

Original Article

Strain-rate effects and dynamic behavior of high entropy alloys



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ABSTRACT

The novel class of multicomponent alloys, also known as high-entropy alloys (HEAs) exhibits excellent properties under low strain-rate conditions. These are especially revealed in the high strength of nanocrystalline CoCrFeMnNi and AlNbTiV alloys, and in the high fracture toughness of AlCoCrCuFeNi and NbMoTaW alloys. Nevertheless, up to now, the dynamic behavior of these high-entropy alloys has not been investigated to the same extent as the quasi-static response. A significantly different mechanical response, such as spallation failure and shear localization, manifests itself when materials are subjected to dynamic loading. Shear localization is an essential precursor to shear failure; studies addressing retardation of its onset are important because of their relevance to applications such as armor for military use. The resistance to shear localization is associated with the extensive work hardening ability enabled by dislocation slip, twinning, and phase transformations which override thermal softening. Apart from shear localization, in contrast to the conventional fracture failure dictated by fracture toughness, spallation resulting from tensile pulses and involving propagating micro-cracks and/or micro-voids also plays an important role in dynamic performance. Although distinctive behaviors have also been reported for other conventional metallic materials under dynamic loading, the unique characteristics of HEAs warrant this review.

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1. Introduction

Numerous studies have been directed towards a novel class of alloys with multiple-principal elements, also known as highentropy alloys (HEAs). Conventional alloys normally contain one single principal element with addition of other elements in varying amounts to tailor properties. In contrast, high entropy alloys [1,2] are comprised of five or more principal elements to maximize configurational entropy ΔS_{config} ($\Delta S_{config} = RlnN$, where N is the number of elements in system), which promotes formation of liquid or random solid solution phases with simple structures such as body-centered cubic (BCC) and face-centered cubic (FCC) and deters the formation of brittle intermetallic phases. High-entropy alloys (HEAs) as novel metallic materials were first discovered in 2004 [1,2]. Based on their constituent elements, HEAs can be classified into five different classes: 1) 3-D transition metals only, 2) transition metals with

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elements possessing larger atomic radii, 3) HEAs composed of refractory metals, 4) ceramic HEAs comprised of oxygen, boron and other anions and 5) others which are comprised of 4f elements Dy, Lu, Tb, Y. Due to multiple constituent elements in HEAs, unique properties, i.e., high hardness/strength, excellent wear resistance and damping capacity, have been discovered. Although mechanical properties at low strain-rates have been the focus of many research efforts and reviews [3–8], properties when the HEAs are subjected to dynamic loading have yet to be fully investigated [3–6].

1.1. Core effects of HEAs

In comparison with conventional alloys, four core effects, which are: (1) High-entropy effect; (2) Sluggish diffusion effect; (3) Severe lattice distortion effect; (4) Cocktail effect, are proposed as possible reasons that render HEAs unique, as shown in Fig. 1. Discussion of each individual effect will be presented first. The high entropy effect describes the thermodynamics which can influence equilibrium structures and microstructures, thereby playing an important role in alloy design. Sluggish diffusion effect is involved in other aspect, which is phase transformation. As for severe lattice distortion effect, it is of great importance to mechanical properties (i.e., dislocation movement) and it is closely related to three other effects. The cocktail effect derives its name from synergy of compositions and microstructures. Finally, the short-range ordering effect, as a newly added crucial effect of HEAs, originates from distinctive energetic favorability between different atomic pairings.

The high entropy effect, which is explained via thermodynamic theory, illustrates the relationship between enhanced system stability and increasing number of elements. According to $\Delta G_{mix} = \Delta H_{mix} - T\Delta S_{mix}$ [9] (where T is the temperature, ΔG_{mix} is mixing Gibbs free energy, ΔH_{mix} is mixing enthalpy and ΔS_{mix}



Fig. 1 – Schematic illustration of four core effects in HEAs.

is mixing entropy), an increment of mixing entropy promotes smaller Gibbs free energy and enhances solid solution phase stability, especially at elevated temperature. Although random solid solution is considered as the stable phase in HEAs, the formation of multiphases, such as nanoprecipitates and second phases, is commonly found, which can be attributed to local stress/strain, varying interatomic interaction, enthalpy and other effects [10]. One recent study [11] discovered a secondphase formation when the CoCrFeMnNi HEA was subjected to a 700 °C annealing treatment for 1000 h, which questioned the stability of random solid solution phase in HEAs. In addition, owning to high configurational entropy, solid solution phase is believed to be stable status of HEAs. That is, nonconfigurational entropy and mixing enthalpy are thought to be negligible in HEAs. However, such assumption can only be valid in ideal solid solutions, where total Gibbs free energy is governed by configurational entropy. Otto [12] proposed in his work that none of the equiatomic alloys comprised of five principal elements displayed a single solid solution phase, suggesting a higher enthalpy and non-configurational entropy that can no longer be overlooked. Nevertheless, An et al. [13] achieved a near single phase in CoCrCuFeNi-based HEAs, regardless of positive mixing enthalpy, by applying radio frequency sputtering combined with rapid quenching.

A certain lattice potential energy barrier (LPE) needs to be overcome for atomic migration to occur from one site to another. For HEAs, significantly different diffusional behavior in comparison with conventional alloys, namely sluggish diffusion effect, has been put forward by Yeh [1]. Such effect is mainly derived from a substantial variety of neighboring atoms of each lattice site, thereby generating different LPEs. In contrast, pure metals have constant LPEs throughout all lattice sites due to identical surrounding atmosphere of atomic arrangement. One can imagine that if an atom jumps from a high-energy site to a lower energy vacancy site, it can be "trapped". Meanwhile, if an atom is initially at a low-energy site, then jumps to a highenergy vacancy site, it tends to return to the original site. Both situations promote sluggish diffusion and inhibit atomic migration.

The lattice distortion effect is mainly derived from a variety of atomic sizes in the lattice. Due to the nature of vast amounts of constituent compositions in HEAs, the differences in atomic sizes and bonding energies contribute to high tendency to form the severe lattice distortion effect. Assuming no chemical ordering, all elements are randomly placed in the lattice and the atoms with larger radii will push the smaller ones away from their original positions, thereby generating strain energy and affecting the HEAs' properties. Atomic mismatches can be large in some circumstances, thereby forming a severe lattice distortion effect. Such effect has been observed in both BCC [14,15] and FCC [16,17] matrix HEAs.

The term "cocktail effect" was first put forward by Ranganathan [18] to describe metallic alloys. Such effect illustrates that the mechanical and physical properties can be altered unexpectedly by mixing multiple components. For example, with increasing atomic fraction of Al, rising hardness and conversion of FCC matrix to BCC matrix phase have been observed in CoCrCuNiAl_x HEAs [1]. The drastic change of hardness (from 133HV to 655HV) achieved by altering atomic fraction of Al is partly attributed to an increasing atomic mismatch and strong interatomic binding force between Al atoms and other metallic atoms. This effect really represents unpredictable properties caused by the great variety of compositional choices for HEAs.

Short-range ordering (SRO) has been widely observed in binary alloy systems [17,19–21] and is considered to be one of the major strengthening mechanisms. Preceding studies have investigated SRO in both BCC-based [22,23] and FCC-based [24,25] HEAs. This effect is closely related to bonding energy between atoms and local strain energy. Although HEAs are believed to be random solid solutions, some studies [26-29] point out that certain degrees of ordering arrangement of atoms can take place at local domains, resulting in SRO structures. If the mixing enthalpy between certain atoms is lower than others in system, denoting a stronger favorability, then SRO can develop. Due to the complexity of multicomponent systems, energetic favorability among atomic pairings can be vastly different. Regarding the formation of SRO, another important factor is local strain energy. Due to lattice distortion, strain energy at some local domains can be much higher than the average, suggestive of metastable state regions. Thus, in order to mitigate strain energy, atoms need to realign, and this leads to SRO. Unlike long-range ordering, such ordered atomic arrangements can only persist over a few to several interatomic distances. Simulations predicting SRO with corresponding four groups of configurations in CrCoNi alloy were reported by Ding et al. [30] illustrating a strong relationship between local chemical order and stacking-fault energy. In addition to computational simulation, such local ordering structure has also been discovered in CrCoNi alloy via energy-filtered transmission electron microscopy by Zhang et al. [29]. They have concluded that tailoring SRO is of great importance to mechanical properties, such as promoting dislocation planar slip in FCC metals and altering Peierls-Nabarro energy barriers. The interaction between SRO and dislocation carriers, e.g., twinning, dislocation, lattice resistance, is summarized in Fig. 2.

1.2. Deformation mechanisms

Due to intriguing features derived from inherently complex compositions of HEAs, their response to dynamic loading can be very distinctive in contrast to conventional metallic materials. For instance, the SFE of HEAs can be varied significantly from one domain to another due to presence of local chemical order in comparison with uniform SFE throughout conventional metallic materials. Such unique phenomenon can strongly affect dislocation glide mode and their dissociation, which leads to different deformation substructures such as deformation twins, deformation-induced microbands, slip bands, phase transitions and, in some extreme cases, amorphization. Phase transformations in HEAs are significantly affected by residual strains and lattice distortion, which result from atomic size mismatch between different constituent elements. These residual strains can subsequently contribute a "driving force" to induce phase transformation [31]. In addition, the deformation mechanism can also be altered when the loading condition is changed, and the changing path can be different in HEAs. For example, rising strain rate is widely believed to promote wavy to planar glide transition [32]; however, the dislocation substructures in FCC solid solution HEAs may not follow this rule due to "planar softening rule" [33]. One of the main foci in the current review is how the change in loading conditions can actually alter the dynamic properties considering the unique characteristics of HEAs.

2. Mechanical behavior

2.1. Dynamic behavior

The study of the dynamic response of materials involves several disciplines, including physics, chemistry, mechanics, and materials science. The classic definition of dynamic behavior requires that materials undergo a non-static condition of rapidly changing loading; they therefore exhibit very different properties in comparison with materials subjected to static and quasi-static conditions of loading [34]. Besides, in contrast to quasi-static and static deformation that maintain static equilibrium condition, due to higher strain-rate subjected at dynamic deformation, inhomogeneous distribution of stress is often generated. Such is the case with elastic, plastic, and shock-wave propagation.

One important effect that occurs when dislocations are forced to move at high velocities is viscous phonon drag [35–37]. Such effect is due to the rapidly propagating dislocations interacting with the elastic lattice vibrations. Phonon drag is negligible at quasi-static conditions because the dislocation movement is too slow to interact with the phonons. In addition, such viscous drag effect at low temperature, at which electron drag becomes more active and dominant, is prohibited due to small population of phonons [38]. This high-



Fig. 2 – Schematic illustration of the interplay between SROs and different deformation carriers in a representative FCC HEA [226].

velocity effect contributes to the scattering of phonons, which significantly influences the viscosity and the frictional force. Operative phonon drag effect is always coupled with vastly increased yield strength.

Apart from the phonon drag effect, additional deformation mechanisms, for instance, deformation twinning, phase transformations, shear banding, are operative in metallic materials subjected to high strain rate deformation [34,35]. One major downside of dynamic deformation is, often, a deteriorating ductility. Therefore, a comprehensive understanding of high strain-rate plastic deformation behavior is essential for its potential use in structural engineering applications. In terms of experimental methods, strain-rate ranges from 10 to 10^8 s^{-1} can be produced by varying approaches, such as dynamic compression on split-Hopkinson pressure bar, shock compression in gas guns, laser-driven systems, detonation waves from explosives, ballistic impact, etc. However, there is a limit to the strain rates that can be achieved experimentally, and the mechanical response at strain rates higher than on the order of 10⁹ s^{-1} is difficult to investigate. In this regime we have to rely on simulations, especially molecular dynamics.

2.2. Thermally activated process

The deformation behavior of metals and alloys strongly depends on strain rate and temperature, indicative of a thermally activated process [32]. This process is predominantly determined by required stress-dependent Gibbs free energy $\Delta G(\sigma)$ to enable dislocation glide. In order for a dislocation to move, possible obstacles, such as solute atoms, precipitates, need to be bypassed. The activation volume V is usually used to represent how much material is involved in the thermally activated processes. It is defined as:

$$V = \frac{\partial (\Delta G(\tau))}{\partial \tau} \Big|_{T} = -M \frac{\partial (\Delta G(\sigma))}{\partial \sigma} \Big|_{T} = MKT \frac{\partial \ln(\dot{e})}{\partial \sigma} \Big|_{T} = bA$$
(2.1)

where M = 3.06 is an orientation factor, σ is uniaxial stress, τ is shear stress, b is the Burgers vector., T is temperature, \dot{e} is plastic strain and A is activation area swept by dislocation. V is usually expressed in units of b^3 . Given that incremental strength derived from grain size strengthening is usually thought to be independent from temperature and strain rate [39,40], solid solution and dislocation—dislocation interaction provide dominant contributions to the strengthening effect. The activation volume for solid solution FCC HEAs can be expressed as follows:

$$\frac{1}{V} = \frac{1}{V_{ss}} + \frac{1}{V_f}$$
(2.2)

where the two terms denote contributions caused by solid solution strengthening and forest hardening.

The activation volume contributed by solute strengthening can be written as [41–43]:

$$V_{ss} = \frac{0.55\Delta E_b(T)M}{\sigma_{ss}}$$
(2.3)

where $\Delta E_b(T)$ is the energy barrier that needs to be bypassed for the change of shear modulus at temperature T, σ_{ss} is the incremental stress due to solid solution strengthening. The activation volume derived from the multiplication of dislocations can be formulated as [44]:

$$V_f = \frac{M\alpha G b^2 w}{\sigma_f} \tag{2.4}$$

where σ_f is the incremental stress due to forest hardening, w is the length scale of junction barrier overcome by a dislocation, and parameter α represents rate dependence of breaking dislocation from the forest network. In contrast to conventional FCC metals with high activation volumes ranging from 200 to 2000 b^3 , a much smaller activation volume between 7 and 77 b³ was reported [45] for CoCrFeMnNi alloy at temperatures of 77 K and 298 K, respectively. The difference in their activation volume is attributed to much smaller number of solute atoms in CoCrFeMnNi alloy. In addition, deformation mechanisms of micro/nano-crystalline materials can be identified by their activation volume, where dislocation glide and dislocation-dislocation interaction prevail in the regime of ~100 b^3 to 1000 b^3 , and interphase boundaries dominate in the regime of ~10 b^3 . Fan et al. [46] proposed a flow stress "upturn" at the strain rate between 10^3 s^{-1} to 10^4 s^{-1} . This "upturn" has been previously attributed to the onset of non-Arrhenius response [47-50]. They have discovered an ultrasmall activation volume in the high stress region, which causes high strain-rate sensitivity.

2.3. Strengthening mechanisms

Metals are characterized by non-directional metallic bonding and higher probability to form defects. Thus, the development of strengthening mechanisms is essential. Apart from tuning chemical bonding or crystalline structure, interfering with the movement of defects via stress fields or substructures is the most effective method of strengthening metals. The most common strengthening mechanisms are grain-boundary strengthening, precipitation strengthening, two-phase hardening, solute strengthening, and order strengthening [55,66].

2.3.1. Dislocation strengthening

Forest hardening is a strengthening effect that arises from increasing dislocation density coupled with the formation of dislocation junctions during plastic deformation. Such junctions are able to restrain dislocation motion, leading to strain hardening, until it is compromised by sufficiently high local stress. The relationship between critical resolved shear stress to destroy junctions and dislocation density can be expressed by the following equation:

$$\tau_{\rm CRSS} = \alpha G b \sqrt{\rho} \tag{2.5}$$

where α is a parameter, ρ is dislocation density.

With respect to the strain rate effect on dislocation substructures, Sevillano et al. [52] proposed that a rising strain rate can generate comparable effect to reducing the temperature and SFE. Coupled with the rising rate is the shift of dislocation glide mode, from wavy to planar, which thereafter refines the dislocation cells and traps dislocations inside the cell structure [53]. Classic CoCrFeMnNi alloy dislocation movement was found to be governed by planar glide on {111} slip planes [39,54] at strain rates below $3 \times 10^3 \text{ s}^{-1}$, which was also commonly found in the family of low SFE [55,56], shortrange ordered [33], or high frictional stress [57] metallic materials. Although planar slip is favored in low SFE materials, it has also been discovered in some solid solution FCC alloys with high SFE due to the "glide plane softening" effect [33]. This effect relies heavily on the short-range ordering. Wong and his co-workers [58] developed a tunable dislocation glide mode for $Al_{0.3}$ CoCrFeNiMn_x by modifying the atomic fraction of Mn. While low Mn alloys displayed wavy glide dominance, their high Mn counterparts were dominated by planar glide. In the higher strain-rate region (9 × 10³ s⁻¹), Tsai et al. [54] proposed a transition of deformation mechanism, from dislocation-glide to nano-twinning in the CoCrFeMnNi alloy.

2.3.2. Solution strengthening

The strengthening effect derived from interaction between solute atoms and dislocations, more specifically, the former's strain field and the latter's stress field [51,56-59], is widely known as the reason behind solid solution strengthening. As a result, an increasing resolved shear stress is required to effect continuous plasticity by dislocation motion, thereby subsequently leading to an enhanced yield strength. Two sets of models, strong (Fleischer model) [60-63] and weak pinning models (Labusch model) [64,65], as shown in Fig. 3, were developed to describe solid solution strengthening in alloys. In the strong pinning model, solutes are treated as individual point obstacles, resulting in strong pinning effect. Meanwhile, in the weak pinning model, energetic fluctuation due to collective effect in area with randomly distributed solutes, decreases the dislocation energy, thereby providing a weak pinning effect. Contrary to the Fleischer model, in which solute atoms are expected to be sparsely and randomly distributed with low concentration, the Labusch model demonstrates a strengthening effect related to a forest of solute atoms, which is proposed to be more applicable to multi-component solution alloys [67-69] with varied atomic sizes. Assuming a single solute species i, the incremental strength analyzed by Labusch model can be formulated as [64–69]:

$$\Delta\sigma_{\rm ss} = f\mu \left[\left(\frac{\eta_i}{1 + \frac{1}{2} |\eta_i|} \right) + \alpha^2 \delta_i^2 \right]^{\frac{2}{3}} x_i^{\frac{2}{3}}$$
(2.6)

where f and α are dimensionless parameters, μ is shear modulus, η_i , x_i and δ_i are elastic modulus, molar fraction and lattice mismatches of solute species i, respectively. The



Fig. 3 – Schematic of classic solution strengthening mechanisms: (a) Fleischer model and (b) Labusch model.

strengthening effect estimated by the Fleisher model can be expressed as:

$$\Delta \sigma_{\rm ss} = B_{\rm i} X_{\rm i}^{\bar{z}} \tag{2.7}$$

where B_i is a constant related to shear modulus and X_i is the solute content. However, due to variation of composition, atomic size misfit, and interatomic distance, strengthening of HEAs is usually estimated through the Labusch model [68]. Derived from binary system, Toda-Caraballo developed a model named after himself [68,70] to describe solid solution strengthening in HEAs. This model can be formulated as:

$$\Delta \sigma_{\rm ss} = F_{\rm t} Z G B x^n \tag{2.8}$$

where F_t is the orientation factor, Z is scaling factor, G is the shear modulus, n is Labusch dependence and can be determined to be 2/3 for concentrated alloys, and x is solute content. The term B is used to describe the resultant total lattice distortion caused by solute atoms and can be expressed as:

$$B = \left(\xi \sqrt{\{\eta'\}^2 + \{\alpha\delta\}^2}\right)^{\frac{4}{3}}$$
(2.9)

where ξ is number of active slip systems in the alloy. η' represents the elastic modulus mismatch between the solute and solvent. α is a constant indicating whether screw or edge dislocations prevail. A larger α corresponds to an increasing fraction of edge dislocations. δ is the atomic size mismatch.

Solid solution strengthening of FCC HEAs turns out to be comparable to that discovered in conventional alloys, with respect to atomic size mismatch governance and magnitude of activation volume [68,70–72]. However, its BCC counterpart, where deformation is mainly controlled by screw dislocation motion, exhibits both strengthening and softening effects [73,74]. Maresca and Curtin [75] developed a model for solution strengthening specifically in BCC materials, by placing more emphasis on edge dislocation motion. Senkov et al. [76] and and Yao et al. [77], however, have put forward a model incorporating elastic modulus mismatch which can be formulated as:

$$\Delta\sigma b^2 = ZF_3^4 c_3^2 E_L^{-\frac{1}{3}}$$
(2.10)

where Z is dimensionless material constant, b is Burgers vector, E_L is the dislocation line tension, F is the interaction force. Therefore, in general, these three models –Toda-Caraballo, Senkov-Yao, and Maresca– place emphasis on different factors, which are atomic size mismatch, elastic modulus mismatch, and edge dislocation motion, respectively.

Coury et al. [78] studied solution strengthening of NbTaTi, MoNbTaTi, WnbTaTi, HfNbTaTi, CrNbMoTi and CrMoTaTi refractory HEAs by applying these three models, reporting that the Toda-Caraballo could not produce as accurate prediction to refractory HEAs as to FCC HEAs [68,70,79].

2.3.3. Grain size strengthening

Grain boundaries provide pronounced strengthening effect at low temperatures. Traditional grain-boundary strengthening is expressed through the classic Hall-Petch equation:

$$\sigma_{\rm v} = \sigma_{\rm v0} + K_{\rm HP} d^{-0.5} \tag{2.11}$$

where σ_y is yield stress, *d* is the grain size, K_{HP} is the Hall-Petch coefficient. According to this equation, the strength of crystalline materials is inversely proportional to grain size to the power of 1/2. Schuh et al. [80] reported a superior tensile strength obtained in a nanocrystalline Cantor alloy. The nanocrystalline structure provides a high-volume fraction of grain boundaries which act as obstacles to prevent further movement of dislocations. Although significantly enhanced strength can be obtained, an undermined ductility comes along with it. Gangireddy [81] and his co-workers discovered an inversely proportional relationship between strain rate sensitivity and strength when grain cell size is larger than dislocation cell. The grain size is also related to the hardness [82].

2.3.4. Twinning

Twinning is an excellent work hardening mechanism and provides similar strengthening effect as high-angle grain boundaries [68], by impeding the migration of dislocations and by generating additional sites for dislocation nucleation. Interaction between dislocations and twin boundaries has been extensively studied and divided into three routes in FCC metals [69,70]: slip transfer, confined-layer slip and twinning partial slip (Fig. 4). Regarding slip transfer and confined-layer slip, dislocations will rather be converted into partial dislocations and move along twin boundaries and interior domains, or create dislocation junctions, which are also known as hard mode I and II. However, when maximum shear loading is applied along the twin plane, restraints caused by twin boundaries are significantly weakened, which is known as "soft mode". A promising approach to provide a desirable combination of enhanced ductility and strength by introducing twinning has been investigated experimentally [71-77] The formation of twins can be mainly achieved by two routes: mechanical deformation, or appropriate annealing treatment. In the latter, the thickness of annealing twins will be typically larger than that for deformation twins. A large twin width is accompanied by wider spacing between twin boundaries,

thereby weakening the effect of blocking dislocation movement. Thus, a superior hardening effect is always associated with deformation twins due to much smaller distance between twin boundaries. Initiation of mechanical twins is closely dependent on stacking fault energy; in low stackingfault energy (<~20 mJ/m²) materials, twinning turns out to be the dominant deformation mechanism. Certain conditions, such as low strain rate and high deformation temperature can inhibit mechanical twinning. Meyers et al. [92] developed a constitutive description of twinning stress, by which the effect of temperature, strain rate and grain size on twinning stress can be predicted. The twinning stress in FCC metals can be expressed as [78]:

$$\sigma_{\rm T} = \alpha_{\rm T} \left(\frac{\gamma_{\rm SF}}{Gh}\right)^{\frac{1}{2}} \tag{2.12}$$

where α_T is a fitting parameter, γ_{SF} is stacking-fault energy, G is shear modulus and *b* is the Burgers vector. It is often assumed that the temperature and strain rate have only a modest effect on the twinning stress. Twinning occurs when σ_T is equal to or lower than slip stress σ_S , whose relationship with strain rate and temperature can be expressed by the Zerilli-Armstrong equation for FCC metals:

$$\sigma_{\rm s} = \sigma_{\rm G} + C_1 \varepsilon^{\frac{1}{2}} \left\{ exp - \left[-C_2 - C_3 \ln\left(\frac{\dot{\varepsilon}}{\dot{\varepsilon}_0}\right) \right] T \right\}$$
(2.13)

where σ_G is athermal stress, \dot{e}_0 is a reference strain rate, C_1 , C_2 , and C_3 are parameters. In contrast to the twinning stress, the slip stress is closely related to temperature and strain rate. The tendency to form mechanical twins is mainly associated with rivalry between twinning stress and slip stress, while lower slip stress makes it more difficult to form twinning. In high SFE FCC metals, twinning is prohibited at quasi-static loading conditions; however, it is extensively generated under shock loading [94,95]. As for BCC and HCP, increasing strain rate has a comparable effect on twin formation as decreasing temperature [96]. Contrary to FCC, BCC and HCP exhibit lower probability to form twins, due to lack of potential slip systems [93,97]. Despite SFE being of great importance for tendency for twin formation (see Eq. (2.12)), another



Fig. 4 – Interaction between dislocations and twin boundaries in FCC metals [227].

important factor, which is also essential, is the deformation mode. For instance, although deformation twins were found in uniaxially compressed CrMnFeCoNi HEA, none was discovered in tensioned alloys subjected to uniaxial tension. Mostly, decreasing temperature is coupled with lower stacking, which promotes the twinning-induced plasticity (TWIP) effect in FCC metals [98]. Such effect can be tailored by adjusting composition. Deng et al. [99] claimed to successfully design a twinning-induced plasticity non-equiatomic Fe₄₀₋ Mn₄₀Co₁₀Cr₁₀ HEA. By reducing Co, Cr contents and completely removing Ni from CrMnFeCoNi HEA, they discovered a transition of deformation mechanisms from dislocation slip at low strain level to deformation twins plus slip at high strain level and attributed the distinctive behavior at higher strain to reduced SFE in their material. Nano-scale deformation twins, also known as nano-twins have been shown to contribute considerably to strain hardening ability by obstructing dislocation movement and subdividing initial grains into twin-surrounded lamellas [83-91].

2.3.5. Precipitation strengthening

Precipitation hardening derived from second-phase particles enables higher strength by interacting with dislocations. Due to the complex composition of HEAs, studies of precipitation strengthening in HEAs remain inconclusive. Hardening effect can be derived from two routes, namely, by-pass and particle shearing mechanisms, which primary rely on size of the precipitate particles. Regarding precipitation hardening in HEAs, due to the high probability of forming simple solution, this observation is rare. Preceding studies have proposed the presence of second phases in CuCr₂Fe₂NiMn, Al_{0.3}CrFe_{1.5}. MnNi_{0.5} [100] (FCC plus BCC phases), single BCC Al_{0.5}CrFe_{1.5}. MnNi [80], Al_xCrFe_{1.5}MnNi_{0.5} [101], single FCC FeCoNiCr [102] and CoCrFeMnNi with addition of titanium [103]. He et al. [104] also reported incremental strength of over 300 MPa contributed by precipitation hardening in (FeCoNiCr)₉₄Ti₂Al₄ HEAs.

2.3.6. Two-phase hardening

Two-phase hardening can be observed in alloys composed of multiple phases with widely different properties; this can generate a synergy of high strength and excellent ductility. Interestingly, apart from grain-boundary hardening, the nanocrystalline Cantor alloy also promotes phase transformation by providing pathways for atomic diffusion, suggesting that single-phase HEAs can only exist under certain restricted conditions [67]. Phase segregation occurs at grain boundaries and eventually leads to formation of second phase precipitates. Thus, in this case, the strengthening effect caused by a second phase is similar to precipitate hardening. Two of the most common phase transformations observed in HEAs are the FCC to BCC and FCC to HCP transition, which can be activated by either appropriate heat treatment or sufficiently high stress loading. As for its presence in HEAs, superior mechanical properties owing to phase transformations have been proposed for $Co_{20}Cr_{26}Fe_{20}Mn_{20}Ni_{14}$ [105] and FeCoCrNi [106] at cryogenic temperature, CoCrFeMnNi under hydrostatic compression [107] and AlCoCrFeNi [108], etc. Aside from traditional FCC to BCC and FCC to HCP transformation, recent research also uncovered a crystalline to amorphous phase transformation in CoCrFeMnNi alloy

achieved by dynamic compressive [109] and quasi-static tensile [110] tests.

3. Mechanical properties

3.1. Dynamic compression/tensile tests by split-Hopkinson compression bar (SHPB)

The mechanical response at strain-rates ranging from 10^{2} - 10^{4} /s is usually investigated by Kolsky's split-Hopkinson bar system [111,112]. In classic split-Hopkinson compression testing, samples are sandwiched between incident and transmitted bars. When the stress wave propagates through the sample, pulse signals are generated and then captured by strain gages attached at the surface of the bars. Regular stress profiles including engineering stress/strain and strain-rate can be derived from pulse signals through following equations:

$$\dot{\epsilon} = \frac{-2C_0}{L_s} \epsilon_r(t) \tag{3.1}$$

$$\varepsilon_e = \frac{-2C_0}{L_s} \int_0^t \varepsilon_r(t) dt$$
(3.2)

$$\sigma_e = \frac{A_0}{A_s} E_0 \varepsilon_t \tag{3.3}$$

where ε_r is the reflected wave signal, ε_t is the transmitted wave signal, L_s is the length of the cylinder specimen, ε_e is engineering strain, σ_e is engineering stress, C_0 is the wave velocity of maraging steel (~5000 m/s), A_0 and A_s are cross-sectional areas for the bar and specimen, respectively.

Application of the Hopkinson bar is not only limited to compression, but also to torsion and tensile tests. The design of the tension testing setup on split-Hopkinson bar system was proposed by Hauser [113], and Lindholm and Yeakley [114]. Chiddister and Malvern [115] developed a high-temperature setup. The split-Hopkinson bar was properly modified for torsion tests by Baker and Yew [116]. The geometry of specimen used in tests can be classified as continuous and discontinuous. The discontinuous specimens include hat-shaped [117–121], double shear [122–124], single/double notch [125–128], compact forced simple shear specimens [129]. Of the other specimens with continuous geometry, the simplest one is the cylindrical specimen.

Shear localization as an undesired deformation mechanism, leading to catastrophic shear failure of materials, has always been found in dynamically deformed specimens. Such failure mechanism has been extensively investigated in metallic materials [120,130–133] and usually been obtained from high strain-rate loading tests, such as split-Hopkinson compression, ballistic impact, drop-weight tower and shock compression. Initiation of shear localization is mainly associated with shear instability which can be expressed by three different terms [134]:

$$d\tau = \left(\frac{\partial \tau}{\partial \gamma}\right)_{\dot{\gamma},T} d\gamma + \left(\frac{\partial \tau}{\partial \dot{\gamma}}\right)_{\gamma,T} d\dot{\gamma} + \left(\frac{\partial \tau}{\partial T}\right)_{\dot{\gamma},\gamma} dT$$
(3.4)

where first term indicates the work hardening effect, second term represents the strain rate effect on shear stress and third term represents thermal softening. Based on equation (3.4), the occurrence of instability ($d\tau < 0$) is closely related to the work hardening ability, where high strain hardening rate will impede shear instability and poor strain hardening ability will lead to localization. A critical shear strain γ_c is often used as a representative parameter to determine the critical condition to initialize shear localization. This critical strain has been proposed by Culver [134] and can be expressed as following:

$$\gamma_c = \frac{n\rho C_V}{\beta \left| \frac{\partial r}{\partial T} \right|} \tag{3.5}$$

where n is strain-hardening exponent, ρ is density, C_V is specific heat capacity, β is Taylor-Quinney factor. The value of critical strain is always inversely proportional to strain rate

and strain-hardening rate and proportional to strain-rate sensitivity [135,136].

In statically deformed specimens, the heat generation rate is surpassed by the heat diffusion rate. However, in dynamically deformed specimens, due to insufficient time for heat dissipation, heat cannot be successfully dissipated, and drastic temperature rise can take place, which is also known as thermal softening effect. The temperature rise due to adiabatic deformation can be written as:

$$dT = \frac{\beta}{\rho C_{\rm V}} \sigma d\epsilon \tag{3.6}$$

Normally, the stress-strain response is conveniently predicted with a modification of the classic Johnson–Cook equation:



Fig. 5 – (a) True stress vs. true strain curves at different strain rates $(10^{-3}, 10^{-2}, 10^{-1}, 1000, 2600 \text{ s}^{-1})$ (b) Plot of strain hardening rate as function of true plastic strain (c)–(e) Electron backscatter diffraction images and their corresponding (f)–(h) Scanning electron microscope for as-received, quasi-statically deformed and dynamically deformed samples [37].



Fig. 6 – (a) True compressive stress vs. true compressive strain curves of the $Al_{0.3}$ CoCrFeNi high-entropy alloy at different strain rates with corresponding Johnson–Cook fitting curve. (b) Comparison of strain-hardening ability of $Al_{0.3}$ CoCrFeNi HEA with other metallic materials. (c) (d) IPF mapping near deformation tip, showing extensive twins across grains.

$$\sigma = (\sigma_0 + B\varepsilon^n) \left(1 + \operatorname{Clog} \frac{\dot{\varepsilon}}{\dot{\varepsilon}_0} \right) \left(\frac{T}{T_r} \right)^A$$
(3.7)

where A, B, C, n are experimental-based parameters, T and T_r are testing and reference temperatures, respectively. This modified version of Johnson–Cook equation was proposed by Meyers et al. [137], thereby predicting corresponding

temperature as a function of strain, strain-rate, thermal softening, and strain hardening:

$$T = \left[T_r^{1-A} + 0.9(1-A)\varepsilon \frac{1+Cln\frac{\dot{\varepsilon}}{\varepsilon_0}}{\rho C_V T_r^A} \left(\sigma_0 + \frac{B\varepsilon^n}{n+1}\right)\right]^{\frac{1}{1-A}}$$
(3.8)



Fig. 7 – (a) Bright field TEM images of the samples deformed at a strain rate of 3600 s⁻¹ and the corresponding SAED pattern. (b) The HRTEM image of the deformation twins and corresponding SAED pattern [147].

Continuous development of shear localization can lead to the formation of a banded structure, namely adiabatic shear band, accompanied with softening effect [138]. As the strainrate keeps increasing, the thermal softening effect becomes more significant. The evolution of shear bands in thermoviscoplastic materials was studied by Molinari and Clifton [139], and Clifton et al. [135], who presented an idealized model to predict critical condition for shear localization without considering heat conduction, inertia, and elasticity. Ultra-fined/ nanocrystalline grains, which result from dynamic recrystallization (DRX), can always be found within shear bands [118,140-142]. Such ultra-fined/nanocrystalline grains were first reported by Meyers and Pak in titanium [142]. DRX is an important microstructural evolution mechanism associated with temperature rise in local domains. Conventional migrational recrystallization theories are no longer suitable for DRX due to significantly shorter deformation time for grain migration. Therefore, equiaxed recrystallized nano-sized grains are usually observed in shear band. Meyers et al. [144] and Hines et al. [143] pointed out that low angle grain boundaries can rotate when subjected to severe plastic deformation. Thus, occurrence of DRX is mainly attributed to elongated dislocation

cells which subsequently turn into elongated sub grains. Such mechanism is called rotational recrystallization.

3.1.1. Al-Cr-Fe-Co-Ni HEAs

The first ever study on dynamic properties of HEAs was conducted by Kumar et al. [37]. They performed compressive tests on Al_{0.1}CrFeCoNi HEA at both quasi-static (strain rates of 10^{-3} , 10^{-2} and 10^{-1} s⁻¹) and dynamic conditions (strain rate of 2600 s^{-1}), revealing a stronger strain-rate dependence at the latter condition. Such stronger relationship between strain-rate and yield strength is common for low SFE materials. Under high strain rate loading, dislocation planar glide is governed by more active partial dislocation movement in low SFE materials, and the phonon drag effect on dislocation motion becomes more pronounced. They successfully tuned down the SFE (<20 mJ m^{-2}) in CrFeCoNi by introducing Al. The authors attributed the drastic rise in yield strength, as displayed in Fig. 5(a), to the phonon drag effect and intensified partial dislocation movement. With respect to strain hardening, given higher frequency of partial dislocation activity, twinning prevailed and consistently generated a rising strain hardening rate in stage II, as shown in Fig. 5(b, e, h). Hence, Kumar et al. [37]



Fig. 8 – Bright field TEM images and corresponding SADE patterns in: (a) as-cold rolled specimen (b) high temperature annealed and (c) low temperature annealed samples [228].

concluded that the excellent work hardening ability at high deformation velocity was primarily derived from deformation mode governed by twinning, which is similar to other low SFE materials.

The dynamic properties of the Al_{0.3}CoCrFeNi HEA were examined by Li et al. [145]. On the basis of Kumar's research, Li and his co-workers investigated similar Al_xCoCrFeNi systems with a higher atomic fraction of Al (~0.3) to achieve an even lower SFE. This work reported a high resistance to shear localization in this material. The stress-strain curves at varying strain-rates and temperatures are illustrated in Fig. 6(a). The modified Johnson–Cook equation defined earlier (Eq. (3.7)), was applied to quantitatively study the shear instability formation in this alloy. By incorporating the experimentally determined strain hardening and thermal softening parameters, a $\frac{d\tau}{dx}$ with a value of 405 MPa could be determined at a shear strain of 1.1. Thus, shear instability was expected to be suppressed under the conditions at which the specimen was tested. However, Tang et al. [146] observed ultra-fine grains as a result of dynamic recrystallization by applying high-pressure torsion with a pressure of 6.0 GPa for 8 revolutions in Al_{0.3}CoCrFeNi alloy. Thus, the suppression of shear localization may also be related to different deformation mode. In addition, higher shear strains and temperatures are needed for the initiation of shear localization. The retardation of localization was mainly attributed to the extensive formation of mechanical twins, as shown in Fig. 6(c) (d), which benefitted from increase of Al content. As illustrated in Fig. 6(b) which compares HEAs with other conventional metallic alloys, at high strain hardening rate (above 1000 MPa). A significantly higher strain-hardening rate than that in coarse grain pure Al (below 500 MPa at strain over 0.05), was established in Al_{0.3}CoCrFeNi. Thus, this suggests a great potential to be future candidate of impact-resistant materials.

To produce a comprehensive study of dynamic properties of Al_xCoCrFeNi HEAs, Wang et al. [147] carried out dynamic compression tests on Al_{0.6}CoCrFeNi HEA. In contrast to single FCC phase formation in Al_{0.3}CoCrFeNi, the Al_{0.6}CoCrFeNi HEA exhibited a BCC second phase. The dynamic compressive strengths at strain rates of 2800, 3600 and 4000 s^{-1} were 880, 842, 997 MPa, respectively. In addition, this alloy possessed excellent strength and ductility with an ultimate fracture strain approaching 40% at strain rate of 4000 s^{-1} . Therefore, a remarkable work hardening ability was expected to overcome strength-ductility tradeoff. Closely interacting dislocations were observed under TEM, as shown in Fig. 7(a). Under highspeed deformation, the dislocation moving pattern tends to convert from the planar plus wavy glide to wavy glide [148], where predominantly wavy glide is always coupled with the formation of dislocation cells. Operation of wavy glide rendered considerable resistance to dislocation movement, thereby contributing to elevated flow stress of this alloy. In addition, due to disparate stacking fault energy between FCC and BCC phase, profuse deformation twins were observed in the former phase, as shown in Fig. 7(b), while only dislocations were present in the latter phase. Therefore, the operation of multiple deformation modes such as forest dislocation strengthening, solid solution strengthening, and deformation twins leads to remarkable ability to resist the strengthductility trade-off.

The dynamic behavior of Al_xCoCrFeNi with highest Al content was studied by Gangireddy et al. [228]. They examined the dynamic properties of Al_{o 7}CoCrFeNi alloys comprised of FCC matrix plus B2 lamellar structure, which has been discovered by a preceding study [149]. By conducting high (1100 °C) and low (580 °C) temperature annealing treatments, distinctive properties were revealed: while high temperature annealed alloy possessed higher work hardening ability, the low temperature annealed alloy had a higher yield strength. The low temperature treatment contributed to inferior ability for dislocation recovery compared to one annealed at high temperature, thereby promoting formation of ordered L12 precipitation, as shown in Fig. 8(c). The difference in mechanical properties between B2 and FCC phases leads to inhomogeneous deformation, whereby the internal domain of B2 provides higher dislocation storage capability [150,151] and the softer primary FCC phase can accommodate greater plasticity. In general, the distinct dynamic behavior is mainly the result of different degrees of dislocation recovery raised by different annealing temperatures.

The first investigation on dynamic properties of AlCoCrFeNiTi_x was reported by Jiao et al. [152]. Unlike classic AlCoCrFeNi with modification only in Al content, they introduced Ti. The dynamic compressive results of Ti₀, Ti_{0.2}, and Ti_{0.4} alloys are revealed in Fig. 9(a-c), suggesting that the yield strength was proportional to the atomic ratio of Ti, where Ti₀ alloy exhibits the lowest yield strength of 1546 MPa and $Ti_{0.4}$ exhibits highest yield strength of 2090 MPa. The enhanced compressive strength was attributed to the larger atomic radius of Ti atoms leading to severe lattice distortion and significant enhancement of the solid solution strengthening effect. Their results are summarized in Fig. 9(d) and compared with Nibased superalloys and Bulk Metallic Glasses, AlCoCrFeNiTi_x exhibited a high ultimate strength while maintaining a relatively large strain to failure, which is attributed to solid solution strengthening, phase variation, thermally activated mechanisms [153–155], and the phonon drag effect.

Besides the conventional AlCoCrFeNi system that only contains metallic elements, Wang and his co-workers [156] designed a (Al_{0.5}CoCrFeNi)_{0.95}Mo_{0.025}C_{0.025} HEA. Preceding research [157,158] reported that the addition of Mo and C could enhance mechanical properties of Al_{0.5}CoCrFeNiMo_x HEAs by forming a (Cr, Mo)-rich σ phase and promoting precipitation hardening. The yield strengths of (Al_{0.5}CoCrFeNi)_{0.95}Mo_{0.025}- $C_{0.025}$ deformed at strain rates of 550, 1300, 2200 and 3000 $\ensuremath{\text{s}}^{-1}$ and were 754, 878, 958, and 1057 MPa, respectively. In comparison with $Al_{0.3}$ CoCrFeNi HEA [145], a much higher strain hardening rate was observed in (Al_{0.5}CoCrFeNi)_{0.95}Mo_{0.025}-C_{0.025}, but pronounced shear banding was found. This contradictory phenomenon was ascribed to dissolved Mo atoms that slow down thermal diffusion. Hereafter, thermal softening was benefitted from hindered heat diffusion, and promoted shear banding.

3.1.2. Co-Cr-Fe-Mn-Ni HEAs

Surprisingly, the classic CoCrFeMnNi (Cantor alloy) is the first ever discovered HEA but is not the first HEA to be examined dynamically. Its dynamic compressive properties were investigated by Park et al. [159] in 2017. The yield strength under dynamic conditions was reported to be 590, 650,



Fig. 9 – Dynamic compressive engineering stress–strain curves of Ti_0 (a), $TiO_{0.2}$ (b), and $Ti_{0.4}Ni$ Al (c) alloys. (d) Ultimate strength as a function of fracture strain for various materials, under dynamic loading [152].

680 MPa at strain rates of 3000, 3500, 4700 s^{-1} , respectively. Fig. 10 shows the Inverse Pole Figure (IPF) maps in the plane of compression and transverse directions for quasi-statically dynamically deformed specimens. Greater inand homogeneity of strain can be observed in the latter. The strain hardening rate derived from the dynamic stress vs. strain curve, as shown in Fig. 10(b), can be divided into four stages (A, B, C, D), corresponding to steep drop of hardening rate caused by the elastic-plastic transition, slow linear decrease caused by dislocation glide coupled with dynamic recovery [160], steady hardening rate related to formation of mechanical twins, and, finally, steady decrease of hardening rate. Profuse twins were observed under both quasi-static and dynamic conditions. A rather higher density of twins was observed in dynamically deformed samples; this was similar to what was reported in Al_{0.1}CrFeCoNi alloy [37]. As the alloy was further strained, a gradually decreasing work hardening rate was overcome by thermal softening, which led to shear instability, as shown in Fig. 10(e, f, i, j). Contrary to conventional FCC alloys with high activation volume, such as Cu (200–2000 b^3), α -brass (200–400 b^3), an exceptionally low activation volume (~10 b³), indicative of extensive twinning, was measured for CoCrFeMnNi alloy. Thereby, profuse twins contribute to the remarkable work hardening ability of this alloy, resembling Al_{0.3}CoCrFeNi [145].

Li et al. [161] have also examined the dynamic properties of CoCrFeMnNi HEA, but prepared by a different method, spark plasma sintering (SPS) method, and focusing on adiabatic shear localization. Unlike in the arc-melting method, alloys prepared by SPS are likely to contain voids, leading to residual porosity and lower strength. They obtained a high strain hardening rate (~1100 MPa) at a strain of 0.2 when the alloy was deformed at strain rate of 1600 s⁻¹. Moreover, through imposed localized shear deformation in hat-shaped specimens, they successfully established the critical shear strain (~7) to induce shear banding (Fig. 11(b)). The retardation of shear localization suggested a strong work hardening ability, which involved twinning and strong dislocation planar glide. Within the shear band, they observed a similar subgrained structure to the one reported by Park [159], as shown in Fig. 11(c). This grain refinement is due to occurrence of dynamic recrystallization. Fig. 11(d) shows that, compared with titanium [137], Ti-6Al-4V alloy [162], AISI 1006 steel [47], brass [47] and copper [120], the Cantor alloy has the best ability to resist the softening effect in the temperature range of 200 K-1000 K. In addition, some nanotwins were found within those nanograins, indicating that they could still exhibit plasticity at elevated temperature.

Wang et al. [163] provided an interesting insight to dynamic properties of as-cast CoCrFeMnNi HEA. Without homogenization treatment, this alloy showed a dendritic structure. At high strain rate, unlike preceding studies on annealed alloys [159,161], as-cast alloy displayed unique "serration behavior", which can be seen as tooth-like flow stress curves in Fig. 12(a). Zhang et al. [164] reported that the serration behavior is more pronounced at elevated temperatures and under dynamic deformation. They ascribed this continuously fluctuating stress to the formation of a heterogeneous structure, namely, deformation bands, as



Fig. 10 – (a) Engineering stress vs. engineering strain curves under dynamic compression (b) Strain hardening rate as a function of true plastic strain. IPF maps one the plane of compressive direction (c)–(f) and transverse direction (g)–(j) (c, d, g, h) at 10^{-4} s⁻¹ (e, f, i, j) at 4700 s⁻¹ [159].



Fig. 11 – (a) Stress vs. strain curves at different strain; (b) shear stress as function of shear strain and corresponding temperature rise within the shear band; (c) TEM micrograph showing recrystallized grains with dislocations and twins; (d) Critical shear strain for shear localization for multiple materials (Cantor alloy, Ti, Ti–6Al–4V alloy, brass, copper and 1006 steel); (e) Twin boundaries within recrystallized grain [161].

displayed in Fig. 12(b and c). Deformation bands in this case generate the Portevin-Le Chatelier effect [165–168]. The serrated flow stress was a result of interaction between moving dislocations and deformation bands and could become more evident at higher strain rate due to larger amount of such heterogeneous structure, as illustrated in Fig. 12(c and d). Besides the alloy prepared by the traditional arc melting method, Wang et al. [169] also extended their work to serration behavior of CoCrFeMnNi HEA prepared by SPS. In contrast to their previous work [163], where serrated flow stress resulted from formation deformation bands, here they ascribed it to secondary Cr-rich oxide phase. The



Fig. 12 – (a) Compressive true stress vs. true strain curves for CoCrFeMnNi HEA at different strain rates. (b) The Euler plus band contrast image of the deformed specimen at the strain rate of 4600 s⁻¹. Bright field TEM micrograph of specimens deformed at strain rates of (c)1700 s⁻¹, (d)4600 s⁻¹ [163].

dispersed Cr-rich particles within matrix provided a strong pinning effect for dislocations movement by generating the Portevin-Le Chatelier effect (Fig. 13). The role of secondary particles was quite comparable to deformation bands, in terms of slowing down dislocation movement. Each time a



Fig. 13 — The bright field TEM micrograph showing pinned dislocations at the boundary of secondary phase particles [164].

dislocation was pinned and unpinned by second phase particle it causes an increase and decrease in stress [164].

On the basis of Park's and Li's work, Tsai et al. [54] proposed a transition of deformation mode when CoCrFeMnNi alloy was subjected to a high strain rate of 9000 s⁻¹. As mentioned before, dynamic deformation of CoCrFeMnNi alloy seems to be governed by dislocation glide. However, at a strain rate of 9000 s⁻¹, kinked twins were observed, suggesting a deformation mode dominated by twinning, as shown in Fig. 14. Such alteration of deformation mode, generating an enhanced strain hardening rate coupled with rising strain rate, makes further study of CoCrFeMnNi alloy deformed at ultra-high strain rate (shock compression) more interesting.

To provide further insight to the effect of adiabatic heating on mechanical response of CoCrFeMnNi, Soares et al. [170] reported that the Taylor-Quinney coefficient was not equal to 0.9 as was previously assumed for the determination of temperature rise at high strain rate for metallic materials. The Taylor-Quinney coefficient is a parameter denotes efficiency of plastic work converted into heat. This coefficient can be classified into two components: integral beta (β_{int}) and differential beta (β_{diff}), which correspond to how much strain energy is converted into heat, respectively; they can be expressed as:



Fig. 14 – (a) TEM micrograph of 40% strained specimen at a strain rate of $3 \times 10^3 \text{ s}^{-1}$ (b,e) Bright field images of deformation twins present at the strain rate of $6 \times 10^3 \text{ s}^{-1}$ and $9 \times 10^3 \text{ s}^{-1}$, respectively (c,d) dark-field image of bundle of twins in sample deformed at strain rate of $6 \times 10^3 \text{ s}^{-1}$ (f,g) Twin–twin interaction with kinked structures in specimen deformed at strain rate of $9 \times 10^3 \text{ s}^{-1}$ with 40% strain [54].

$$\beta_{\rm int} = \frac{\rho C_{\rm p} \Delta T}{\int dW_{\rm p}} \tag{3.9}$$

$$\beta_{diff} = \frac{\rho C_p \dot{T}}{\dot{W}_p} \tag{3.10}$$

where C_p is heat capacity, dW_p is the increase in plastic work, ρ is density, T is temperature, \dot{T} and \dot{W}_p are differentials of plastic work and temperature with respect to time. Fig. 15(a and b) reveals β_{int} and β_{diff} as a function of strain at different strain rates. The authors discovered that the adiabatic heating in specimens deformed at 1,750, and 2800 s⁻¹ were quite similar and their corresponding β exhibited certain degree of plastic strain dependence.

Apart from most common dynamic test performed by high strain rate compression, Qiao et al. [171] investigated the strain rate dependent microstructural evolution of CoCrFeMnNi via split Hopkinson tensile bar (SHTB) at a strain rate of 7000 s⁻¹. Distinct fracture behaviors were found under different loading conditions, as shown in Fig. 16(a and b), indicating a typical ductile and ductile plus brittle fracture mode in statically and dynamically deformed specimens, respectively, which is suggestive of a transition of fracture mode. Such transition of fracture mode has also been observed in Al_xCrMnFeCoNi alloy by Cao and his co-workers [172]. Similar to CoCrFeMnNi subjected to dynamic compression [159,161], dislocations and twins prevailed in dynamically tensioned specimens, as illus-



Fig. 15 – (a) Integral β and (b) differential β diff as a function of true strain for the CoCrFeMnNi high-entropy alloy under different strain rate (1 s⁻¹, 750 s⁻¹, 2800 s⁻¹) [170].



Quasi-static

Fig. 16 - SEM micrographs of the fracture morphology for (a) quasi-statically deformed and (b) dynamically deformed specimens. EBSD micrograph of (c) statically deformed specimen at strain of 0.34 and (d) dynamically deformed specimen at strain of 0.57 [171].

trated in Fig. 16(c and d), in contrast to the statically deformed one. The differences in deformation structures cause alteration of fracture from purely ductile to a mixture of ductile/brittle at high strain rates.

Amorphization is most often obtained by two routes in metallic materials: rapidly decreasing temperature to below glass transition temperature Tg or introducing structural/ chemical disorder. Recently, a large amount of emphasis has been put on solid-state amorphization (SSA) achieved by severe plastic deformation [173,174]. It is known that accumulation of defects can eventually lead to amorphization. Amorphization in HEAs is being widely studied to broaden their applications in the field of advanced nuclear materials, corrosion and oxidation resistant materials, but is rarely observed experimentally.

Outside of conventional deformation modes of dislocation glide, twinning, microbanding, Zhao et al. [109] reported the formation of an amorphous phase in Cantor alloy achieved via dynamic compressive tests, suggesting an unprecedented deformation-induced amorphization. Associated with amorphization, vast amounts of crystalline defects such as stacking faults, twins and HCP phases have also been observed in dynamically deformed specimens, as shown in Fig. 17(a-e). In order to activate the FCC to HCP phase transition, the required driving force $G_{HCP} - G_{FCC}$ can be established as a function of stacking fault energy $\gamma_{\rm SF}$:

$$\frac{\gamma_{\rm SF}}{2b} = G_{\rm HCP} - G_{\rm FCC} \tag{3.11}$$

Such driving force needs to be fulfilled by generating defect energy G_d , which can be expressed as following:

$$G_d \sim \frac{\rho G b^2}{2} \tag{3.12}$$

where ρ is defect density. Combining Equation (3.11) and equation (3.12), they predicted a critical value of $\rho \sim 8 \times$ $10^{15}m^{-2}$, which was achievable by dynamic loading, to initiate phase transformation. With ongoing plastic deformation, the severely deformed FCC and HCP can be further converted into



Fig. 17 – (a–e) TEM micrograph of the CoCrFeMnNi HEA subjected to dynamic compression/shear, showing the formation of deformation twins, HCP and amorphous phases. (f) Microstructural evolution of CoCrFeMnNi HEAs subjected to increasing stress and strain rate [109].

an amorphous phase, due to continuously rising defect concentration. In addition, the amorphous domains located favorably at the intersections of HCP and twin packets. A relationship between shear strain and volume fraction of amorphous phase was accordingly proposed [109]:

$$f_a = K[1 - \exp(-\alpha\gamma)]^n \tag{3.13}$$

where K is a constant related to various factors (stacking fault energy, volume fraction of HCP martensite in the deformation packets, and so on), $1 - \exp(-\alpha\gamma)$ is a term that represents fraction of deformation bands and γ is shear strain. Accordingly, the rising defect density was coupled with increasing fraction of deformation bands, which had previously been found in as-cast Cantor alloys [163], leading to the propagation



Fig. 18 – (a–c) EBSD local misorientation maps of the $Cr_{26}Mn_{20}Fe_{20}Co_{20}Ni_{14}$ HEA undergoing strain rates of (a) 1000 (a), (b) 2000 and (c) 3000 s ⁻¹, respectively. The twin boundaries are indicated by red lines. (d) Estimated dislocation density (ρ) and volume fraction of twin boundaries as function of strain rates [175].

of amorphization. The authors conclude that as flow stress increased, in addition to the common deformation mechanisms (dislocation motion, twinning, phase transformations), previously reported in CoCrFeMnNi HEA, a newly discovered amorphization was added. This renders a remarkable strain hardening ability under dynamic conditions; the corresponding microstructural evolution is plotted in Fig. 17(f).

Other than investigations on equiatomic CoCrFeMnNi HEA, a systematic study on dynamic properties of non-equiatomic Cr₂₆Mn₂₀Fe₂₀Co₂₀Ni₁₄ alloy was performed by Jiang et al. [175]. Preceding studies [176] suggested that this alloy possessed a low stacking fault energy of 24 mJ m⁻² and synergy of excellent strain hardening ability and considerable ductility (maximum strain: 73%) at quasi-static strain rate. They subjected the alloy to dynamic compression at three strain rates (1000, 2000 and 3000 s^{-1}). Noteworthy that continuously increasing flow stress was obtained at the three strain rates (1000, 2000 and 3000 s^{-1}), indicative of a remarkable strain hardening ability at various loading conditions. The local misorientation map shown in Fig. 18 reveals that the volume fractions of twins and stacking faults increase with strain rate. The highest temperature rise caused by adiabatic deformation was estimated to be 64.7 K, which was far from the recrystallization temperature (640 K) of this alloy. Thus, no shear band was observable under TEM. In addition, a smaller activation volume (11 b^3) was found in the present alloy, compared with CoCrFeMnNi HEA (76.8 b³ [177]). Profuse twins and twin-dislocation reactions were attributed to such low activation volume, which showed a good agreement to previously reported value for twin-dislocation reactions $(1-100 b^3 [177])$. Moreover, the low stacking-fault energy of this alloy contributes to restricted cross-slip and dislocation climb and increasing density of stacking faults, thereby promoting dislocation and stacking fault strengthening. Therefore, the authors conclude that increasing strain rate from $1000 s^{-1}$ to $3000 s^{-1}$ facilitated an evolution of deformation mechanism from dislocation glide to stacking faults plus nano-twinning, then eventually to dense dislocation arrays plus nano-twin bundles, which resemble the CoCrFeMnNi HEA deformed at increasing strain from $3000 s^{-1}$ to $9000 s^{-1}$ [54]. This synergy of multiple deformation mechanisms guarantees a retained hardening ability and retards shear localization.

Accordingly, the study of the classic Cantor alloy has been extendedly to strain rates ranging from 1000 s^{-1} to 9000 s^{-1} . A relevant question is: how would the addition of a sixth metallic element change properties of Cantor alloy at high strain rates? Cao et al. [172] examined the dynamic tensile behavior and developed a microstructural evolution of Al_xCrMnFeCoNi (x = 0, 0.4, 0.6) HEAs. It was revealed that the dynamic tensile strength was proportional to the fraction of Al, which is mainly due to the formation of B2 precipitates (Fig. 19(a and b)). The interphase boundaries between B2 precipitates and FCC matrix, shown in Fig. 19(c)–(e), capture large amounts of dislocations and provide a strong boundary strengthening effect. Increasing



Fig. 19 – Dynamic compressive stress strain curves of (a) $Al_{0.4}$ CrMnFeCoNi (b) $Al_{0.6}$ CrMnFeCoNi alloys. (c) TEM micrographs of dislocation configuration of (c–e) $Al_{0.6}$ CrMnFeCoNi tested at strain rate of 10^{-3} s⁻¹ (c,d,e) and 1300 s⁻¹ (f,g,h) [172].

the fraction of Al not only promoted formation of precipitates but also restrained grain growth of the matrix phase, leading to considerably refined grains in Al_{0.6}CrMnFeCoNi. In comparison with CoCrFeMnNi [178], Al_{0.4}CrMnFeCoNi exhibited small activation volume (10–24 *b*³), close to the one reported for Cr₂₆-Mn₂₀Fe₂₀Ni₁₄ [175], indicative of intensive dislocation motion over obstacles raised by solute atoms. In addition, the Al_{0.4}-CrMnFeCoNi revealed a higher strain rate sensitivity (m = 0.039–0.048) in comparison with Al_{0.6}CrMnFeCoNi, which resulted from the positive relationship between Al content and activation volume. Thus, the authors concluded that the high strength of Al_{0.6}CrMnFeCoNi was chiefly due to the multiple strengthening mechanisms, including precipitation hardening, solid solution strengthening, and boundary strengthening.

3.1.3. Refractory HEAs

Although FCC HEAs have received a vast amount of attention due to their excellent mechanical properties at relatively low temperatures, the inherently low melting points of their constituent elements make them less desirable for application in a wider temperature range. For instance, many aerospace materials require exposure to extremely high temperatures and require sufficiently high strength to bear external dynamic loading. For the time being, Ni-based superalloys are widely



Fig. 20 — (a) The compressive true compressive stress vs. true compressive strain curve of RHEA

Ti₂₀Hf₂₀Zr₂₀Ta₂₀Nb₂₀. (b) IPF map showing the localized deformation in area adjacent to shear band. (c) Magnified grain boundary map of the area marked by A in (b). (d) KAM map of (c) indicating lattice distortion and lenticular structures in interior and the neighboring area of the shear band [179].

used in critical components of modern-day jet engines. Novel refractory high-entropy alloys (RHEAs) are being considered as a potential replacement for superalloys due to their excellent resistance against softening at elevated temperature. Dynamic properties of $Ti_{20}Hf_{20}Zr_{20}Ta_{20}Nb_{20}$ with average grain size of ~200 μ m were investigated by Dirras et al. [179] using a Hopkinson bar system with strain rate up to ~3.4 \times 10³ s⁻¹. They are among the first to investigate dynamic properties of BCC HEAs

comprised of refractory elements. The stress-strain curve at high strain-rate condition illustrated in Fig. 20(a), shows a progressively decreasing flow stress at strains >0.05, denoting the occurrence of instability and an intensive softening effect after yielding. Formation of shear bands was clearly revealed in Fig. 20(b), with a highly distorted neighboring needle-like features indicated by black arrows with misorientation profile displayed in Fig. 20(c). Such needle-like structure is considered as thin shear bands at internal areas of grains. The lamellar structure was previously defined as a combination of twins and shear bands [180]. However, a misorientation angle lower than 60° has been revealed across the boundaries of needle-like structures, as shown in Fig. 20(c), suggesting other possibilities than twin boundaries and demanding additional work to provide additional clarification. Although preceding studies reported that the occurrence of shear band is always coupled with DRX. The formation of DRX in the present alloy is strongly confined due to slow diffusion rate in Ta and Ta-W alloys [181]. Therefore, the authors proposed that the DRX is unlikely to form in the $Ti_{20}Hf_{20}Zr_{20}Ta_{20}Nb_{20}$ alloy with such short duration of deformation time.

On the basis of Dirras's work [179], the dynamic behavior of $Ti_{20}Hf_{20}Zr_{20}Ta_{20}Nb_{20}$ RHEA at elevated temperature was further investigated by Hu et al. [182]. The true stress–strain curves at various strain-rates and temperatures are displayed in Fig. 21. As shown in Fig. 21(a), the strain rate hardening effect at room temperature was not pronounced under dynamic conditions. At elevated temperature (673 K), however, the yield strength of the alloy exhibited stronger dependence to strain rate. Such behavior could be explained in terms of strain-rate sensitivity, *m*:

$$m = \frac{\partial \sigma}{\partial ln\dot{e}} = \frac{\sigma_2 - \sigma_1}{\ln(\dot{e}_2 - \dot{e}_1)}$$
(3.14)

 σ_2 and σ_1 are the flow stresses in compression at different strain rates of $\dot{\epsilon}_2$ and $\dot{\epsilon}_1$. A stronger strain rate dependence of stress occurs at elevated temperature. Such behavior was chiefly associated with more intensive viscous drag effect [183,184]. The high strain-rate sensitivity established at elevated temperature suggests enhanced resistance to shear localization. In addition, similar to conventional alloys, the yield strength and flow stress was inversely proportional to the temperature increase. Such behavior is often found in BCC metals and is mainly due to a change of Peierls-Nabarro stress, which is the predominant factor controlling screw dislocation propagation. One can also see that the Ti₂₀Hf₂₀Zr₂₀Ta₂₀Nb₂₀ RHEA retains a relatively high strength at elevated temperature. The temperature sensitivity parameters η_a which revealed the thermal softening effect of materials are listed in Table 1, and are expressed as:

$$\eta_a = \left| \frac{\sigma_2 - \sigma_1}{T_2 - T_1} \right| \tag{3.15}$$

where σ_1 and σ_2 are stresses at temperature T_2 and T_1 , respectively. The authors conclude that the enhanced viscous drag effect was coupled with elevated temperature, and was reflected in rising strain-rate sensitivity of dynamic tests at 673 K.



Fig. 21 – Dynamic compressive true stress true strain curves of $Ti_{20}Hf_{20}Zr_{20}Ta_{20}Nb_{20}$ RHEA samples at various strain rates and temperatures [182].

Table 1 — Variation of temperature sensitivity η_{α} with strain rate [183].			
True strain	Strain rate	Temperature(k)	Temperature sensitivity, η_a (Mpa/K)
0.03	$1600 s^{-1}$	293 K—473 K 473 K—673 K	0.83 0.79
	$2600s^{-1}$	673 K—873 K 77 K—293 K	0.54 2.19
		293 K—673 K	0.55

Particular HEA systems have been investigated extensively, for example, the Al-Cr-Fe-Co-Ni and Co-Cr-Fe-Mn-Ni. Based on the dynamic properties examined by Hopkinson pressure bar, most HEAs comprised of three d-transition elements exhibited both high strength and high strain hardening ability when subjected to dynamic loading. Despite evident shear banding observed widely in dynamically deformed HEA, the critical shear strain to induce shear localization was significantly deterred. Further microstructure characterization suggested that although traditional approaches to improve work hardening ability, like twinning and dislocation multiplication, were detected in HEAs, their presence could be altered by local chemical order, second phase formation, solution atoms and so on. RHEAs, however, provide another path to design high-temperature alloys. With the addition of refractory elements, their strength at elevated temperatures was well maintained.

3.2. Ballistic impact response

Understanding of dynamic behavior at strain rates ranging on the order of 10^5 s^{-1} , beyond the usual range of Hopkinson bars, is mainly achieved by ballistic impact tests [185]. In general, ballistic impact test is accomplished by high-speed collision between projectile and target materials. This idea was first proposed by Taylor [186] in 1948 by accelerating a flat-head cylindrical projectile to a rigid target. This test is also known as Taylor impact test. It is very useful in obtaining important information on the dynamic behavior of materials. The plastic deformation generated during impact is concentrated in the impact region, which is which is separated from undeformed material. In Taylor's analysis, the dynamic yield strength can be estimated by the following equation:

$$\sigma_{y} = \frac{\rho v_{0}^{2}}{2} \cdot \frac{l_{0} - l_{f}}{l_{0} - L_{f}} \cdot \frac{1}{\ln(l_{0} - l_{f})}$$
(3.16)

here ρ is cylindrical rod density, l_0 is initial length of the rod, v_0 is projectile flying velocity, l_f is final undeformed section length, L_f is the overall final length of the specimen.

During this process, the leading elastic compressive wave is followed by the plastic compressive wave [187]. When this amplitude exceeds the yield strength of the material, severe deformation in the form of radial motion outward away from the specimen axis occurs.

Distinctive shear condition can take place with different geometries of projectile, such as cylindrical, conical, and blunt. Investigations of impact response have been conducted on various metallic materials, including 7039 aluminum alloy [188], Al_{0.1}CoCrFeNi [189] and Fe₂₇Co₂₄Ni₂₃Cr₂₆ [190]. However, due to substantial amounts of variants and uncertainties in ballistic impact tests, the experimental results can vary significantly from one source to another.

The ballistic response of Mn-free $Fe_{27}Co_{24}Ni_{23}Cr_{26}$ alloy, as one of the variants close to CoCrFeMnNi HEA, was examined [190]. As shown in Fig. 22, EBSD mapping revealed the deformation response in three different domains: (a) undeformed, (b) intermediate deformation, and (c) severe deformation. In the domain prior to deformation, only annealing twins were discovered, while nano-sized twins could not be distinguished (Fig. 22(a, d)). As shown in Fig. 22(e) (f), specimens subjected to both intermediate and heavy deformation revealed twin–twin interaction, while the latter revealed formation of nanotwin bundles and interaction between deformation nanotwins due



Fig. 22 – (a) EBSD IPF maps and TEM images of $Fe_{27}Co_{24}Ni_{23}Cr_{26}$ alloy (a, d) non-deformed area (b, e) intermediate deformation area and (c, f) severely deformed area. Annealing nanotwins are revealed in (d, e, f), and bundle of deformation nanotwins (DNTs) are revealed in (e, f) [190].

to increasing twin density. Therefore, multiplication of twins significantly confines dislocation motion and deters the activation of shear localization.

Muskeri et al. [189] reviewed the ballistic impact response of single-phase Al_{0.1}CoCrFeNi HEA, which is one of the first HEAs examined dynamically. By applying EBSD analysis, distinctive microstructures were observed in areas with different distances to the crater wall, as shown in Fig 23. In the condition of partial penetration, both low angle misorientation (~2-3°) and high angle misorientation (~60°) deformation corresponding to deformation microbands and deformation twins, respectively, were revealed in Fig 23(a). Deformation microbands were formed as a consequence of planar dislocation arrays that had been previously observed in FeNiMnAlCr alloy [191], Cr₁₀Mn₅₀Fe₂₀Co₁₀Ni₁₀ alloy [192], and $Al_4Mo_4Nb_8Ti_{50}Zr_{34}$ alloy [193]. They can provide similar effect as strain hardening due to mechanical twins through dense dislocation boundaries. One can see that the domains close to the crater wall exhibited much higher hardness (by a factor of 2.5) than the undeformed domains, as illustrated in Fig. 23(e). In addition, there was no evident formation of shear bands at low-velocity impact, indicative of the excellent strain hardening ability associated with twins and microband formation. However, increasing impact velocities (to 739 m s⁻¹ and 994 m s⁻¹) were coupled with pronounced formation of shear bands and recrystallized grains, suggestive of dominant thermal softening over hardening effect. In full penetration condition, the highest hardness (~5 GPa) was measured at the domain near the entry and exit holes.

Moreover, Choudhuri et al. [194] reviewed the ballistic performance of a series of nickel-based HEAs, reporting an excellent classic strength-ductility tradeoff. They fabricated AlCoCrFeNi_{2.1} HEA with dual-phase (FCC-B2 phases) via thermomechanical processing. The ballistic testing was performed with tungsten-carbide projectiles. Varying impact velocities were applied to investigate deformation features in three different conditions (partially penetrated/plugged/fully penetrated) as shown in Fig. 24(b-d). The EBSD micrograph of partially penetrated specimen (Fig. 24(f)), displays a restrained crack growth path within B2 lamellar domains. In addition, shear bands were also observed, denoting severe shear deformation in local domains. Such phenomenon was likely associated with residual stresses provided by crack formation. However, with increasing striking velocity (higher than plug condition), shear localization becomes the dominant failure mode and guides crack growth, similar to other HEAs mentioned earlier.

Instead of studying the impact resistance, the penetration performance, essential to anti-armor application, was examined by Xing-Fa Liu et al. [195]. Besides HEAs only comprised of 3 d-transition metals, such as Fe, Co, and Ni, the emphasis has shifted to HEAs containing refractory elements such as Mo, Ta, and W. A "self-sharpening" multiphase WFeNiMo HEA fabricated by electromagnetic levitation melting technology was investigated by Liu et al. [195]. There is a constant competition between ballistic ("bullet-proof") materials and materials to defeat armor. The concept of "self-sharpening" is associated with the specific increase in the depth of the



Fig. 23 – (a) Microstructure as a function of distance from the crater wall for partially penetrated $Al_{0.1}$ CoCrFeNi HEA plate showing highly deformed grains, microbands, and undeformed grains; (b) IPF map of the domain in the black box marked in (a); (c) Magnified IPF map of the selected domain in green box marked in (b) showing formation of deformation twins; (d) misorientation profile along the arrow shown in (c), indicating boundaries with 60° misorientation that is classic trait of twinning; (e) nano-indentation hardness contour map indicating maximum hardness at the domain close to crater wall [189].

penetration (DOP) by sharpening the penetrator rod into acute shape. Conventional materials used to fabricate rod penetrators are tungsten heavy alloys (WHAs) and depleted uranium alloys (DUAs). While the WHA are relatively less effective due to high penetration resistance plus decreased DOP, the DUA penetrators are widely banned because of many relative long-term health problems. In DUAs the sharp tip is ensured by sequential shear-band formation. On the other hand, WHAs tend to form mushroom shape upon impact, significantly compromising penetration performance. Therefore, novel metallic alloys, that can efficiently penetrate through armors, are desperately in need. Preceding studies proposed applying severe plastic deformation, such as cold torsion [196], swage molding [197], and hot-hydrostatic extrusion [198,199]. Other researchers claimed a "self-sharpening" achieved by inducing shear localization [200-206]. Contrary to traditional uniaxial dynamic compression tests on cylindrical specimens, whose stress conditions are unfavorable for shear band formation [207], the unique shape of rod penetrator endows desired conditions for shear localization to take place. The shear failure localized at the edge of the penetrator heads during deformation consistently pulls off and sharpens them, leading to high depth of penetration. In addition to the softest FCC matrix phase, tungsten BCC and Fe/Ni concentrated μ phases can provide stiffer and stronger

properties of the present alloy. Substantially different properties among three phases enabled the excellent synergy of high strength and ductility under dynamic conditions, with yield strength, fracture strength and fracture strain equal to 1200 MPa, 1900 MPa, 0.28, respectively. As for the ballistic performance, in contrast to conventional 93 W alloy, the WFeNiMo HEA exhibited a superior penetrating performance, represented by the acute head shape, as shown in Fig. 25. The achievable "self-sharpening" ability was mainly attributed to unique microstructure of WFeNiMo HEA, which promoted higher susceptibility to shear localization. As the shear bands continue to develop, the deformed parts are peeled off, resulting in the sharp shape of the penetrator. μ phase plays an essential role in shear band formation by promoting inhomogeneous deformation and higher strain gradients. The shear localization mechanism proposed in Fig. 26 (c-g) illustrates that the distribution of μ precipitates contributes to the intense inhomogeneous deformation and strain gradient.

3.3. Shock compression response

The shock response of materials is an indication of their mechanical properties at ultra-fast deformation speed. Via explosive detonation, projectile impact, high-power pulsed-laserdriven shock compression, ultra-high-speed deformation can



Fig. 24 - (a-d) Optical micrographs showing (a) sections of the impacted EHEA plate along with the WC projectile. Cross sectional areas of specimens with different impact conditions: (b) partial penetration, (c) plugged condition, (d) full penetration. (e) BSE (back scattered electron) image, and corresponding (f) EBSD IPF mapping for area within black box, showing formation of crack along FCC/B2 interphase (indicated by white arrow) and shear band (indicated by yellow arrow).

be generated. The propagation of a plane shock is denoted by the (U, u) Hugoniot curve (U is shock velocity and u is particle velocity), which is one of the primary methods to investigate shock compression response. In addition, the yield strength derived from the curves which is also known as HEL (Hugoniot Elastic Limit), which represents strength at the point when relaxation



Fig. 25 – (a) Depth of penetration of WFeNiMo rod and 93 W rod versus kinetic energy per volume calculated by $\frac{\partial V^2}{2}$, with images of the retrieved remnants, respectively; longitudinal sections of medium carbon steel targets impacted by (b) a WFeNiMo penetrator and (c) a 93 W penetrator, with SEM micrographs of the remnant in the corresponding insets, respectively [195].

wave has not arrived, and target is only deformed along lateral direction. The yield strength in shock compression can measured through von Mises yield criterion [35]:



Fig. 26 – (a) SEM and (b) TEM images showing microstructures of shear band. (c-g) Schematic drawing showing the formation mechanism of shear band in WFeNiMo alloy; (c) Undeformed microstructure with low dislocation density; (d) Formation of strain gradient due to geometrically necessary dislocation (GND); (e) Formation of elongated subgrains due to cotinuously rising dislocation density; (f) Elongated subgrain evloves into equiaxed grains and rotates to accommodate further deformation; (g) Onset of shear band cc [194].

$$\sigma_{\rm Y} = \sigma_{\rm H} \left(\frac{1 - 2\nu}{1 - \nu} \right) \tag{3.17}$$

where v is Poisson's ratio, σ_H is elastic limit under uniaxial strain condition. Therefore, the HEL value can be expressed as a function of yield strength σ_Y and Poisson's ratio v, and this relationship can only be applied to dynamic conditions. The plate impact geometry is illustrated in Fig. 27.

An investigation on shock compression response of the FCC-matrix CoCrFeMnNi alloy and BCC-matrix NiCoFeCrAl alloy was performed by Jiang et al. [208]. The measured free surface velocity profiles in Fig. 28 illustrate that high elastic limits (HEL) could be obtained from both HEAs (2.3–2.9 GPa and 3.5–3.9 GPa), suggesting a surprisingly high capacity to resist high speed damage. Fig. 29 displays the TEM



micrographs for undeformed and deformed specimens, suggesting low densities of dislocations and stacking faults before shock compression in both CoCrFeMnNi alloy and NiCoFeCrAl HEAs. The TEM images of deformed samples recovered from 11.7 GPa to 10.4 GPa are shown in Fig. 29(a) and Fig. 29(b), respectively. As shown in Fig. 29(a), mechanical twinning and dense dislocations are observed, therefore suggesting a dominant deformation mechanism l by dislocation movement and twinning for FCC-matrix CrMnFeCoNi alloy. In contrast, no visible twins could be found in NiCoFeCrAl alloy and only tangled dislocations can be observed, denoting a plasticity primarily led by dislocation movement. A much higher shock yield strength was measured in the latter, which could be attributed to precipitate hardening and nano-



Fig. 28 – The measured free surface particle velocity as a function of time for (a) CoCrFeMnNi and (b) NiCoFeCrAl HEAs [208].



Fig. 29 — TEM micrograph showing distinct microstructure in the deformed and undeformed samples: (a) comparison between the pre-loading manganese-containing HEA (CrMnFeCoNi) sample and sample recovered from 11.7 GPa, indicating the formation of tangled dislocations and deformation twins; (b) comparison between the pre-loading aluminumcontaining HEA (NiCoFeCrAl) sample and sample recovered from 10.4 GPa, indicating formation of tangled dislocations [208].



Fig. 30 – (a) (b) Typical dislocation structures at the wave front for bcc NiCoFeTi and TiZrNb HEAs. The atomic configuration is identified by common neighbor analysis (CNA), blue: bcc, white: defect. The red line shows the contour of moving dislocations. The inserts show the magnification of the corresponding nano-twin nucleus at the highest shock velocity. (c) Dislocation width as a function of shock velocity. NiCoFeTi and TiZrNb samples are prepared with random solid-solution while TiZrNb-Aging shows the TiZrNb samples with local chemical short-range order [209].

composite strengthening effect resulting from the formation of Cr-rich and Al-rich BCC phases [1]. The authors concluded that the exceptionally high HEL could be possibly associated with inherent server lattice distortion coupled with enhanced lattice frictional stress.

Molecular dynamics (MD) [209] using the common neighbor analysis (CAN) for defect identification pointed out the shock strength dependence of dislocation structure for BCC HEAs, illustrating a strong relationship between shock velocity and dislocation width, as shown in Fig. 30(a-c). An extended edge dislocation structure was observed in TiZrNb and NiCoFeTi HEAs which was different from BCC elemental materials Nb and Mo. Such unique dislocation structure was mainly attributed to the strong dependence of dislocation behavior on shear stress. The high shear stress was coupled with high dislocation mobility and extended morphology, while a compact dislocation core structure was associated with a low shear stress. Extended features of edge dislocations were correlated to four stages of shock response that correspond to compact dislocation core, dislocation with elastic energy radiation, extended dislocation core and twinning, as shown in Fig. 30(d).

3.3.1. Spall behavior

Spallation is a type of fracture that occurs at strain rates larger than 10^2 s^{-1} . When materials are subjected to shock waves, a tensile pulse is generated when waves reach to the free surface. During this process, if the tensile stress exceeds the strength, spall failure occurs [210,211]. The evolution of spall failure involves the nucleation and propagation of microvoids or micro-cracks resulting from the transient tensile stress. Curran [212] developed a model in 1987 to describe the formation of voids and cracks. Several properties can be measured in a spalling test including spall strength and the time to failure. This technique has received a vast amount of attention due to its importance to materials applications in military and industrial fields. In contrast to the static loading, the fracture modes and deformation mechanisms are significantly different [35,212,213]. The pressure in the shock wave can be expressed by the Rankine-Hugoniot relationship [214]:

$$P = \rho_0 U u \tag{3.18}$$

where ρ_0 is the density of specimen, u is the particle velocity, and U is the wave velocity. It can be approximated as:

$$U = c_b + su \tag{3.19}$$

where c_b is the bulk sound speed and s is linear Hugoniot slope coefficient. These two parameters are material dependent, and the equation is known as equation-of-state. Another important factor to be considered is the free-surface velocity which represents the velocity change of the free surface during impact test. This parameter is usually used to describe the initial damage and the damage evolution rate. Similar to the stress—strain curve in dynamic tensile/compressive test that provides stress profile during deformation, free-surface velocity profiles are used to investigate shock wave motion, void nucleation kinetics, and spall strength. According to Stepanov [215], the spall strength can be expressed as:

$$\sigma_f = \rho_0 c_l \Delta u \left(\frac{1}{1 + \frac{c_l}{c_b}} \right)$$
(3.20)



Fig. 31 – Optical microscope images of spall damaged (a) hot-rolling plus quenching (HRQ) sample; (b) cold-rolling plus quenching sample (CRQ). Schematic drawing of void evolution in the (c) HRQ and (d) CRQ samples [216].

where ρ_0 is the sample density, c_1 is sound velocity along longitudinal direction, Δu is equal to difference between maximum and minimum particle velocity. Therefore, it is crucial to study HEAs that can resist spallation due to their potential value to be applied to nuclear engineering, which are demanding in terms of resistance to intense ion irradiation. The spall behavior of metastable dual-phase $Fe_{50}Mn_{30}$. CoCr₁₀ HEA at two impact velocities (190 m/s and 420 m/s) has been reported by Yang et al. [216] by using a single-stage light gas gun; the formation of voids and microcracks was affected by phase composition. Alloys subjected to different manufacturing processes (hot-rolling + quenching, cold-



Fig. 32 – Evolution of micro-cracks under 420 m/s impact loading: (a) HRQ sample subjected to impact velocity of 420 m/s; (b) CRQ samples subjected to velocity of 420 m/s. Schematic drawing of the microcrack propagation: (c) evolution of microcrack in HRQ sample; (d) evolution of microcrack in CRQ sample; coalescence path at the interface of adjacent martensite phase; (e) the martensitic area with two different variant orientations; (d) the formation of coalescence path at the interface of two variants [216].



Fig. 33 — EBSD (a, c) IPF maps and (b, d) KAM maps of void coalescence and crack propagation in Al_{0.1}CoCrFeNi HEA [221].

rolling + quenching) exhibit distinctive phase components. One major important factor was the formation of martensite. The size of martensite plates for hot-rolled specimens was much larger than the cold-rolled ones, however its weight ratio was much higher and its distribution in matrix was more uniform in cold-rolled specimens. Due to a deficiency of driving force caused by neutralization between compressive residual stress and external tensile pulse, the interior of martensite was unavailable for void nucleation. However, interfaces between matrix and martensite were preferred sites for void nucleation [217-220]. Fig. 31 shows the microstructure of (a) hot-rolled and (b) cold-rolled samples subjected to 190 m/s impact loading, denoting a higher density but less intensively nucleation of voids, faster initial damage rate, and lower spall strength in cold-rolled sample. Void nucleation, growth and coalescence at an impact velocity of 420 m/s are summarized in Fig. 32, suggesting that refined and evenly distributed martensite constrains coalescence which can lead to the formation of microcracks. Phase content can also affect the propagation of microcracks. Large-sized martensite plates can serve as obstacles to deviate or block microcrack propagation while small-sized martensite plates only have very limited effect in influencing microcrack propagation, resulting in the generation of large-sized cracks.

Contrary to $Fe_{50}Mn_{30}CoCr_{10}$ HEA, whose spall strength is mainly contributed by second-phase formation, Zhang et al. [221] demonstrated a superior ability to resist spallation in $Al_{0.1}CoCrFeNi$ alloy in a single-stage gas gun experiment, despite its inferior yield strength and tensile strength. The weaker spall strength in the $Fe_{50}Mn_{30}CoCr_{10}$ HEA is the result



Fig. 34 – EBSD IPF maps showing cracking propagation and void nucleation in FeCrMnNi alloy [224].





of a soft matrix phase and rapidly growing cracks along interfaces between phases. It is noteworthy that at impact velocities below 511 m s⁻¹, deformation mode was governed by stacking faults and dislocation glide, while nano-twinning was benefited from rising shock velocity (>511 m s⁻¹). Moreover, an optimized spall strength can also be derived from approach of crack formation in $Al_{0.1}$ CoCrFeNi. Intragranular voids dominated; they tend to emerge in regions adjacent to high angle grain boundaries. These voids are then connected by high misorientation bands and eventually lead to cracking, as shown in Fig. 33.

In contrast to ductile failure found in $Al_{0.1}$ CoCrFeNi alloy, Hawkins et al. [222] observed a brittle spallation failure in FeMnCrNi alloy. They discovered that at the preliminary stage of spallation, relatively brittle ferrite phase was favored for void nucleation, as shown in Fig. 34. Although relatively same conditions (projectile velocity, tilt, etc.) were applied while spalling test, scattering results of HEL and spall strength were obtained. They conjectured that such disparate properties could possibly be caused by inhomogeneous microstructure and presence of Cr-rich ferrite phase. Given by its brittle failure mode, FeMnCrNi alloy was believed to be less desirable to be used in extreme loading conditions.

Thürmer et al. [223] provided experimental results combined with simulation for the classic equiatomic Cantor alloy. A spall strength of ~8 GPa at the strain rate of $\sim 10^7 \text{ s}^{-1}$ was found in this alloy, which was possibly caused by homogeneous distribution of composition and lack of phase segregation; the latter is detrimental to the buildup of spall strength. Moreover, traditional strengthening methods including nanotwins and precipitation also help to enhance strength.

Outside of the standard Cantor alloy composition, Xie et al. [224] reported a retardation of spallation in CoCrFeNiMn alloys charged with hydrogen. They introduced hydrogen into the alloy by using a cathodic electrochemical charging technique and investigated its effect on spallation. With the addition of hydrogen, the sizes of micro-cracks and micro-voids were



Fig. 36 – (a) Stress–strain curves and (b) Strain hardening rate vs. strain of various of high entropy alloys and ultrafinegrained-titanium (UFG-Ti).

reduced, as revealed in Fig. 35. The preliminary stage of vacancy and vacancy-hydrogen complex formation is mainly achieved by removing metallic atoms and replacing them with hydrogen atoms. Such optimized behavior primarily led to a higher diffusion barrier in hydrogen-vacancy complex. On the other hand, coalescence of micro-voids was greatly constrained by the pinning effect produced by hydrogen.

4. Summary and conclusions

Due to the complexity of multi-components systems, HEAs have highly tunable physical properties. However, systematic

understanding of HEAs subjected to dynamic loading is yet to be achieved. Here, we report on mechanical behavior of HEAs under dynamic conditions, such as spall, shock wave compression, and split Hopkinson pressure bar compression/ tension. Outstanding properties are found in FCC-based HEAs at room temperature and BCC-based HEAs at elevated temperatures. Nevertheless, the dynamic response of HEAs at cryogenic temperature is not included in present report.

Multiple deformation mechanisms contribute to the complex behavior: microband strengthening, solution strengthening, forest hardening, precipitation hardening, and martensitic transformation, which partially account for the extraordinary properties and deterred formation of shear bands. In addition, twinning is a prominent deformation mechanism in the FCC-based HEAs due to their inherently low SFE. Shear banding resulting from adiabatic deformation derived from deteriorating strain hardening rate is widely present in many HEAs. HEAs with second phases exhibit outstanding dynamic properties, perhaps better than the single-phase HEAs. The inherently hard BCC and soft FCC phases generate an exceptional work hardening ability for Al_xCoCrFeNi and WFeNiMo alloys. In general, the FCC-based alloys show high strain hardening rate, especially, the classic CoCrFeMnNi alloy. We summarize several HEAs with their corresponding dynamic stress strain curves and strain hardening rate in Fig. 36; in comparison with ultra-fine grain Ti (UFG), many HEAs exhibit much better work hardening ability in spite of the higher strength.

In comparison with FCC HEAs, the dynamic properties of refractory HEAs at room temperature are less attractive due to strongly limited ductility. However, they can retain strength at elevated temperatures due to the inherently high melting temperature of refractory elements. Therefore, their unrealized potential at high temperatures and high loading speeds makes them promising and applicable for extreme environments.

With respect of spallation failure, similar to dynamic compression/tensile tests, most of the research was performed on 3d-transition HEA, while this field of study stays vastly inadequate for BCC RHEAs. Existing studies claim that spallation in HEAs is significantly influenced by second-phase formation. Further studies need to be done to thoroughly understand how voids and cracks evolve within HEA alloys during spallation.

4.1. Applications

In contrast to the conventional materials, unique characteristics and complex constituent compositions renders HEAs superior in many aspects, such as great resistance to strength-ductility trade-off, spallation resistance, thermal stability, and highly tunable properties, which make them versatile for a broad range of applications. Some of them are discussed below.

4.1.1. Armor materials

Armor is classified, in a simplified manner, into body (personal) and heavy (armored vehicles). The requirements and the projectile-defeating mechanisms are very different. Armor materials are known to be extremely demanding in terms of their resistance to the high strain-rate loading which not merely requires strength but also involves complex thermomechanical processes, in order to minimize damage from a projectile. Superior modern armor is composed of different layers including ceramic, metallic, and polymeric materials. Armors use a range of stratagems: specially developed steels, polyamide, polyethylene, and ceramics are some of the materials. Kevlar (poly-para-phenylene terephthalamide) is a flexible body armor material with a very low density and enables the wearer to retain excellent agility. However, the materials mentioned above have to provide protection against a projectile with mass and flying speeds over 10 g and 850 m/s, respectively, if they are to be satisfactory body armors. With superior strength found

in dynamic compression/tensile tests and excellent hardening capacity found in ballistic impact tests, HEAs undoubtedly become one of most competent candidates for the metallic component of armor. The transition in deformation mode at higher strain-rate endows excellent strain hardening ability of HEAs under ultra-high loading rate. Moreover, the high strain rate sensitivity also contributes to enhanced strength at high strain rate condition.

4.1.2. Nuclear energy materials

Given by its high efficiency, economy, environmental friendliness, nuclear energy has become more and more popular, with vast amounts of effort devoted to the structural materials in nuclear reactors. Ni-based alloys, zirconium-based alloys, and stainless steels are widely used. Under atmosphere of intense irradiation with flux of neutrons and protons, severe swelling and spalling failure can occur. Irradiation damage can induce defects such as hydrogen and helium bubbles (voids). Refractory HEAs have obtained substantial amount of attention for their relatively inert characteristics due to lack of activity of the refractory elements, which make them promising candidates as next generation nuclear energy materials. The good spallation resistance of HEAs suggests the unrealized potential to endure extreme radiation damage.

4.1.3. Aerospace materials

In the aerospace industry, the structural components of turbine engines are among the most demanding environments for materials, requiring high stability, high strength, and excellent fracture strength. Contemporary aerospace materials including Fe-based, Ni-based, Ti-based and Co-based superalloys exhibit poor stability at elevated temperature. Refractory HEAs such as TaNbHfZrTi HEA and TiHfZrTaNb, have intrinsically high melting points and high configurational entropies. These endow them with outstanding stability at high temperature. One major shortcoming for the application of refractory HEAs is the undeniably high expense and their challenging processing. Therefore, besides traditional fabrication method, laser additive manufacturing as one of the latest developed processing techniques, provides defect-free components, which make HEAs more durable [225].

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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