of the deformed samples at the strain rates of (a) 10^{-4} s^{-1} ; (b) and (c) 1800 s^{-1} . (d) Schematic sketches showing the deformation mechanisms of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy under dynamic loading.

impact toughness of this alloy are the result of mechanical nanotwinning instead of planar slip dislocations²⁶. This has been also observed by Gludovatz *et al.*⁵ in the CrMnFeCoNi high-entropy alloy. Deformation-induced nanotwinning delays the onset of necking instability (i.e., localized plastic deformation that can lead to premature failure) to higher strains.

In order to understand the strain hardening mechanisms of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy, it is important to establish the microstructure evolution of deformed samples. This was examined at strain rates of 10^{-4} s^{-1} and 1800 s^{-1} by transmission electron microscopy (TEM) and is shown in Fig. 3. Figure 3(a) shows tangles of randomly distributed dislocations in the sample deformed at 10^{-4} s^{-1} , revealing that forest dislocations can lead to steady strain-hardening during plastic deformation. Ma *et al.*²⁷ reported a superior high tensile elongation (of ~80%) for a monocrystalline $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy due to continuous dislocation motion. Figure 3(b) indicates that the high density of dislocations tends to align in one direction and entangle under dynamic loading. Such a high dislocation density configuration has been also observed for the laser-induced shock compression of monocrystalline copper²⁸. Mechanical twins form in many metallic materials, especially those with low stacking-fault energy (SFE). Some fcc single phase materials with high SFE (such as Al and Ni) may not easily form twins except under extreme conditions, such as low temperatures and high strain rates²⁹. Figure 3(c) shows that the parallel bands are mechanical twins, which suggests that mechanical twinning is an important plastic deformation mechanism in the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy at high strain rates. Figure 3(d) summarizes the deformation mechanism of the $\text{Al}_{0.3}\text{CoCrFeNi}$ HEA under dynamic loading, indicating that the dislocation slip and twinning can lead to the excellent strain-hardening ability.

Proportionality between the twin-boundary energy and the stacking-fault energy (SFE) has been reported for most metals³⁰. Some fcc single phase materials with high SFE^{31,32} (Al with SFE $\approx 86 \text{ mJ/m}^2$ and Ni with SFE $\approx 120\text{--}130 \text{ mJ/m}^2$) may not easily form twins except under extreme conditions, such as low temperatures and high strain rates. Kumar *et al.*³³ reported low SFE (below 30 mJ/m^2) of the $\text{Al}_{0.1}\text{CoCrFeNi}$ alloy. He attributed it to the larger atomic size difference between Al and other elements in the alloy. Low stacking-fault energy values ($\sim 30 \text{ mJ/m}^2$) have been reported in NiFeCrCoMn alloy³⁴. Hence, one would expect that addition of Al would cause a high lattice strain per atomic percent in CrFeCoNi as compared to Mn. Mishra *et al.*³⁵ proposed that lattice strain in HEAs might play an important role in dislocation core energy which in turn will have a bearing on stacking-fault formation and its energy. The lattice strain in HEAs raises the base energy of the crystal and thereby reduces the additional energy required to nucleate dislocations and twins. Since $\text{Al}_{0.1}\text{CoCrFeNi}$ has $\sim 1.2 \text{ wt.}\%$ of Al, it is assumed that Al addition might have caused a decrease in SFE from $\sim 30 \text{ mJ/m}^2$ to a lower value. Thus, it is proposed that the SFE of $\text{Al}_{0.3}\text{CoCrFeNi}$ is lower than that of the $\text{Al}_{0.1}\text{CoCrFeNi}$ alloy, leading to the formation of profuse mechanical twins in high strain-rate deformation.

Shear localization has been found to be an important and sometimes dominant deformation and fracture mode in metals, granular ceramics, polymers, and metallic glasses at high strains and strain rates¹⁰. For metals, thermal softening is the first stage of this process, leading to processes of dynamic recovery and recrystallization with associated drops in the flow stress. Calculations by Meyers *et al.*³⁶ revealed that the break-up of the elongated sub-grains and diffusive rotation of the grain boundaries can occur during the deformation process. The complex

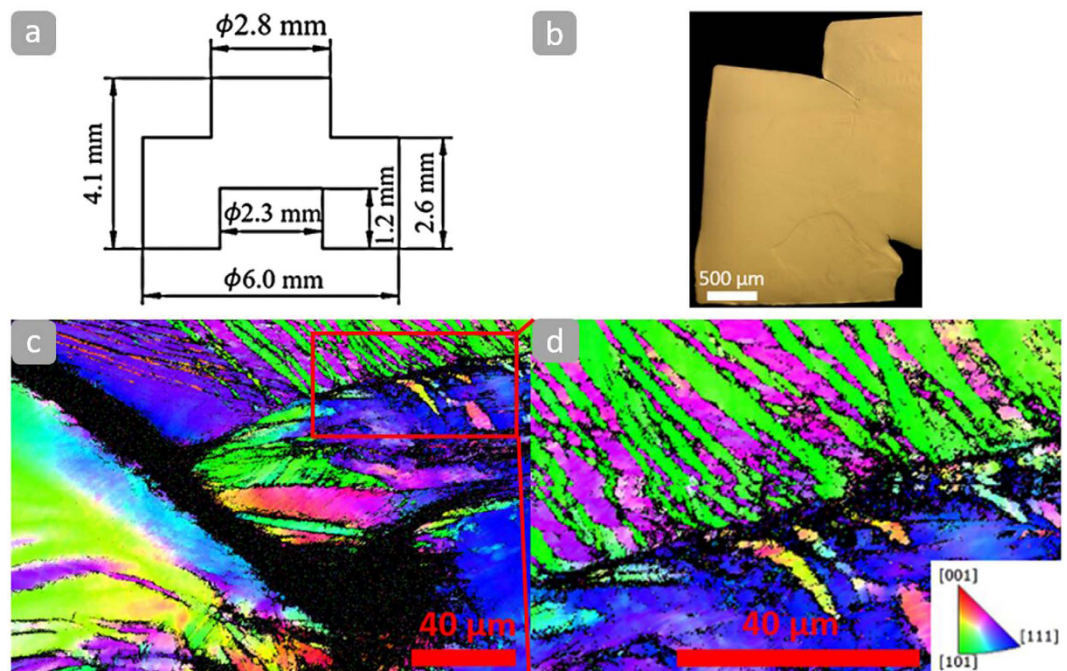


Figure 4. (a) Dimension of the hat-shaped sample. (b) Optical microscopy image of the deformed hat-shaped sample. (c) IPF mapping near the deformation tip. (d) IPF mapping showing deformation twinning in the vicinity of the tip.

inter-relationships between stress, stress state, strain, strain rate and temperature, have been used for pursuing a better design of materials with the objective of postponing and even avoiding localized shear deformation.

An adiabatic shear band can form when the material starts to “soften”. The condition for instability is $\frac{d\tau}{d\gamma} \leq 0$, is obtained from the general function $\tau = f(\gamma, \dot{\gamma}, T)$:

$$\frac{d\tau}{d\gamma} = \left(\frac{\partial \tau}{\partial \gamma} \right)_{\gamma, T} + \left(\frac{\partial \tau}{\partial \dot{\gamma}} \right)_{\gamma, T} \frac{\partial \dot{\gamma}}{\partial \gamma} + \left(\frac{\partial \tau}{\partial T} \right)_{\gamma, \dot{\gamma}} \frac{dT}{d\gamma} \leq 0 \quad (4)$$

The normal stress and strain shown in Equation (1) at a constant strain rate of 10^4 s^{-1} can be converted to the corresponding shear stress and shear strain³⁷ by: $\tau = \frac{\sigma}{2}$ and $\gamma = \sqrt{2e^{2\varepsilon} - 1} - 1$. For a shear strain of 1.1, at constant $\dot{\gamma}$ and T , $\frac{\partial \dot{\gamma}}{\partial \gamma} = 0$, the strain hardening parameter is $\sim 440 \text{ MPa}$, and the thermal softening parameter is $\sim -0.2 \text{ MPa/K}$. Consequently, $\frac{dT}{d\gamma} = 175 \text{ K}^{-1}$. Neglecting the second term in Equation (4):

$$\frac{d\tau}{d\gamma} = \left(\frac{\partial \tau}{\partial \gamma} \right) + \left(\frac{\partial \tau}{\partial T} \right) \frac{dT}{d\gamma} \approx 440 - 0.2 \times 175 = 405 \text{ MPa} \gg 0 \quad (5)$$

Thus, shear instability is not predicted from the constitutive response of this alloy.

The resistance of the $\text{Al}_{0.3}\text{CoCrFeNi}$ alloy to shear localization was then experimentally studied by dynamically-loading a hat-shaped sample to induce forced shear localization. The dimension of the hat-shaped sample is shown in Fig. 4(a). Figure 4(b) shows the optical image of the deformed hat-shaped sample. The electron backscatter diffraction (EBSD)-inverse pole figure (IPF) mapping in Fig. 4(c) indicates that the hat-shaped sample was highly deformed under high strain rates but not fractured. The formation of mechanical twins can be observed in Fig. 4(d) near the deformation tip. No observable shear band can be identified in the hat-shaped sample at an imposed shear strain of ~ 1.1 . The microstructure is severely deformed in the “forced” shear region. Xu *et al.*³⁸ studied shear localization and recrystallization in dynamic deformation of a CG single fcc 8090 Al-Li alloy and observed a recrystallized equiaxed structure with an average grain size of $\sim 0.2 \mu\text{m}$. Dynamic recrystallization, which has been widely observed inside adiabatic shear bands¹⁰, is absent in the shear region of the hat-shape sample, indicating the alloy continues to harden under dynamic loading instead of softening by recrystallization. Jiao *et al.*³⁹ reported pile-ups around the indents in the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy and observed a highly localized severe plastic deformation under the nanoindentation. However, they did not mention shear bands. Indeed, pile-ups are a natural result of indentations because of the volume constraining requirement. Tang *et al.*⁴⁰ found that the grain size of the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy decreased severely from $\sim 350 \mu\text{m}$ to nanocrystalline size due to dynamic recrystallization via high-pressure torsion processing method. These observations indicate that dynamic recrystallization of this alloy can happen under severe plastic deformation. However, adiabatic shear localization of this alloy can only be activated by higher strains and temperatures than the ones imposed

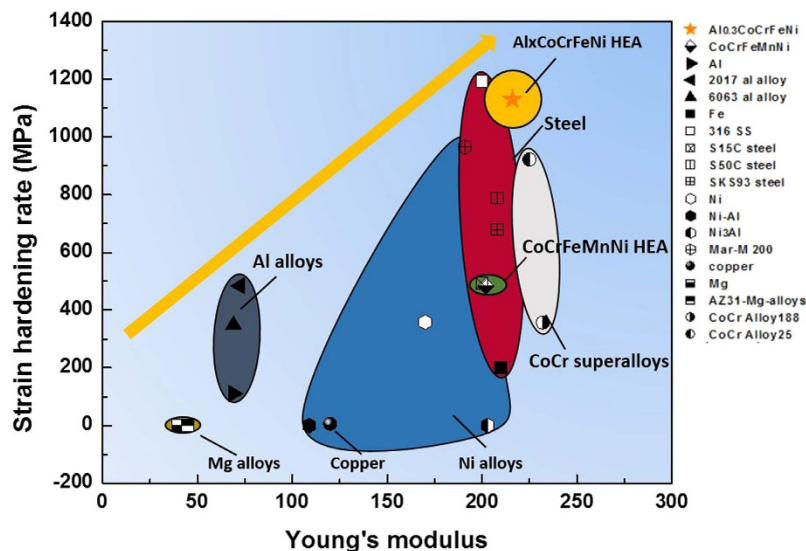


Figure 5. Comparison of strain-hardening ability of $\text{Al}_{0.3}\text{CoCrFeNi}$ HEA with other structural materials. For consistency, strain-hardening rates were calculated at a fixed true strain of 0.2 (in compression) for all materials except for pure Mg which usually fractured at this point.

by our experiments. The enlarged Fig. 4(d) with a $\sim 36\%$ twin area fraction and a $\sim 2.4 \mu\text{m}$ average twin thickness confirms that their formation can play an important role on high strain-rate deformation. Consequently, the dislocation slip and twin-twin reaction can lead to the excellent strain-hardening ability; this, in turn, results in the its extraordinary resistance to shear localization.

In summary, we investigated the mechanical response and microstructural evolution of a high-entropy alloy with emphasis on the high strain-rate regime. The yield stress σ_y increases from 216 MPa to 503 MPa with increasing strain rate from 10^{-4} s^{-1} to 1800 s^{-1} , showing a significant strain-rate sensitivity of 0.053. The strain hardening rate of the alloy is also significantly higher than a CG pure Al and retained in the dynamic regime. The TEM images and EBSD analysis of dislocations and mechanical twinning reveal that (1) the high strain-hardening ability, enabled by solid solution hardening, forest dislocation hardening and twinning hardening, (2) the high strain-rate sensitivity and (3) modest thermal softening give rise to the high resistance to shear localization. Figure 5 shows the excellent performance of the $\text{Al}_{0.3}\text{CoCrFeNi}$ HEA studied here, in comparison with other alloys. Our results suggest that the $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy maintains the remarkable mechanical properties at high strain rates, rendering it to be of great potential for impact-protection (ballistic) applications.

Methods

Sample Processing. The $\text{Al}_{0.3}\text{CoCrFeNi}$ high-entropy alloy was fabricated by vacuum-induction melting with the Al, Co, Cr, Fe and Ni elements to cast a plate of $\sim 127 \text{ mm} \times 305 \text{ mm} \times 19 \text{ mm}$. Then the plate underwent the hot-isostatic-pressing (HIP) at $1,204^\circ\text{C}$ and 103 MPa for 4 hours to reduce defects formed during the casting and cooling processes. Samples were cut from the center of the bulk materials and underwent homogenization at $1,200^\circ\text{C}$ for 2 hours, followed by water quench.

Mechanical testing. The quasi-static compression tests of cylinders were performed in an Instron Universal Testing Machine with the dimensions of 4 mm in diameter and 6 mm in length. Dynamic compression tests were performed using a Split Hopkinson Pressure Bar (SHPB). The SHPB was used to test the dynamic response and shear deformation using two kinds of specimens: cylinders and hat-shaped samples, respectively. The cylinders for dynamic tests have a length of 6 mm and a diameter of 6 mm and were placed in a sealed temperature controlled chamber. Temperature inside the chamber was recorded by thermocouples. Hat-shaped specimens were used to generate high localized shear strain and induce forced shear localization.

Microstructure characterization. Neutron-diffraction experiments were performed at room temperature on the Nanoscale-ordered Materials Diffractometer (NOMAD) at the Spallation Neutron Source (SNS) located at Oak Ridge National Laboratory. This aerodynamic levitator provides a containerless environment, in which HEA samples (2 mm diameter spheres) are suspended above a conical nozzle by flowing argon gas. Samples for the electron backscattered scanning diffraction (EBSD) observation were prepared by standard mechanical grinding and polishing with the colloidal Al_2O_3 . The sharp-tip specimens for atom-probe tomography (APT) were prepared in a FEI Nova 200. The APT experiments were conducted using a CAMECA local electrode atom probe (LEAP), 4000X HR, equipped with an energy-compensated reflection lens. The APT measurements were performed in voltage mode at the temperature of 50 K, pulse frequency of 200 kHz, and pulse fraction of 20%, respectively. At least 5 million ions were collected for each sample to ensure adequate data statistics. The datasets were reconstructed and analyzed using the IVAS 3.6.8 software (CAMECA Instruments). To prepare TEM

specimens, deformed samples were cut to slices of $\phi 3 \times 1$ mm using a linear cutting machine and then slices were mechanically polished to a thickness of 70 μm . Foils with a diameter of 3 mm for the TEM examination were prepared by TenuPol-3 with an etching solution of 30% volume nitric acid and 70% volume methanol. TEM samples were characterized by transmission electron microscopy (TEM) using a model Tecnai G² Polara FEI electron microscope operating at 200 kV.

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Author Contributions

Z. Li and M.A. Meyers formulated the project. H. Diao synthesized Al_{0.3}CoCrFeNi high-entropy alloy. Z. Li carried out the mechanical testing. S. Zhao and Z. Li did the TEM experiments. H. Diao and P.K. Liaw contributed the EBSD, neutron-diffraction pattern and APT analysis. Z. Li and M.A. Meyers wrote the paper, and all authors contributed to revising the paper.

Additional Information

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