Recent Advances in Additive Manufacturing of High Entropy Alloys and Their Nuclear and Wear-Resistant Applications

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Abstract: Alloying has been very common practice in materials engineering to fabricate metals of desirable properties for specific applications. Traditionally, a small amount of the desired material is added to the principal metal. However, a new alloying technique emerged in 2004 with the concept of adding several principal elements in or near equi-atomic concentrations. These are popularly known as high entropy alloys (HEAs) which can have a wide composition range. A vast area of this composition range is still unexplored. The HEAs research community is still trying to identify and characterize the behaviors of these alloys under different scenarios to develop high-performance materials with desired properties and make the next class of advanced materials. Over the years, understanding of the thermodynamics theories, phase stability and manufacturing methods of HEAs has improved. Moreover, HEAs have also shown retention of strength and relevant properties under extreme tribological conditions and radiation. Recent progresses in these fields are surveyed and discussed in this review with a focus on HEAs for use under extreme environments (i.e., wear and irradiation) and their fabrication using additive manufacturing.

Keywords: high entropy alloys (HEAs); additive manufacturing (AM); wear; nuclear applications; irradiation

1. Introduction

1.1. The History of High Entropy Alloys

Since the first copper-based alloy was developed around 7000 years ago, numerous metallic alloys have been utilized in various applications [1]. In traditional alloying engineering, the principal metal is used as a matrix to incorporate other alloying elements as solute. In most cases, alloying has been done to improve the strength and hardness of ductile metals. Until now, around 30 alloy systems have been introduced, based on the principal element alloying concept [2]. Increasing demands for advanced materials under harsher environments led to the innovative alloying strategies which improved the performance of existing materials against high temperatures, impact, fatigue fracture, corrosion, or wear. Heat treatments have also been used along with alloying to tailor the properties of materials for desired applications. In the 1970s, a new class of materials, named intermetallics, were developed to increase the specific hot hardness. In 1980s, another class of materials named super-alloys were developed. Inconel, Waspalloy, Hastelloy, MP35N, MP98T, Rene alloys, TMS alloys and CMSX single crystal alloys are widely used commercial superalloys. Figure 1 shows how engineering materials evolved over human history. In the beginning of the 21st century, when the alloying technology reached maturity and so did the capability of materials for more advanced applications, a new alloying concept emerged. These alloys were initially called by several different names, such as multi-principal elements alloys, equi-molar alloys, equi-atomic ratio alloys, substitutional alloys and multicomponent alloys. The most common name of these alloys is high entropy alloys (HEAs) given by J. W. Yeh [3], because these alloys have higher mixing entropy in their liquid or solid solution states than any other alloying systems. Attractive structural
properties, wide composition ranges and higher probability to find simpler microstructures enabled HEAs to gain rapidly growing attention from researchers. HEAs are considered one of three innovations in the alloying techniques along with bulk metallic glasses and metal rubbers [2].

![Figure 1. Evolution of Engineering Materials](image)

The German scientist Franz Karl Archard could be called the predecessor for HEAs research [2]. In the 18th century, Archard studied equi-mass multicomponent alloys containing five to seven different elements from Fe, Sn, Pb, Zn, Bi, Ag, Co, Sb, As and Cu. He tested these compositions for ductility, hardness, impact resistance, wear and density, etc. He published his work in a French book entitled, *Recherches sur les propriétés des alliages métalliques* [4,5]. In 1981, Brian Cantor and his student Alain Vincent started to work on Archard’s idea again at University of Sussex in Sussex, England [6]. They explored various equi-molar combinations out of 20 different elements and found that the CoCrFeMnNi alloy formed a single face-centered cubic (FCC) structure. In another independent research (MS thesis of National Tsing Hua University, Taiwan, 1996 [7]), J. W. Yeh developed different HEAs based on a concept that high entropy of an alloy system reduces the number of phases that appear in the final product. S. Ranganathan is another notable pioneering researcher in the field who wrote a review paper on the concept of HEAs and talked about the possibility of fabricating HEAs in 2003 [8]. In 2004, two independent studies by Cantor et al. [9] and Yeh et al. [3] introduced the concept of HEAs properly with the experimental results. They developed metallic alloys having nearly equi-atomic composition of more than five elements.

The basic idea of HEAs is to use five or more principal elements in or near equi-atomic compositions. According to the Gibbs phase rule, the number of phases increases with the increasing number of elements. Most of these phases are expected to be intermetallics due to their strong negative enthalpies. Binary/ternary phase diagrams also indicate
that an alloy having several principal elements would develop several phases including brittle intermetallics, resulting in complex microstructures which will limit their practical applications [8, 10]. For instance, Cu-Zn, Al-Cu and Al-Cu-Zn phase diagrams have 5, 13 and 20 intermetallics or intermediate phases, respectively [11]. By this reasoning, scientists had been reluctant to work with equi-atomic or near-equi-atomic composition made of several principal elements. This notion began to change as it was observed that the number of phases in HEAs was far less than predicted by the Gibbs phase rule [12]. The high configurational entropy of HEAs allowed for forming solid solutions rather than intermetallics, and thus, resulted in much simpler microstructures [3, 9]. The main hypothesis to explain the much smaller number of phases in HEAs is that by forming an atomic configuration of high randomness (high entropy, $\Delta S$), the Gibbs free energy of the system will be lowered ($\Delta G = \Delta H - T\Delta S$), and therefore, the formation of intermetallics is thermodynamically suppressed [3, 13, 14].

The performance of a HEA is difficult to predict as it often depends on the complicated interactions among the constituents instead of the rule of mixtures. Hence, the characterization of structure and stability under operational conditions is important before putting them into practical applications. Most of the literature mentions HEAs have been developed by “trials and errors” with an aim to get a single phase. The properties of HEAs are then compared to those of traditional alloys. For example, the wear resistance of FeCo$_{1.5}$CrNi$_{1.5}$Ti and FeAl$_{0.2}$Co$_{1.5}$CrNi$_{1.5}$Ti was reported to be higher by a factor of two compared to that of steels [15].

Since 2004, two books [2, 11] and thousands of research papers on HEAs of numerous compositions have been published. Most of the review papers focused on basic understanding and development [12, 13, 15–26], four core effects [27, 28], physical metallurgy [14], design strategies [29–31], phase stability [32, 33], or thermodynamics [34] and microstructures [35–39]. As research on the properties of HEAs progressed, various research groups reviewed the mechanical performances [40–42], heat resistance [43, 44], magnetic [45] and physical properties of HEAs [46, 47]. As synthesis techniques for HEAs advanced, various reviews on simulations and modeling [48, 49], fabrication methods [50, 51], welding techniques [52–55], high pressure technology [56] were published. Due to the possibility of having a wide composition range, new HEAs and their properties were reported continuously beyond the existing literature. In recent years, more reviews focused on diffusion studies [57], deformation behavior [58, 59], corrosion [60, 61], fracture and fatigue [62, 63], defects and radiation resistance [64–66], refractory HEAs [67], HEAs composites [68] and ceramics [69] were published. Moreover, high entropy alloys have applications in different fields such as biomedical [70, 71], energy [72], wear [73, 74], nuclear [75, 76] and creep [63], corrosion [60, 77–79].

In this context, this paper discusses as illustrated in Figure 2 recent updates on the fabrication of HEAs by additive manufacturing (AM) and the HEAs for applications under extreme environments (i.e., wear behavior and nuclear applications). Moreover, unlike previous reviews on these topics, this review would provide more convenience to readers who have just stepped in this field as well, since the reviewed research publications on AM, wear behavior and nuclear applications of HEAs are enlisted in a detailed tabular form with their results. Section 2 discusses additive manufactured HEAs in terms of their composition, microstructure and their mechanical properties, such as ultimate tensile strength (UTS), tensile elongation ($\varepsilon$), yield strength (YS), hardness (H), compressive strength (CS), compressive yield strength (CYS) and the amount of compression (C). In Section 3, the behaviors of HEAs under ion irradiation are analyzed in terms of dislocation, microstructure, irradiation resistance, hardness, phase stability, swelling resistance and self-healing. Furthermore, the tribological studies of HEAs are surveyed in terms of HEAs content variation, particle reinforcement, media and nitriding/carburizing/sulfurizing, in comparison with conventional materials. The wear behaviors of HEAs at higher temperatures and oxides formation are also reviewed.
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1.2. The Definitions of High Entropy Alloys

The first ever definition of HEA was given by Yeh et al. [3] as a class of alloys composed of five or more principal elements having concentration between 5% to 35% for each element. The second definition was also proposed by the same group [13]. In the second definition, the three categories of alloys were introduced on the basis of the configurational entropy: low entropy alloys (configurational entropy alloys (ΔS_conf) ≤ 0.69R), medium entropy alloys (0.69R ≤ ΔS_conf ≤ 1.61R) and high entropy alloys (ΔS_conf ≥ 1.61R) [30], where R is the universal gas constant. Here, the low entropy alloys are mostly conventional alloys with one or two major elements and the medium entropy alloys have two to four major elements. The high entropy alloys contain five or more major elements. The second definition does not require equi-atomic composition. For example, Ti₂ZrHf₀.₅Mo₀.₂ [80], FeCoNiCrTi₀.₂ [81] and Al₀.₃CoCrFeNi [82,83] are categorized as HEAs according to the second definition.
Moreover, these definitions are not strict, and it is not clarified which one should be used to categorize an alloy. For example, an alloy having composition of 5% A, 5% B, 20% C, 35% D and 35% E has the configuration entropy of 1.36R according to Equation (1) derived from Boltzmann’s entropy formula [30].

\[
\Delta S_{\text{conf}} = -R\sum c_i \ln c_i
\]  

where \( c_n \) is the atomic fraction of the \( n \)th element. In case of equi-atomic composition, Equation (1) reduces to [30]:

\[
\Delta S_{\text{conf}} = R \ln(n)
\]  

For example, an alloy having 25 components with equi-atomic concentration has \( \Delta S_{\text{conf}} = R \ln(n = 25) = 3.22R \). This material has the concentration of each element out of the range suggested by the first definition (between 5% to 35%), but it has sufficiently high entropy according to the second definition [15].

Considering both definitions together may often be confusing. In the past, researchers have also limited HEAs to have equi-atomic compositions or single-phase microstructures [26]. HEAs lack a standard definition that embraces all possible conditions. Both definitions are used frequently but neither clarifies the conditions of its usage. Generally, a metallic alloy with multiple principal elements and high configurational entropy is considered as a HEA.

2. Manufacturing of HEAs
2.1. Background and Conventional Methods

Brian Cantor estimated the total number of possible metallic alloys with different compositions to be up to around 10^78 [12]. This means many new alloys are yet to be discovered. For the manufacturing of HEAs, the initial synthesis strategy was to choose equi-atomic concentration of principle elements to maximize the entropy of the system. However, later, HEAs in non-equi-molar ratios were also developed for various applications. Arc melting was mostly preferred to produce HEAs thanks to its convenience, availability and simplicity. Furthermore, developing a HEA became more complex as more non-equi-atomic compositions were considered and several other manufacturing techniques were used. Alshataif et al. [84] covered almost all kinds of processing techniques used so far for HEAs synthesis. They detailed solid state processing (i.e., powder atomization methods, ball milling, cold/hot pressing, sintering, spark plasma sintering), liquid state processing (i.e., arc melting, vacuum induction melting, directional solidification, infiltration, electromagnetic stirring), thin film deposition (i.e., magnetron sputtering, pulsed laser deposition, plasma spray deposition) and additive manufacturing. Most of these manufacturing techniques are commercially available. That means most HEAs would not require a special manufacturing process and mass-producing HEAs would be possible with the existing alloying technologies and facilities.

The influence of process parameters, such as temperature and pressure, on the properties of HEAs were also studied. The effects of temperature on the properties of HEAs were studied through processes such as: annealing and heat treatments [85–104] and thermomechanical processing [105–108]. A number of research groups reported how temperature affected the microstructures and mechanical properties of HEAs in various manufacturing processes [96,109–112]. Moreover, the physical or chemical responses of various HEAs under a variety of thermal histories during manufacturing were studied: thermal aging behavior [86,113–115], TaNbHfZrTi synthesis by hydrogenation–dehydrogenation reaction and thermal plasma treatment [116], martensite formation [117–120], Al_xCoCrFeNi formation with high gravity combustion from oxides [121], laser surface melting [122], precipitation behavior [123–126] and WTaMoNbV synthesis using inductively coupled thermal plasma [127].

Researchers have also attempted to alter the microstructures and properties of HEAs by high pressure treatments. Regulating pressure during fabrication of HEAs can consider-
ably alter the interaction between the atoms by changing the interatomic distance, bonding nature and packing densities. These changes often convert the microstructures and affect the mechanical and structural properties. Dong et al. [56] reviewed the applications of high pressure technology for HEAs. They reviewed the use of dynamic high pressure, diamond anvil cells, high pressure torsion and hexahedron anvil press. Zhang et al. [128] reviewed high pressure induced phase transitions in HEAs. Application of high pressure torsion [37,129–142] is more frequent than other pressure techniques [136,143–150].

Furthermore, various researchers successfully welded/brazed HEAs [52–55,151]. Guo et al. [52] reviewed arc welding, laser welding, electron beam welding, friction stir welding to join HEAs and conducted the microstructural analysis on the welded structures. Filho et al. [54] gave a general review on the properties of welded HEAs parts and Tillmann et al. [151] reviewed HEAs brazing. Lopez et al. [53] reviewed fusion based welding (i.e., for CoCrFeNiMn and other related HEA systems) and solid state welding. Scutelniciu et al. [55] reviewed friction stir, electron and laser beam, tungsten inert gas welding techniques for CoCrFeMnNi, AlCoCrCuFeNi, AlCrFeCoNi and CoCrFeNi alloys.

2.2. Additive Manufacturing of HEAs

3-D printing in manufacturing industries, when properly applied, not only makes a design phase more efficient and economic but also brings thoughtful impacts on product design. Recent advances in additive manufacturing (AM) made it more influential throughout the supply chain which generates revenue as well [152]. The additive manufactured HEAs showed improvement in their mechanical properties in comparison to as-cast HEAs [153–160]. Higher cooling rates in AM processes help suppress diffusional phase transformation and increase the chemical homogeneity of HEAs [161]. Under certain circumstances, AM gives a better control over the material processing and helps tailor application-specific microstructures which become more important for the parts for applications under extreme environments. For example, it was demonstrated that fine and tailorable microstructures in HEAs were obtained using AM techniques [162–169], which implies AM can improve the mechanical performance of at least some HEAs. However, this may not be a trivial task as a good understanding of the AM technique and material behavior during the AM process is required [170].

AM of HEAs has been discussed briefly in a few review papers [51,161,171,172] and books [2,173]. Xiaopeng Li [161] discussed the requirements and challenges of AM of HEAs and bulk metallic glasses. Chen et al. [51] examined the microstructural evolution and mechanical properties of AM-processed CoCrFeNi, AlxCoCrFeNi, CoCrFeMnNi and Ti25Zr50Nb50Ta25. Fabricating HEAs by spark plasma sintering (SPS) and their property analyses were discussed in the book chapter “Spark Plasma Sintering of High Entropy Alloys” of [174]. SPS followed by mechanical alloying has largely been used to develop HEAs, which was reviewed in detail by Vaidya et al. [175]. Therefore, SPS studies are not included here.

In this review, studies on the AM of HEAs are tabulated and the mechanical properties of these HEAs are discussed. Tables 1–3 detail the HEAs synthesized by selective laser melting (SLM), electron beam melting (EBM) and direct energy deposition (DED), respectively. The performances of these HEAs are discussed in terms of their composition, their microstructure and their mechanical properties, such as ultimate tensile strength (UTS), % elongation at fracture (ε), yield strength (YS), hardness (H), compressive strength (CS), compressive yield strength (CYS), bending strength (BS), bending elongation (δb) and % compression at fracture (C).
Table 1. The compositions, microstructures and mechanical properties of SLM manufactured HEAs.

| Source                          | Alloy Composition      | Microstructure (Grain Size)                  | Result UTS (MPa), YS (MPa), BS (MPa), $\delta$ b (mm), $\epsilon$ (%), H, CS (MPa), C (%) |
|---------------------------------|------------------------|----------------------------------------------|----------------------------------------------------------------------------------------|
| Chen et al. [176]               | CoCrFeMnNi            | FCC (53.1 $\mu$m)                            | UTS = 281 $\pm$ 18, YS = 12.5 $\pm$ 0.5, H = 261 $\pm$ 7 HV                           |
| Niu et al. [169]                | CoCrFeMnNi            | FCC (<5 $\mu$m)                              | CS = 2447.7                                                                           |
| Li et al. [177]                 | CoCrFeMnNi + TiNp nanoparticles | FCC                                | UTS = 601–1036, $\epsilon$ = 12–30                                                 |
| Li et al. [178]                 | CoCrFeMnNi + Fe based metallic glass | FCC                                | UTS = 916–1517                                                                     |
| Li et al. [179]                 | CoCrFeMnNi + TiN nanoparticles | -                                     | -                                                                                     |
| Kim et al. [180]                | (CoCrFeMnNi)C        | FCC (180–330 nm)                             | YS = 800–900, $\epsilon$ = 25–30                                                   |
| Li et al. [181]                 | CoCrFeMnNi + 12 wt% nano-TiNp | FCC (<2$\mu$m)                      | UTS = 1100                                                                           |
| Li et al. [182]                 | CoCrFeMnNi + TiN nanoparticles | FCC                                  | -                                                                                     |
| Zhu et al. [153]                | CoCrFeMnNi            | FCC                                          | H = 212 HV                                                                           |
| Xu et al. [183]                 | CoCrFeMnNi            | FCC (1–2 $\mu$m)                            | $H = 2.84 \pm 0.13$ GPa                                                              |
| Park et al. [154]               | CoCrFeMnNi + 1 at%C | FCC (20–35 $\mu$m)                         | UTS = 829–989, YS = 741, $\epsilon$ = 24.3                                         |
| Ren et al. [184]                | CoCrFeMnNi            | -                                            | -                                                                                     |
| Dovgyy et al. [185]             | CoCrFeMnNi            | FCC & cubic (0.2–0.8 $\mu$m)                | -                                                                                     |
| Zhou et al. [155]               | CoCrFeMnNi + 0.5 at%C | FCC (40–50 $\mu$m)                        | UTS = 776–797, YS = 630–656, $\epsilon$ = 7.7–13.5                                 |
| Wu et al. [186]                 | CoCrFeMnNi + 0.5 at%C | FCC (40–50 $\mu$m)                        | UTS = 795, YS = 638                                                                 |
| Lin et al. [198]                | CoCrFeNi              | FCC                                          | -                                                                                     |
| 0.5 at%C                       | CoCrFeNi              | FCC                                          | -                                                                                     |
| Sun et al. [187]                | CoCrFeNi              | $\approx$ 3 mm in length and $\approx$ 200 $\mu$m in width | UTS = 676.7–691, YS = 556.7–572, $\epsilon$ = 12.4–17.9|
| Song et al. [188]               | CoCrFeNi + N (1.8%)   | FCC                                          | UTS = 600–853, YS = 520–650, $\epsilon$ = 27                                         |
| Zhou et al. [189]               | (CoCrFeNi)$_{1-x}$($WC_x$) | FCC (1 $\mu$m)                       | H = 603–768 HV                                                                     |
| Brif et al. [156]               | CoCrFeNi              | FCC                                          | -                                                                                     |
| Niu et al. [190]                | AlCoCrFeNi            | Disordered (A2) + Ordered (B2) BCC           | H = 632.8 HV                                                                         |
| Karlsson et al. [170]            | AlCoCrFeNi            | FCC & BCC (<20 $\mu$m)                     | -                                                                                     |
| Peyrouzet et al. [157]           | Al$_{0.3}$CoCrFeNi    | FCC (width=13 and length=70–120 $\mu$m)     | UTS = 896, YS = 730, $\epsilon$ = 29                                               |
| Sun et al. [158]                | Al$_{0.3}$CoCrFeNi    | FCC & BCC (1 $\mu$m)                        | UTS = 878, YS = 609, H = 270HV                                                      |
| Zhou et al. [160]                | Al$_{0.3}$CoCrFeNi    | FCC                                          | UTS = 721, YS = 579, $\epsilon$ = 22                                               |
| Luo et al. [191]                 | AlCrCuFeNi            | BCC (avg. width=4 $\mu$m)                  | CS = 1655.2–2052.8, C = 6.5–6.8                                                     |
| Luo et al. [192]                 | AlCrCuFeNi$_x$ (2 $\leq x \leq$ 3) | FCC (thickness=490 nm) & BCC                  | UTS = 957, $\epsilon$ = 14.3                                                        |
| Li et al. [112]                  | AlCoCuFeNi            | BCC                                          | YS = 744, $\epsilon$ = 13.1, CS = 1600                                              |
| Yao et al. [193]                 | AlCrFeNiV             | FCC (width=15 $\mu$m, length=75–200 $\mu$m) | UTS = 1057.47, $\epsilon$ = 30.3                                                   |
| Wang et al. [194]               | AlCoCrCuFeNi          | FCC & BCC                                    | H = 710.4 HV                                                                         |
| Wang et al. [195]                | AlMgScZrMn            | FCC                                          | $H = 42.8–86.7$ HV                                                                  |
| Sarawat et al. [196]             | AlCoFeNiV$_{0.9}$Sm$_{0.1}$ | FCC                                | UTS = 394, $\epsilon$ = 10.5                                                       |
| Agrawal et al. [197]             | Fe$_{49}$Mn$_{23}$Cr$_{9}$Si$_{5}$ | HCP                                      | UTS = 1100, YS = 530, $\epsilon$ = 30                                              |
| Zhang et al. [198,199]           | NbMoTaW                | BCC (13.4 $\mu$m)                          | H = 826 HV                                                                           |
Table 1. Cont.

| Source            | Alloy Composition | Microstructure (Grain Size) | Result UTS (MPa), YS (MPa), BS (MPa), $\delta_b$ (mm), $\varepsilon$ (%), H, CS (MPa), C (%) |
|-------------------|-------------------|-----------------------------|--------------------------------------------------------------------------------------------------|
| Yang et al. [200,201] | Ni$_6$Cr$_4$WFe$_9$Ti | FCC (300–1000 nm) + unknown phase | UTS = 972, YS = 742, $\varepsilon$ = 12.2 |
| Chen et al. [202]   | CoCrFeNiMn        | FCC + Mn$_2$O$_3$ particles | YS = 620, UTS = 730, $\varepsilon$ = 12 |
| Litwa et al. [203]  | CoCrFeNiMn        | FCC                         | H = 320 HV |
| Zhang et al. [204]  | CoCrFeNiMn        | FCC                         | YS = 729.6 |
| Kim et al. [205]    | CoCrFeNiMn        | FCC                         | YS = 752.6 |
| Choi et al. [206]   | CoCrFeNiMn        | FCC                         | - |
| Su et al. [207]     | CrCuFeNi$_2$, Al$_{0.5}$CrCuFeNi$_2$, Al$_{0.5}$CrCuFeNi$_2$, AlCrCuFeNi$_2$ | FCC, FCC + BCC/B2, FCC + BCC/B2 | - |
| Peng et al. [208]   | CoCrFeNi + Ti coated diamond | FCC + diamond particles, FCC + Cr$_2$C$_3$ + diamond particles | H = 622 HV, BS = 530, $\delta_b$ = 0.64, H = 615 HV, BS = 925, $\delta_b$ = 0.48 |
| Wang et al. [209]   | CoCrFeNiMn        | FCC                         | H = 164–370 HV |
| Sun et al. [210]    | Al$_{0.5}$CrCuFeNi, Al$_{0.5}$CrCuFeNi, AlCrCuFeNi | FCC, FCC | - |
| Ishimoto et al. [211] | Ti$_{1.4}$Nb$_{0.6}$Ta$_{0.4}$Zr$_{1.4}$Mo$_{0.6}$ | BCC | YS = 1690 |
| Park et al. [212]   | (CoCrFeMnNi)$_{0.5}$C$_1$ | FCC | YS = 741, UTS = 874 |
| Lin et al. [213]    | CoCrFeNi         | FCC                         | YS = 701 ± 14, UTS = 907 ± 25 |
| Kim et al. [214]    | CoCrFeNiMn       | FCC                         | - |
| Jin et al. [215]    | CoCrFeNiMn       | FCC                         | YS = 520 ± 10, UTS = 770 ± 10, $\varepsilon$ = 25 |
| Lin et al. [216]    | Al$_{0.5}$Co$_{0.5}$CrFeNi$_{1.5}$Ti$_{0.5}$ | FCC + $\sigma$ + L12 | YS = 1235, UTS = 1550 |
| Peng et al. [217]   | CoCrFeNiMn       | FCC                         | - |
| Vogiatzief et al. [218] | AlCrFe$_2$Ni$_2$, Heat treatment (750–950 °C, 3 h & 6 h) | FCC + BCC | H = 276–483 HV |
| Liao et al. [219]   | Al$_{0.5}$FeCrNi$_{1.5}$V$_{0.2}$ | FCC | H = 220–240 HV |
| Guo et al. [220]    | CoCrFeNiMn       | FCC                         | YS = 622, UTS = 763, $\varepsilon$ = 16 |
| Kim et al. [221]    | (CoCrFeNiMn)$_{100-x}$C$_x$ | FCC (15–22 µm) | YS = 653–753, UTS = 766–911 |
| Zhao et al. [222]   | CoCrFeNi         | FCC                         | H = 238–525 HV |
| Gu et al. [223]     | CoCr$_{2.5}$FeNi$_2$TiW$_{0.5}$ | FCC | YS = 449–581, CS = 823–893, $\varepsilon$ = 4.4–9.9, H = 436.7–499.2 HV |

Table 2. The compositions, microstructures and mechanical properties of EBM manufactured HEAs.

| Source            | Alloy Composition | Microstructure (Grain Size) | Result UTS (MPa), YS (MPa), $\varepsilon$ (%), H, CS (MPa), C (%) |
|-------------------|-------------------|-----------------------------|--------------------------------------------------------------------------------------------------|
| Peng et al. [224] | CoCrFeNiMn        | FCC                         | YS = 196 |
| Wang et al. [225] | CoCrFeMnNi        | FCC                         | YS = 497, 205, H = 157.1 HV |
| Kuwabara et al. [226] | AlCoCrFeNi     | BCC & FCC                   | UTS = 1073, YS = 769, $\varepsilon$ = 0–1.2, YS = 944–1015, CS = 1447–1668, C = 14.5–26.4 |
| Wang et al. [227] | AlCoCrFeNi        | BCC                         | - |
| Fujieda et al. [228] | CoCrFeNiTi      | FCC + Cubic                 | UTS = 1178, YS = 773, $\varepsilon$ = 25.8 |
| Popov et al. [229] | Al$_{0.5}$CrMoNbTa$_{0.5}$ | FCC                         | - |
Table 3. The compositions, microstructures and mechanical properties of DED manufactured HEAs.

| Scheme | Alloy Composition | Microstructure (Grain Size) | Result |
|--------|-------------------|----------------------------|--------|
| Guan et al. [230] | CoCrFeMnNi | FCC (13 µm) | UTS (MPa), YS (MPa), ε (%) = 517, ε = 26 |
| Melia et al. [231] | CoCrFeMnNi | FCC (~4 µm) | UTS = 647–651, ε = 232–424 |
| Li et al. [232] | CoCrFeMnNi | FCC | |
| Gao et al. [233] | CoCrFeMnNi | FCC (30–150 µm) + BCC | UTS = 620, ε = 448 |
| Xiang et al. [234,235] | CoCrFeNiMn | FCC | UTS = 400–600 |
| Chew et al. [236] | CoCrFeNiMn | FCC (3.68 ± 0.85 µm) | UTS = 660, ε = 518 |
| Qiu et al. [237] | CoCrFeMnNi | FCC | UTS = 891, ε = 564 |
| Li et al. [238] | CoCrFeMnNi + WC (0–10 wt%) | FCC | UTS = 550–845, ε = 300–675, ε = 9 |
| Amar et al. [239] | CoCrFeMnNi + TiC (0–5 wt%) | FCC | UTS = 550–723, ε = 300–385, ε = 32 |
| Guan et al. [240] | CoCrFeMnNi AlCoCrFeNiTi0.5 | FCC (24 µm) | YS = 888–1100, H = 197–657 HV |
| Wang et al. [241] | CoCrFeMnNiMo0.2 | FCC | UTS = 532–928, ε = 37 |
| Zhou et al. [242] | CoCrFeNiNbx | FCC | UTS = 400–820, ε = 220–750 |
| Gwalani et al. [243] | AlxCoCrFeNi (x = 0.3–0.7) | FCC | |
| Zhou et al. [244] | CoCrFeNiNb | FCC | |
| Gwalani et al. [245] | AlxCoCrFeNi (x = 0.3–0.7) | FCC + BCC | H = 170–380 HV |
| Nartu et al. [246] | AlxCoCrFeNi | FCC | UTS = 400–820, ε = 220–750 |
| Mohanty et al. [247] | AlCrFeMoV | FCC | |
| Gwalani et al. [248] | AlxCoCrFeNi | FCC + BCC | H = 350 HV, |
| Malatji et al. [249] | AlxCoCrFeNiCu | FCC + BCC | |
| Moorehead et al. [250,251] | NbMoTaW | BCC | |
| Kunc et al. [252] | TiZnNbMoV | BCC | |
| Dobbelein et al. [253] | TiZnNbHfTa | BCC | H = 509 HV0.2 |
| Peggues et al. [254] | CoCrFeNiMn | FCC | |
| Li et al. [255] | CoCrFeNiMn | FCC | |
| Tong et al. [256] | CoCrFeNiMn | Vacuum arc melting | |
| Shen et al. [257] | CoCrFeNi (SiC) | FCC + Cr3C2 (1 µm) | UTS = 2155–2499, ε = 125–713, H = 139–310 |
| Cai et al. [258] | CoCrFeNi | FCC | |
| Zhang et al. [259] | NbMoTaNbx | FCC + B2 | |
| Peng et al. [260] | AlxCoCrFeNi | FCC | |
| Kuzminova et al. [261] | CoCrFeNi | FCC | UTS = 456–551, ε = 208–259 HV |
| Malatji et al. [262] | NiCrFeNi | FCC | |
| Dong et al. [263] | CoCrFeNi | FCC | |
| Zhou et al. [264] | CoCrFeNi | Solution treatment (2–1250 °C) | |
| Zheng et al. [265] | CoCrFeNi | FCC + HCP (Laves C14) + Nb rich carbide | |

Cantor alloy (i.e., CoCrFeMnNi) and its variants have been largely investigated. Apart from SPS, SLM is the most widely studied AM technique for HEAs [196–231]. HEAs that were successfully fabricated by SLM include CoCrFeNiMn [153,154,169,176–185,188,202–206,209,212,214,215,217,220,221], AlCrFeNiV [193], AlCoCrFeNi [170,190], AlCoCrCuFeNi [194], CoCrFeNi [98,155,186,187,208,213,222], CoCr2.5FeNi2TiW0.5 [223],
Various HEAs have exhibited significant improvement in their mechanical properties after AM synthesis as compared to the as-cast structures of the same compositions [153,154]. Zhou et al. [155] reported that arc-melted CoCrFeNi had the YS of 225 MPa whereas SLM-manufactured CoCrFeNi had the YS of 656 MPa. Brié et al. [156] observed that SLM-manufactured CoCrFeNi showed noticeable improvement in YS from 188 MPa (as-cast) to ~142 MPa to ~713 MPa, respectively.

Improvement in mechanical properties was reported when HEAs were manufactured CoCrFeNiMn showed noticeable improvement in YS from 188 MPa (as-cast) to ~139 HV to ~310 HV and ~2270 MPa, respectively. The UTS for x = 0.5, 1, 1.5 was measured to be 653, 752 and 753 MPa respectively. The UTS for x = 0.5, 1, 1.5 was found to be 766 ± 318.5, 895 ± 22.3 and 911 ± 125.1 MPa, respectively.

Shen et al. [258] discussed the effect of SiC particles added to CoCrFeNi. They noticed that adding SiC particles changed the microstructure from the FCC phase to the FCC/Cr23C6 dual phase. The hardness and YS improved significantly from ~139 HV to ~310 HV and ~125.1 MPa, respectively.

Meanwhile, EBM was used to manufacture CoCrFeNiMn [225], AlCoCrFeNi [226,227], CoCrFeNiTi [228] and Al0.5CrMoNbTa0.5 [229]. DED techniques were used to fabricate CoCrFeNiMn [230–240,255–258], CoCrFeNi [258,259,262], Al0.5CoCrFeNi [261], CoCrFeNiMo0.2 [241], CoCrFeNiNb0.3 [242], Al1CoCrFeNi [243–245,259,264], AlCoCrFeNi21.2 [246,264], Al2CoCrFeMoV0.4 [247], Al2CoCrFeNiTi0.5 [248], AlCrCuFeNi [249,263], Al2CoCrFeNiCu/AlTiCrFeCoNi [250,251], NbMoTaW [252], TiZrNbMoV [253], NbMoTaTi3Ni6 [260], CoCrFeNb0.2Ni2.1 [265] and TiZrNbHfTa [254].

The microstructures and mechanical behaviors of the HEAs produced by different AM processes are still under investigation by several research groups. The HEAs listed in Tables 1–3 mainly have either FCC or BCC microstructures except CoCrFeNiMn which has HCP. Improvement in mechanical properties was reported when HEAs were fabricated with AM [158,169,177,180,188,193,268]. These improvements are mostly attributed to grain refinement. Grain refinement in HEAs is claimed to be due to the high cooling rates as it happens in various other materials [158,200,230]. Moreover, the wear behavior [189,249], thermo-mechanical analysis [199,246], effect of annealing [98], creep behavior [183], residual stresses [232], corrosion behavior [176,198,226,228,231,241,249], strengthening mechanisms [153] and deformation mechanism [237] of additive manufactured HEAs have also been reported.

Particle reinforcement in a HEA matrix with AM has been an area of interest for many researchers lately who expect microstructure refinement and mechanical properties enhancement [123,177,208,212,220,221,269–280]. Li et al. [177] introduced nano TiN ceramic particles in a CoCrFeMnNi matrix, which led to equiaxed grains of 5 µm. The same group [179] also fabricated the same composition with SLM followed by laser remelting and obtained ultrafine grains (80% grains less than 2 µm and 90% grains less than 3.5 µm). Song et al. [188] showed that the YS and ductility of CoCrFeNi increased by 25% and 34%, respectively, when doped with 1.8 at% nitrogen. Fu et al. [279] noticed that adding TiC particles increased the YS and UTS up to 1100 MPa. Rogal et al. [271] increased the UTS of CoCrFeNiMn up to 1600 MPa by introducing nano-Al2O3 particles. Carbon doping was attempted [154,155,180,186] to enhance the mechanical properties of HEAs. Peng et al. [208] added diamond particles into CoCrFeNi and found out the bending strength was 925 MPa. Park et al. [212] added carbon into CoCrFeNiMn ((CoCrFeNiMn)0.9C1) and noticed that the YS and UTS were ~741 MPa and ~874 MPa, respectively. Similarly, Kim et al. [221] also added carbon into CoCrFeNiMn in a ratio (CoCrFeNiMn)100-xC x = 0.5–1.5. The YS for x = 0.5, 1, 1.5 was measured to be 653, 752 and 753 MPa respectively. The UTS for x = 0.5, 1, 1.5 was found to be 766 ± 318.5, 895 ± 22.3 and 911 ± 125.1 MPa, respectively.

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to 600 MPa and in UTS from 457 MPa (as-cast) to 745 MPa. Peyrouzet et al. [157] showed that the YS of Al0.3CoCrFeNi increased from 275 MPa (as-cast) to 730 MPa and the UTS from 502 MPa (as-cast) to 896 MPa when manufactured with SLM. The UTS of as-cast Al0.3CoCrFeNi was 522 MPa and it was increased to 878 MPa with SLM processing [158]. Arc-melted Al0.3CoCrFeNi had the YS of 334 MPa and the UTS of 709 MPa [159]. SLM increased the YS up to 579 MPa and the UTS up to 721 MPa [160].

Moreover, the CS of AlCrCuFeNi was 2052 MPa when fabricated with SLM and 1750 ± 15 MPa [281] with arc-melting. The hardness of AlCoCrCuFeNi improved from 500 to 710 Hv [194] by using SLM. The YS of AlMgScZrMn manufactured with arc melting, SPS, and SLM is 188 ± 2.3 MPa, 231 ± 3 Mpa and 394 Mpa respectively [195]. Agrawal et al. [197] reported that the YS of as-cast and SLM-printed Fe40Mn20Co20Cr15Si5 was 420 ± 20 Mpa and 530 ± 40 Mpa, respectively. The YS of CoCrFeNiMn was 2.5 times higher (around 518 Mpa) [26] with DED in comparison to that of cast parts (209 Mpa) [282] at room temperature (RT). Furthermore, the as-cast AlCoCrFeNi had the UTS of 956 Mpa, and the EBM specimen had the UTS of 1073 MPa [226]. Similarly, Fujieda et al. [228] reported that EBM-synthesized CoCrFeNiTi showed the improved tensile strength of around 1178 MPa, which is much stronger than various commercial high corrosion resistant materials such as duplex stainless steel: 655 MPa, super duplex stainless steel: 750–800 MPa and Ni-based super alloys (i.e., Alloy C276: 690 MPa, Alloy 718: 1275 MPa).

Refractory HEA NbMoTaW has shown a drastic reduction in grain size when made with AM. The average grain size of BCC phase was 200 µm in as-cast sample [283] and 13.4 µm in SLM-processed sample. Additionally, this alloy did not follow the rule of mixtures. Instead, it showed the cocktail effect for the hardness of the final structure. The hardness of Nb, Mo, Ta and W was in the range of 85–410 HV but the final hardness of SLM processed NbMoTaW was measured to be 826 HV [198]. Senkov et al. [284] commented that NbMoTaW did not have any abrupt hardness changes at high temperatures, consistently exhibiting better hardness properties than superalloys. Moreover, SLM-processed Ni6Cr4WFe9Ti (UTS = 972 MPa, YS = 742 MPa, ε = 12.2%) had ~93% increase in YS, ~50% increase in UTS, and ~77% increase in tensile ductility as compared to the vacuum arc melted samples (UTS = 649 MPa, YS = 385 MPa, ε = 6.9%) [200,201].

In summary, various studies have successfully manufactured SLM, EBM and DED techniques. They have also shown that the properties of HEAs could be altered by changing the input parameters for AM process. For example, CoCrFeNiMn was manufactured with SLM by multiple researchers [153,154,169,176–185,188,202–206,209,212,214,215,217,220,221] and many of them acquired different mechanical properties for CoCrFeNiMn by changing input parameters in AM processes (refer Tables 1–3).

3. Applications under Extreme Environments

3.1. Nuclear Applications

Nuclear energy is contributing to around 13% of electricity demand worldwide [285] with negligible carbon emission. The safety, reliability and economy of these nuclear power plants depends heavily on the performances of advanced structural materials under high-energy irradiation and elevated temperatures [286,287]. Radioactive waste handling units also require radiation-tolerant materials. Not to mention nuclear applications, radiation-resistant materials are in great demand in medical and aerospace fields as well.

The typical range of operating temperatures of nuclear reactors spans from 350 to 900 °C as listed in Table 4 [288]. At high temperatures, several effects come into play such as thermal expansion, vacancy concentration, diffusion rate, phase transformation, precipitation, recovery, recrystallization, dislocation climb, creep, grain weakening/migration/growth, oxidation and intergranular oxygen dispersion. With conventional alloys, design strategies for nuclear reactor materials were mostly concerned with tuning the microstructures by various heat treatments, precipitation, cold working and solute atoms to get desired properties. HEAs, though, introduce the concept of modifying compositional complexity of the structural materials to make them suitable for nuclear applications.
Currently, reduced activation ferritic/martensitic steels (RAFM) (e.g., F82H, EUROFER 97), are the most popular option for irradiation-resistant structural materials. Oxide dispersion strengthened (ODS) RAFM steels (i.e., EUROFER 97 reinforced with 0.3 wt.% Y$_2$O$_3$ particles), C/C, SiC/C, SiC/SiC, refractory metals/alloys (W, Cr), V and Ti-based alloys are also being used [289,290]. HEAs are considered to be potential candidates for nuclear applications [2,291–293]. Yeh et al. [294] mentioned that HEAs are potential candidates for structural materials of the 4th generation nuclear reactor. Previously, the irradiation responses and defect behaviors [65,66], intrinsic transport properties [66], irradiation induced structural changes [295] of HEAs were reviewed. Building upon these reviews, this section mainly focuses on ion irradiation resistance of HEAs.

The majority of the previous ion irradiation studies on HEAs are listed in Table 5 where phases, irradiation conditions and important findings are summarized. These HEAs were studied under Ni, Au, Ag, Ar, He, Kr, or Xe ions irradiation. The most popular strategy to design single-phase HEAs of high irradiation resistance used elements having low activation or thermal neutron absorption cross section [296–300].

### Table 5. Summary of irradiation studies on HEAs.

| Source | Material (Fabrication) | Phase | Irradiation Conditions (Energy, Ion, Fluence, Temperature) |
|--------|------------------------|-------|----------------------------------------------------------|
| Jawaharram et al. [301] | CoCrFeNiMn | FCC | 2.6 MeV, Ag $^{+}+$, $1.5 \times 10^{-3}$ & $1.9 \times 10^{-3}$ dpa$^{-1}$ s$^{-1}$, 23–500 °C |
| Lu et al. [302] | NiCoFeCr, CoCrFeNiMn | FCC | 3 MeV, Ni$^{2+}$, $5 \times 10^{16}$ ions·cm$^{-2}$, 500 °C |
| Barr et al. [303] | CoCrFeNiMn | FCC | 3 MeV, Ni$^{2+}$, $3 \times 10^{15}$ ions·cm$^{-2}$, 500 °C |
| Lu et al. [304] | CoCrFeNi, CoCrFeNiMn | FCC | 1.5 MeV, Ni$^{+}$, $4 \times 10^{14}$ & $3 \times 10^{15}$ ions·cm$^{-2}$ (peak dose~4 dpa), 500 °C 3 MeV, Ni$^{+}$, $5 \times 10^{16}$ ions·cm$^{-2}$ (peak dose~60 dpa), 500 °C |
| Tong et al. [305] | CoCrFeNiMn, CoCrFeNi | FCC | 16 MeV, Ni$^{5+}$, 8 MeV Ni$^{3+}$, 4 MeV Ni$^{1+}$ & 2 MeV Ni$^{1+}$, 0.1–1 dpa, 420 °C |
| Jin et al. [306] | CoCrFeNi, CoCrFeNiMn | FCC | 3 MeV, Ni$^{2+}$, $5 \times 10^{16}$ ions·cm$^{-2}$ (peak dose~53 dpa), 500 °C |
| Chen et al. [307] | CoCrFeMnNiAl$_{0.3}$CoCrFeNi | FCC | 1 MeV, K$_{r}$ ions, $6.3 \times 10^{15}$ ions·cm$^{-2}$, 300 °C |
| Wang et al. [308] | CoCrFeNiCu | FCC | 100 keV, He$^{+}$, $2.5 \times 10^{17}$, $5 \times 10^{17}$ & $1 \times 10^{18}$ ions·cm$^{-2}$, RT |
| He et al. [309] | CoCrFeNi, CoCrFeNiMn, CoCrFeNiPd | FCC | electrons, $5 \times 10^{18}$ e·cm$^{-2}$·s$^{-1}$, 400 °C |
| Yang et al. [310] | CoCrFeNiMn, CoCrFeNiPd | FCC | 3 MeV, Ni$^{2+}$, $5 \times 10^{16}$ ions·cm$^{-2}$, 420, 500 & 580 °C |
| Yang et al. [311] | CoCrFeNiMn | FCC | - He ion, - RT & 450 °C |
| Hashimoto et al. [312] | CoCrFeNiMn, CoCrFeNiAl$_{0.3}$ | FCC | 1250 keV, 1.5 dpa, 300–400 °C |
| Zhang et al. [313] | CoCrFeNiCu | FCC | 3 MeV Ni$^{2+}$, $10^{14}$ ions·cm$^{-2}$, RT |
| Yang et al. [314] | CoNi, FeNi, CoCrFeNi | FCC | 3 MeV, Ni$^{2+}$, $1.5 \times 10^{16}$ ions·cm$^{-2}$, 420, 500 & 580 °C |
| | | | $5 \times 10^{16}$ (peak dose~17 dpa) & $5 \times 10^{16}$ (peak dose~53 dpa) ions·cm$^{-2}$, 500 °C |
| Source                  | Material (Fabrication)          | Phase | Irradiation Conditions (Energy, Ion, Fluence, Temperature) |
|-------------------------|---------------------------------|-------|----------------------------------------------------------|
| Abbaya et al. [315]     | CrCoFeNi                         | FCC   | 1.5 MeV, Ni$_{2+}$, $1 \times 10^{15}$ (peak dose=2 dpa) & 5 $\times 10^{15}$ (peak dose=96 dpa) ions·cm$^{-2}$, RT |
| Sellami et al. [316]    | CoCrFeNi                         |       | 1.5 MeV, Ni$_{2+}$, $1 \times 10^{15}$–$1 \times 10^{14}$ ions·cm$^{-2}$, 21 MeV, Ni$_{2+}$, $2 \times 10^{15}$ & $1 \times 10^{16}$ ions·cm$^{-2}$, RT |
| Chen et al. [317]       | CoCrFeNi                         | FCC   | 275 keV, He$^+$, 5.14 $\times 10^{15}$ ions·m$^{-2}$, 250, 300, 400 °C |
| Kombaiah et al. [318]   | CoCrFeNi, Al$_{12}$CoCrFeNi     | FCC   | 3 MeV, Ni$_{2+}$, $1 \times 10^{17}$ ions·cm$^{-2}$ (peak dose~100 dpa), 500 °C |
| Lu et al. [319]         | CoCrFeNiS                        | FCC   | 3 MeV, Ni$_{2+}$, 5 $\times 10^{16}$ ions·cm$^{-2}$, 580 °C |
| Tunes et al. [320]      | CrFeNiMn                         | FCC   | 30 keV, Xe$^+$, 2.6 $\times 10^{16}$ ions·cm$^{-2}$, 500 °C |
| Edmonson et al. [321]   | CoFeNi                           | BCC   | 30 keV, Xe$^+$, 9.3 $\times 10^{18}$ ions·cm$^{-2}$, 6 keV He$^+$, 6.4 $\times 10^{16}$ ions·cm$^{-2}$, RT |
| Fan et al. [322]        | CoCrFeNi                         | FCC   | 3 MeV, Ni ions, 5 $\times 10^{16}$–8 $\times 10^{16}$ ions/cm$^{-2}$, 580 °C |
| Chen et al. [81]        | CoCrFeNiTi$_{0.2}$              | FCC   | 275 keV, He$^{2+}$, 5.14 $\times 10^{20}$ ions·m$^{-2}$, 400 °C |
| Lyu et al. [323]        | CoCrFeNiMn$_{0.2}$              |       | 27 keV, electrons, -, RT |
| Xu et al. [324]         | (CoCrFeNi)$_{3.0}$Ti$_{0.1}$Nb$_{0.1}$ | FCC   | 2.5 MeV, Fe ions, 1.5 $\times 10^{19}$ ions·m$^{-2}$, RT-500 °C |
| Cao et al. [325]        | (CoCrFeNi)$_{0.8}$Ti$_{0.2}$Al$_{1.4}$ | FCC   | 4 MeV, Au ions, 10–49 dpa, RT |
| Tolstolutskaya et al. [326] | Cr$_{13}$Fe$_{0.7}$Mo$_{0.9}$Ni$_{0.2}$ | FCC   | 1.4 MeV, Ar ions, 0, 0.3, 1 & 5 dpa, RT |
| Kumar et al. [327]      | Fe$_{0.27}$Ni$_{0.95}$Mo$_{0.5}$Cr$_{0.18}$ | FCC   | 3 MeV, Ni$_{2+}$, 4.2 $\times 10^{14}$, 4.2 $\times 10^{14}$ & 4.2 $\times 10^{15}$ ions·cm$^{-2}$, RT & 500 °C |
| Li et al. [328]         | Cr$_{0.16}$Fe$_{0.63}$Mo$_{0.2}$ | FCC   | Neutron, 8.9 $\times 10^{14}$ n·cm$^{-2}$, s, 60 °C |
| Voyevodin et al. [329]  | Cr$_{13}$Fe$_{0.4}$Mo$_{0.1}$Ni$_{0.1}$ | FCC   | 1.4 MeV, Ar ions, 2.2 $\times 10^{15}$ ions·cm$^{-2}$, RT |
| Dias et al. [330]       | Cu$_{0.5}$CrFeTi (x = 0.21–1.7) | BCC + FCC | 300 keV, Ar$^+$, 3 $\times 10^{19}$ at·m$^{-2}$, RT |
| Yang et al. [328]       | Al$_{0.5}$CoCrFeNi              | FCC   | 3 MeV, Au ions, 6 $\times 10^{15}$ ion·cm$^{-2}$ (peak dose ~31 dpa), 250–650 °C |
| Gromov et al. [331]     | AlCoCrFeNi                       | -     | 18 keV, electrons, -, RT |
| Zhang et al. [299]      | Al$_{0.5}$CrMoNi$_{0.5}$Zr$_{0.5}$ (AtCrMoNi$_{0.5}$Zr$_{0.5}$) | FCC   | 400 keV, He$^+$, 8 $\times 10^{15}$ & 8 $\times 10^{16}$ ion·cm$^{-2}$, RT |
| Yang et al. [82]        | Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi | FCC, FCC + B2, A2 + B2 | 3 MeV, Au ions, 1 $\times 10^{14}$–1 $\times 10^{16}$ ions·cm$^{-2}$, RT |
| Xia et al. [83]         | Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi, Al$_{0.5}$CoCrFeNi | FCC, FCC + B2, B2 + A2 | 3 MeV, Au ions, 1 $\times 10^{14}$–1 $\times 10^{16}$ ions·cm$^{-2}$, RT |
| Yang et al. [332]       | Al$_{0.5}$CoCrFeNi              | FCC   | 3 MeV, Au ions, 6 $\times 10^{15}$ ions·cm$^{-2}$, 250–650 °C |
| Zhou et al. [333]       | Al$_{0.5}$CoCrFeNi (x = 0–2)    | FCC + BCC | 1 MeV, Kr$_{2+}$, - RT |
| Zhou et al. [334]       | Al$_{0.5}$CoCrFeNi, HfNbTaTiZrV | FCC, Amorphous | MeV Kr & 200 keV, electrons, 2 dpa, RT & 150 °C |
| Zhou et al. [335]       | HfNbTaTiZrV                     | BCC   | 1 MeV Kr$_{2+}$, - RT-150 °C |
| Moschetti et al. [336]  | HfNbTaTiZr                      | BCC   | 5 MeV, He$^{2+}$, 1.6 $\times 10^{12}$–4.4 $\times 10^{17}$ ions·cm$^{-2}$, s, 50 °C |
| Sadeghilardjani et al. [337] | HfTaTiZrV                   | BCC   | 4.4 MeV, Ni$_{2+}$, 1.08 $\times 10^{17}$ ion·cm$^{-2}$, RT |
| Li et al. [338]         | HfNbTiZr                        | BCC   | 1.5 MeV, He ions, 5 $\times 10^{15}$–1 $\times 10^{17}$ ions·m$^{-2}$, 700 °C |
| Karer et al. [339]      | TaTaIVZr, TaTaIVZr, TaTaIVNb   | BCC   | 2 MeV, V$^+$, 2.26 $\times 10^{15}$ ions·cm$^{-2}$, 500 °C |
| Wang et al. [340]       | ZrTiHfCuBe, ZrTiHfCuBeNi, ZrTiHfCuNi | Amorphous | 100 keV, He ions, 5.0 $\times 10^{17}$, 1.0 $\times 10^{18}$ & 2.0 $\times 10^{18}$ ions·cm$^{-2}$, RT |
3.1.1. Dislocation

Neutrons generated from fission/fusion reactors induce atomic displacements in structural materials and may introduce point defects. Development of interstitial and vacancy defects will change local lattice parameters of the original phase. This will lead to deterioration of structural materials, namely, hardening, phase instability, irradiation-induced segregation, irradiation-induced creep, volumetric swelling and H/He embrittlement [290,348,349]. For instance, Yang et al. [332] noticed various defects (i.e., dislocation loops, long dislocations and stacking-fault tetrahedra) were induced by irradiation of 3 MeV Au ion on Al$_{0.3}$CoCrFeNi but they did not observe void formation. They noticed that defect density decreased and defect size increased with the increasing temperature. Chen et al. [81] investigated the effect of irradiation of He$^+$ 275 KeV on FeCoNiCrTi$_{0.2}$ at 400 °C. They reported that high pressure He bubbles generated at the peak damage region and faulted dislocation loops (1/3<111>) formed. Perfect loops were rarely noticed and the size of the faulted loops was observed to be abnormally large. They also determined that the SFE and reported the upper limit for SFE at 400 °C for FeCoNiCr and FeCoNiCrTi$_{0.2}$ was estimated approximately to be ~80 mJ·m$^{-2}$ for the largest radius of a faulted loop of ~15 nm and ~32 nm at 500 °C. Hence, this showed that adding Ti significantly reduced the SFE.

Yang et al. [314] studied the irradiation behavior of CoCrNiFe with 3 MeV Ni$^{2+}$ ions at 500 °C. They investigated the defects (dislocation loops and void distribution) as a function of depth. The defects were found at the depth of 200–600 nm and 1100–2000 from the surface. Overall, the average defect diameter was measured to be <10 nm. Lu et al. [302] observed the irradiation behavior of FeNi, CoFeNi, CoCrFeNi and CoCrFeNiMn under 3 MeV Ni$^{2+}$ ions at 500 °C. In SEM images, they noticed faulted (1/3<111>) dislocation loops in all of these alloys as a result of irradiation. These loops increased with the increasing number of principal elements. Yang et al. [298] studied Al$_{0.3}$CoCrFeNi under 3 MeV Au ion irradiation. Average dislocation loop size was found to be ~12 nm at 250 °C and ~32 nm at 500 °C. Dislocation loop density was ~18 × 10$^{21}$ m$^{-3}$ at 250 °C and ~2 × 10$^{21}$ m$^{-3}$ at 500 °C. Hence, the average defect size increased and defect density decreased with irradiation temperature.

3.1.2. Hardness

Various researchers claimed that ion irradiation increased the hardness of the HEAs [306,307,326,327,329,337,343] although there are a few cases that reported some deviant behaviors [80,336,341]. HEAs showed better resistance to hardening by irradiation than stainless steels [326,329]. For instance, Sadeghilaridjani et al. [337] reported that the hardness of HfTaTiVZr increased by 20% under Ni$^{2+}$ ion irradiation, but under the same condition, the hardness of SS304 increased by around 50%. Such superior resistance to irradiation hardening was attributed to the reduced mobility of the point defects due to sluggish diffusion and self-healing ability. Similarly, Tolstolutskaya et al. [326] irra-
diated CrFeNiMn, Cr$_{0.18}$Fe$_{0.4}$Ni$_{0.28}$Mn$_{0.14}$ and Cr$_{0.18}$Fe$_{0.28}$Ni$_{0.27}$Mn$_{0.28}$ with 1.4 MeV and found out that the hardness increased by 22–45%. They also reported that the hardness of austenitic steels, such as X18H10T and SS316, almost doubled under similar irradiation conditions.

### 3.1.3. Phase Stability

The microstructure of a material under irradiation could be affected by formation of defect clusters, dislocation loops, stacking faults, precipitates, voids, or He bubbles. These phenomena may even alter local chemical compositions. Such microstructural changes could lead to deterioration in properties such as conductivity, ductility, fracture toughness or creep strength. Irradiation damage also can cause material swelling, irradiation-induced creep (IIC), irradiation-assisted stress corrosion cracking or irradiation growth. The attractive properties of the HEAs typically come from specific phases acquired with suitable compositions. Moreover, there are various potential intermetallics compositions present in HEAs. Therefore, the phase stability analysis becomes of utmost importance. Taking advantage of recent advancement in microstructure characterization technologies, most of the studies mentioned in Table 5 focused on the phase stability and microstructural changes after ion irradiation. Phase stability after irradiation was observed for several HEAs such as CoCrFeNiMn, CoCrFeNiPd and high entropy metallic glasses such as ZrTiHfCuBe and ZrTiHfCuBeNi [340] and Al$_x$CoCrFeNi (x = 0.1–1.5) [82,83]. Such high phase stability was mainly attributed to sluggish diffusion and high configurational entropy that kinetically restrains precipitation by reducing thermodynamic driving force. Moreover, Kumar et al. [327] reported that Fe$_{0.27}$Ni$_{0.28}$Mn$_{0.27}$Cr$_{0.18}$ alloy exhibited higher phase stability and higher resistance to radiation swelling and void formation in comparison to austenitic stainless steels. No micro-void formed in CoCrFeNi irradiated by 1.5 MeV Ni$^{2+}$ ion. Thanks to its superior interface stability during irradiation, AlCrMoNbZr/ (AlCrMoNbZr)N was considered a promising candidate for an accident-tolerant fuel cladding material [299].

On the other hand, Wang et al. [308] noticed a reduction in crystallinity resulted from He ion irradiation on CoCrFeCuNi alloy. Yang et al. [298] observed that precipitation of the L12 phase was suppressed due to irradiation-induced ballistic mixing at temperatures less than or equal to 500 °C. Additionally, precipitation of the B2 phase was favored at 650 °C due to improved diffusion by irradiation. They found that the swelling resistance of CoCrFeNiMn was less than 0.5% for temperatures up to 680 °C. They attributed the improved resistance to the void-induced swelling of these alloys to the complex arrangements of different atoms in the lattice structure. Gandy et al. [343] reported irradiation-induced phase transformation in SiFeVCrMo from the tetragonal sigma phase to BCC. This BCC phase formed at high temperatures as well as after ion irradiation. Under irradiation of 3 MeV Ni$^{2+}$ ion, NiCoFeCr maintained its phase stability; however, Al$_{0.12}$NiCoFeCr showed phase change. Al$_{0.12}$NiCoFeCr microstructure changed from single FCC to FCC matrix, nanoprecipitates (i.e., Ni$_3$Al) and ordered structure (i.e., L12) [318]. Irradiation of 3 MeV Ni$^{2+}$ ion on the CoCrFeNiMn alloy induced Mn depletion and Co/Ni enrichment at grain boundaries [303]. He et al. [309] studied the phase stability of CoCrFeNi, CoCrFeNiMn and CoCrFeNiPd under electron irradiation. They concluded that Cr, Fe, Mn and Pd elements were most likely to deplete and Co/Ni preferred to accumulate at defect clusters (i.e., dislocation loops). Atwani et al. [341] irradiated CrTaVW HEA with 1 MeV Kr$^{2+}$ ions at 800 °C. This HEA showed the segregation of Cr and V at the triple junction and grain boundaries after irradiation. Yang et al. [332] and Lu et al. [302] also noticed that Ni and Co tended to enrich, but Cr, Fe and Mn preferred to deplete at defect clusters in Al$_{0.1}$CoCrFeNi under 3 MeV Au ion irradiation.

### 3.1.4. Irradiation-Induced Creep (IIC)

Irradiation-induced creep is another important aspect since nuclear reactors operate at high temperatures as shown in Table 5. Jawaharram et al. [301] studied both thermal creep and irradiation-induced creep (IIC) of Cantor alloy (CoCrFeMnNi). For IIC measurement,
CoCrFeNiMn was irradiated with 2.6 Mev Ag$^{3+}$ ion in the temperature range of 23–650 °C. They reported that over the temperature range investigated, ICC was more dominant than thermal creep.

3.1.5. Swelling Resistance

Density reduction by volume increase due to formation of voids and defects is the mechanism behind material swelling. In order to maintain the structural integrity and mechanical strength, reasonably high swelling resistance is required for structural materials for nuclear applications. Several HEAs showed adequate swelling resistance under irradiation. Yang et al. [332] reported negligible void formation in Al$_{0.1}$CoCrFeNi under 3 MeV Au ion irradiation. Moreover, Yang et al. [314] noticed that dislocation loops and void distribution varied as a function of depth in CoCrFeNi under 3 MeV Ni$^{2+}$ ion irradiation. Jin et al. [306] suggested that compositional complexity, including the number and type of components, be taken into consideration to improve swelling resistance. They concluded that adding Fe and Mn would be more effective to reduce swelling than adding Co and Cr. Their results showed that NiCoFeCrMn exhibited 40 times higher swelling resistance than Ni.

3.1.6. Self-Healing

HEAs showed the ability to absorb and heal radiation-induced damages. Self-healing of HEAs was explained by Egami et al. [350] in detail. Xia et al. [83] reported that defect clustering in disordered FCC or BCC happened in smaller size than in the ordered B2 phase. They attributed this effect to the reduced defect mobility and large atomic stress in the disordered phases which might have led to self-healing. Patel et al. [344] investigated the phase stability of V$_{2.5}$Cr$_{1.2}$WMoCo$_{0.04}$ under 5 MeV Au$^+$ ion irradiation. They noticed that 96% of the BCC phase of the as-cast alloy remained intact up to the irradiation dose of 42 displacement per atom (dpa). The remaining 4% converted to another BCC phase with a little larger lattice parameter. Their energy dispersive X-ray (EDX) analysis detected no element segregation. Such phase stability was attributed to the self-healing capability [350]. Tong et al. [305] concluded that the local lattice distortion could be relaxed by lattice expansion with low dose of irradiation. Sellami et al. [316] irradiated CoCrFeNi with 1.5 MeV and 21 MeV Ni Ions. They reported elastic strain values of ~0.035% and 0% for 1.5 MeV Ni ion and 21 MeV Ni ion irradiations, respectively. They suggested that the complex composition of HEAs induced higher chemical disorder that reduced the mobility of defects generated by low energy (i.e., 1.5 MeV Ni) ion irradiation; therefore, a small elastic strain of ~0.035% was obtained. Furthermore, when low energy irradiation CoCrFeNi alloy was further subjected to high-energy (i.e., 21 MeV Ni) irradiation, elastic strain relaxation (resulted into ~0% elastic strain) was observed which could be attributed to the mechanisms such as defect annealing, recombination and rearrangement.

3.1.7. Miscellaneous

Lu et al. [80] reported an unexpected decrease in lattice parameter by 0.676% after He ion irradiation on Ti$_2$ZrHfV$_{0.5}$Mo$_{0.2}$. Similarly, TiZrNbHfTa showed insignificant changes in hardness but a considerable increase in UTS and YS without loss of ductility [336]. Li et al. [328] noticed an increase in electrical resistivity of Fe$_{0.27}$Mn$_{0.27}$Ni$_{0.28}$Cr$_{0.18}$ after neutron irradiation.

3.2. Wear Behavior

The wear properties of HEAs were studied mostly with pin/ball on a disc set up with antagonist materials such as Al$_2$O$_3$, steels (i.e., SKH51, GCr15, 100Cr6), Si$_3$N$_4$, SiC, ZrO$_2$, 1Cr18Ni9Ti, BN, inconel-718 and WC. For lubrication, mostly dry conditions were used but some studies also used H$_2$O$_2$, deionized water and acid rain (pH = 2). Previously, Tsai and Yeh et al. [351], Kasar et al. [352], Senkov et al. [67], Sharma et al. [16], Zhang et al. [37], Li et al. [42], Menghani et al. [353] and Ayyagari et al. [354] discussed the wear behaviors
of HEAs. In this review, we will analyze the tribological studies of HEAs in terms of HEAs content variation, particle reinforcement, media and nitriding/carburizing/sulfurizing, temperature effects and oxide formation. Table 6 provides the details of the compositions, microstructures, methods and results (i.e., wear rate or wear resistance, hardness, friction coefficient) of the wear studies performed so far on HEAs.

Table 6. Wear studies of HEAs.

| Source                          | Composition            | Microstructure | Method, Medium, Antagonist Material, Temperature, Wear Rate |
|---------------------------------|------------------------|----------------|-------------------------------------------------------------|
| Joseph et al. [355]             | CoCrFeNiMn            | FCC            | Pin-on-disc, dry, Al₂O₃, 600–800 °C, RT, 0.5 × 10⁻⁶–3.8 × 10⁻⁴ mm³ N⁻¹ m⁻¹ |
| Wang et al. [356]               | CoCrFeNiMn            | FCC            | Ball-on-disc, MoS₂-oil lubrication, GCr15, RT-140 °C        |
| Xiao et al. [357]               | CoCrFeNiMn            | FCC            | Ball-on-flat, dry, WC-Co, RT, 0.5 × 10⁻⁴–5.4 × 10⁻⁴ mm³ N⁻¹ m⁻¹ |
| Jones et al. [358]              | CoCrFeNiMn            | FCC            | Rotary tribometer, γ, γ, 0.5 × 10⁻⁶ mm³ N⁻¹ m⁻¹             |
| Zhu et al. [359]                | CoCrFeNiMn, CoCrFeNiMnV, CoCrFeNiMnNb, CoCrFeNiMnNbV | FCC + HCP (Laves) + α | Ball-on-disc, dry, Si₃N₄, RT, 1.85 × 10⁻⁵–6.39 × 10⁻⁵ mm³ N⁻¹ m⁻¹ |
| Deng et al. [360]               | CoCrFeNiMoₓ (x = 0–0.3) | FCC            | Ball-on-disc, dry, GCr15, RT, 0.33 × 10⁻³–0.53 × 10⁻³ mm³ N⁻¹ m⁻¹ |
| Lindner et al. [361]            | CoCrFeNiMn, CoCrFeNi | FCC            | Ball-on-disc, dry, Al₂O₃, RT                                |
| Sha et al. [362]                | (CoCrFeNiMn)N         | FCC + BCC      | Ball-on-disc, dry, ruby, RT, 1 × 10⁻⁷–1.4 × 10⁻⁸ mm³ N⁻¹ m⁻¹ |
| Xiao et al. [363]               | CoCrFeNiMnCₓ (x = 0–1.2) | FCC            | Ball-on-disc, dry, Si₃N₄, RT, 0.47 × 10⁻⁵–6.5 × 10⁻⁵ mm³ N⁻¹ m⁻¹ |
| Zhu et al. [277]                | CoCrFeNiMn + TiN-Al₂O₃ | FCC + TiN     | Ball-on-disc, dry, 440C steel, RT                           |
| Cheng et al. [364]              | CoCrFeNiMn, AlₓCoCrFeNiMn, AlCoCrFeNiMn | FCC + BCC | Ball-on-disc, dry, Si₃N₄, RT-800 °C, 0.5 × 10⁻⁴–3.8 × 10⁻⁴ mm³ N⁻¹ m⁻¹ |
| Joseph et al. [365]             | CoCrFeNiMn, Alₓ₀.₃CoCrFeNi, Alₓ₀.₄CoCrFeNi, AlCoCrFeNi | FCC + BCC | Pin-on-disc, dry, Al₂O₃, 25 & 900 °C                          |
| Liu et al. [366]                | CoCrFeNiMn + Y₂O₃ (particles) | FCC + Y₂O₃ | Ball-on-disc, dry, GCr15, RT                                |
| Wang et al. [367]               | (CoCrFeMnNi)₉₅Ti₁₅ | FCC + BCC      | Ball-on-disc, dry, Si₃N₄, RT-800 °C, 4 × 10⁻⁶–2.23 × 10⁻⁵ mm³ N⁻¹ m⁻¹ |
| Zhang et al. [368]              | CoCrFeNi + (Ag or BaF₂/CaF₂) | FCC            | Ball-on-disc, dry, Inconel-718, RT, -4 × 10⁻⁵–40 × 10⁻⁵ mm³ N⁻¹ m⁻¹ |
| Geng et al. [369]               | CoCrFeNi                | FCC            | Pin-on-disc, vacuum (4 Pa) & air, Inconel 718, RT, 0.6 × 10⁻⁴–0.4 × 10⁻⁴ mm³ N⁻¹ m⁻¹ |
| Zhang et al. [370]              | CoCrFeNi + (graphite or MoS₂) | FCC            | Ball-on-disc, dry, Si₃N₄, RT-800 °C, -1 × 10⁻⁵–2.3 × 10⁻⁵ mm³ N⁻¹ m⁻¹ |
| Zhou et al. [371]               | CoCrFeNiMoₓ₀.₈₅, Alₓ₀.₅CoCrFeNi | FCC            | Slurry jet test-rig, HCl+NaCl, -40 °C, -                   |
| Zhang et al. [372]              | CoCrFeNiMo                | FCC            | Ball-on-disc, dry, - RT                                    |
| Huang et al. [373]              | FeCoCrFeNiSᵢₓ         | FCC + BCC      | Ball-on-disc, dry, GCr15, RT                                |
| Cui et al. [374]                | CoCrFeNiMo, Sulfurized at 260 °C for 2 h | FCC + Fe₅/MoS₂ film | Pin-on-disc, dry, GCr15, RT                                |
| Li et al. [375]                 | CoCrFeNiMoₓ₀.₂        | FCC            | Ball on disc, dry, GCr15, RT, 3.9 × 10⁻⁴–5.4 × 10⁻⁴ mm³ N⁻¹ m⁻¹ |
| Source            | Composition                                                                 | Microstructure                        | Method, Medium, Antagonist Material, Temperature, Wear Rate                      |
|-------------------|------------------------------------------------------------------------------|----------------------------------------|---------------------------------------------------------------------------------|
| Ji et al. [376]   | CoCrFeNiCu + 2% MoS₂                                                          | FCC + MoS₂ (particles)                 | Ball-on-disc, dry, Si₃N₄, RT                                                  |
|                   | CoCrFeNiCu + 5% MoS₂                                                          | FCC + MoS₂ (particles)                 |                                                                                 |
|                   | CoCrFeNiCu + 20% WC                                                          | FCC + WC (particles)                   |                                                                                 |
|                   | CoCrFeNiCu + 50% WC                                                          | FCC + WC (particles)                   |                                                                                 |
|                   | CoCrFeNiCu + 80% WC                                                          | FCC + WC (particles)                   |                                                                                 |
| Verma et al. [377]| CoCrFeNiCu₅₆ (x = 0–1)                                                        | FCC                                    | Pin-on-disc, dry, -, RT & 600 °C, \(-1.3 \times 10^{-4}-2.5 \times 10^{-3} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\) |
| Liu et al. [378]  | CoCrFeNiB₅ (x = 0.5–1.5)                                                     | FCC + Borides                          | Roller friction wear tester, dry, W₃C₅V, RT                                |
| Jiang et al. [379]| CoCrFeNiNb₅ (x = 0–1.2)                                                       | FCC + HCP (Laves)                      | Ball-on-disc, dry, BN, RT                                                    |
| Yu et al. [380]   | CoCrFeNi + WC                                                                 | FCC                                    | Pin-on-disc, dry, Si₃N₄, RT-800 °C, \(-1.8 \times 10^{-4}-9 \times 10^{-4} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\) |
| Liu et al. [381]  | Co₁₀Cr₁₀Fe₉₀Mo₁₀₀ + graphene nanoplatelets (0.2–0.8 wt%)                    | FCC                                    | Ball-on-plate, dry, GCr15, RT                                               |
| Wang et al. [382] | Co₁₀Cr₁₀Fe₉₀Mo₁₀₀ + WC (10 wt%)                                              | FCC + WC (Laves)                       | Ball-on-disc, dry, Si₃N₄, RT                                                |
| Derimov et al. [383]| (CoCrCuTi)₁₀₀₋ₓM₀ₓ (x = 5–10)                                               | FCC + BCC                              | Ball-on-disc, dry, GGr15, RT                                               |
| Guo et al. [384]  | CoCrFeNiCu₀.₂₂Ti (Ti or Cr)ₙ (x = 0–1.5)                                     | FCC + TiC                              | Brooks sliding friction & wear tester, dry, RT                               |
| Zhang et al. [385]| (CoCrFeNiTi₀.₃₅C₀.₅₅) (x = 3–12 wt%)                                        | BCC + Cr₂₃C₈ + TiC                     | ML-100 friction and wear tester, -, -, RT                                   |
| Erdoğan et al. [386]| CoCrFeNiTi₁₀₀₋ₓA₀ₓ₅                                                        | FCC                                    | Ball-on-disc, dry, WC                                                     |
| Liu et al. [387]  | CoCrFeNiMo₁₀₅                                                               | FCC                                    | Pin-on-disc, dry, -, GCr15, RT                                             |
| Moazzen et al. [388]| CoCrFeNiMo₁₀₅ (x ≥ 0.3)                                                       | FCC                                    |                                                                                           |
| Yang et al. [389] | CoCrFeNiMo₅₅ (x ≥ 1)                                                         | FCC + σ                               |                                                                                           |
| Li et al. [390]   | CoCrFeNi₂₅V₁₀₅Ti₂ (x = 0.5–1.25)                                             | BCC + (Co,Ni)Ti₂                      | Ball-on-disc, dry, Si₃N₄, 4.4 \times 10^{-5}. 37.5 \times 10^{-3} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\     |
| Islak et al. [391]| CrFeNiMoTi                                                                  | FCC                                    | Ball-on-flat, dry, 100Cr6, RT                                              |
| Wen et al. [392]  | CrCoNiTiV                                                                    | FCC + BCC + TiO                       | HT-1000 tribometer, -, WC, RT & 600 °C                                    |
| Wang et al. [393] | CuNi₅₅Ti₂                                                                  | BCC                                    | CJS111A wear tester, dry, -, RT                                             |
| Cheng et al. [394]| (Fe₂₅Co₂₅Ni₂₅K₂₅S₂₅)₁₀₀₋ₓNbₕ (x = 0–4 wt%)                                 | BCC + HCP (Laves) + FCC               | Ball-on-disc, dry, GCr15, RT                                               |
| Yadav et al. [395]| (CuCrFeTiZn₂₀₋ₓPb₂₀₋ₓBu₂₀₋ₓ (x = 0–10)                                     | FCC + BCC + Pb (particles)            | Ball-on-disc, dry, -, SAE 52100, RT, 1.17 \times 10^{-5}-50 \times 10^{-5} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\     |
| Gou et al. [396]  | CoCrFeNi + WC + Mo₂C + NbC                                                    | FCC                                    | Ball-on-disc, dry, GCr15, 700 °C                                           |
| Yadav et al. [397]| (CuCrFeTiZn₂₀₋ₓPb₂₀₋ₓBu₂₀₋ₓ (x = 0–10)                                     | FCC + BCC                            |                                                                                           |
| Cui et al. [398]  | Al₅₀CoCrFeNi₅₅ (x = 0–0.75)                                                  | FCC + BCC                            | MDW-02 abrasive wear tester, RT                                               |
| Gwalani et al. [399]| Al₀₅₀CoCrFeNi                                                             | FCC + B2                              |                                                                                           |
| Chen et al. [400] | Al₀₅₀CoCrFeNi                                                                | FCC + BCC                            |                                                                                           |
| Du et al. [401]   | Al₀₂₅CoCrFeNi                                                                | FCC                                    | Universal wear testing machine, dry, Si₃N₄, 20–600 °C, \(-1.5 \times 10^{-4}-3.5 \times 10^{-4} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\) |
| Chen et al. [402]| Al₀₅₀CoCrFeNi                                                                | FCC + BCC                            | Ball-on-block, deionized water & acid rain (pH = 2), seawater, GCr15, RT, 1.58 \times 10^{-4}-6.52 \times 10^{-4} \text{mm}^{3} \text{N}^{-1} \text{m}^{-1}\ |
| Source               | Composition                                      | Microstructure | Method, Medium, Antagonist Material, Temperature, Wear Rate |
|----------------------|--------------------------------------------------|----------------|----------------------------------------------------------|
| Ji et al. [409]      | Al<sub>3</sub>CoCrFeNi                          | FCC            | Jet erosion testing machine, water and 15 wt% SiO<sub>2</sub> particles (350–600 mm), RT |
| Haghdadi et al. [404] | Al<sub>3</sub>CoCrFeNi, AlCoCrFeNi             | FCC, BCC       | Scratch testing, dry, RT                                  |
| Fang et al. [405]    | Al<sub>3</sub>CoCrFeNi                          | FCC            | Pin-on-disc, dry, 900 °C                                  |
| Wu et al. [406]      | Al<sub>3</sub>CoCrFeNi                          | FCC            | Ball-on-block, dry and deionized water, Si₃Ni₄, RT, -0.2 x 10⁻⁶ - 1.86 x 10⁻⁴ mm²N⁻¹ m⁻¹ |
| Nair et al. [407]    | Al<sub>0.1</sub>CoCrFeNi, AlCoCrFeNi, Al₃CoCrFeNi | FCC + BCC (B2) | Ball-on-disc, dry, WC, RT                                 |
| Kumar et al. [408]   | Al<sub>0.5</sub>Co<sub>0.5</sub>CrFeNi, Al<sub>0.5</sub>Co<sub>0.5</sub>CrFeNi (x = 0–1) | FCC + BCC (B2) + A₂ + σ | Pin-on-disc, demineralized water & (demineralized water + 3.5 wt% NaCl), EN-31, RT, 0.81 x 10⁻⁴ – 1.86 x 10⁻⁴ mm³N⁻¹ m⁻¹ |
| Mu et al. [409]      | AlCoCrFeNi                                       | BCC + FCC      | Ball-on-disc, dry, Si₃Ni₄, RT                             |
| Wu et al. [410]      | AlCoCrFeNi                                      | BCC            | Ball-on-disc, dry, Si₃Ni₄, RT                             |
| Zhao et al. [411]    | Al<sub>0.5</sub>CoCrFeNi                          | FCC + BCC      | Ball-on-disc, dry, deionized water + 0.5 wt% NaCl, RT, -2 x 10⁻⁵–7.5 x 10⁻⁵ mm³N⁻¹ m⁻¹ |
| Kumar et al. [412]   | Al<sub>0.5</sub>Co<sub>0.5</sub>CrFeNi (x = 0–0.5) | FCC + BCC      | Pin-on-disc, engine oil (SAE Grade/20W-40), EN-31 steel, RT, 2.1 x 10⁻⁵–11 x 10⁻⁵ mm³N⁻¹ m⁻¹ |
| Li et al. [413]      | Al<sub>0.5</sub>CoCrFeNiCu<sub>0.5</sub>Si<sub>x</sub> (x = 0–0.5) | FCC + BCC1 + BCC2 | - , - , CG₁₅₃, RT, 0.9 x 10⁻⁶–1.19 x 10⁻⁶ mm³N⁻¹ m⁻¹ |
| Li et al. [414]      | (AlCoCrFeNi)<sub>100-x</sub>Nb<sub>x</sub> (x = 0–30 wt%) | FCC + BCC      | Reciprocating tester, dry, N₄Si₃s, RT                    |
| Kafexhiu et al. [415] | AlCoCrFeNi<sub>2</sub>                           | BCC + FCC      | Ball-on-plate, dry, 100Cr6 steel, RT, 7 x 10⁻²–11 x 10⁻⁵ mm³N⁻¹ m⁻¹ |
| Miao et al. [416]    | AlCoCrFeNi<sub>2</sub>                           | FCC (L.12) + BCC (B2) | Ball-on-disc, dry, Al₂O₃/Si₃Ni₄/SiC/Gr15, RT-900 °C, -1 x 10⁻⁴–4.2 x 10⁻⁴ mm³N⁻¹ m⁻¹ |
| Ye et al. [417]      | AlCoCrFeNi<sub>2</sub> + TiC (0–15 wt%)         | FCC + B2 + TiC | MM-200 wear testing machine, dry, RT                      |
| Wang et al. [418]    | (AlCoCrFeNi)N                                    | BCC + nitrides (AlN, CrN, Fe₄N) | Ball-on-block, dry, deionized water & acid rain (pH = 2), Si₃Ni₄, RT, 2.8 x 10⁻³–7.5 x 10⁻⁶ mm³N⁻¹ m⁻¹ |
| Liu et al. [419]     | AlCrCuFeNi<sub>2</sub>                           | BCC            | Ball-on-block, dry, simulated rainwater & deionized water, Si₃Ni₄, RT, 2.163 x 10⁻³–0.23 x 10⁻³ mm³N⁻¹ m⁻¹ |
| Kong et al. [420]    | Al<sub>1.5</sub>CrCuFeNi<sub>2</sub>            | BCC            | MMS-2A roller friction wear tester, dry, RT               |
| Malati et al. [421]  | AlCrCuFeNi                                       | FCC + BCC      | Ball-on-disc, dry, SiC, RT                                |
| Wang et al. [422]    | Al<sub>1.5</sub>CoCuFeNi<sub>2</sub>            | FCC + BCC      | Ball-on-block, dry, deionized water & acid rain (pH = 2), Si₃Ni₄, RT, 1 x 10⁻⁴–12 x 10⁻⁴ mm³N⁻¹ m⁻¹ |
| Xiao et al. [423]    | Al<sub>1.5</sub>CoCrFeNiSi<sub>1.1</sub> (x = 0.5–1.5) | FCC + BCC      | Ball-on-flat, distilled water, WC-12Co, RT, 6.7 x 10⁻⁶–5.5 x 10⁻⁵ mm³N⁻¹ m⁻¹ |
| Liu et al. [424]     | AlCoCrFeNiSi<sub>1.1</sub> (0–0.5)               | BCC            | Pin-on-disc, dry, ZrO₂, RT, 1.3 x 10⁻⁴–5.1 x 10⁻⁴ mm³N⁻¹ m⁻¹ |
| Hsu et al. [425]     | Al<sub>1.5</sub>CoCrFeNiCu<sub>2</sub>B<sub>x</sub> (x = 0–1) | FCC + boride precipitates | Pin-on-disc, dry, Al₂O₃, RT                               |
| Chen et al. [426]    | Al<sub>0.5</sub>CoCrFeNiCu<sub>2</sub>Ti<sub>x</sub> (x = 0–0.2) | FCC            | Pin-on-disc, dry, Al₂O₃, RT                               |
| Lobel et al. [427]   | AlCoCrFeNiTi                                    | BCC            | Ball-on-disc, dry, Al₂O₃, RT                              |
| Source               | Composition                                      | Microstructure             | Method, Medium, Antagonist Material, Temperature, Wear Rate |
|---------------------|-------------------------------------------------|----------------------------|-------------------------------------------------------------|
| Lobel et al. [426]  | AlCoCrFeNiTi                                    | BCC                        | Ball-on-plate, dry, 100Cr6 Steel, RT                         |
| Wu et al. [427]     | AlCoCrFeNiTi(x = 0.5–1) AlCoCrFeNiTi(x = 1.5) AlCoCrFeNiTi(x = 2) | FCC + BCC FCC + BCC + Ti2Ni FCC + BCC + Ti2Ni + ordered BCC | Cavitation erosion tests, Distilled water+ 3.5 wt% NaCl, RT |
| Erdogan et al. [428]| Al0.14Co0.88Fe0.9Ni0.1Ti0.18 (x = 0–0.5, y = 0–0.5) | FCC                        | Ball-on-disc, dry, WC, RT, 0.25 × 10⁻¹⁰–7.8 × 10⁻¹⁴ mm²·N⁻¹·m⁻¹, 0.25 × 10⁻¹⁰–7.8 × 10⁻¹⁴ mm²·N⁻¹·m⁻¹ |
| Xin et al. [429]    | Al0.3Co0.7Fe0.5Ni0.25Ti0.5 + TiC                | FCC                        | Ball-on-disc, dry, Si3N4, RT, 0.3 × 10⁻⁵–12.6 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Gouveia et al. [430]| Al0.2Co0.8Fe0.5Ni0.25Ti                         | FCC                        | Ball-on-plate, dry, AISI 52,100 steel, RT, 1.6 × 10⁻⁸–7.5 × 10⁻⁹ mm²·N⁻¹ |
| Chuang et al. [431]| Al0.4Co0.6Fe0.5Ni0.25Ti                         | FCC                        | Pin-on-disc, dry, SHK511 steel, RT, 4 × 10⁻⁴–1.8 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Liu et al. [432]    | AlCoCrFeNiTi0.8 | BCC + B2                   | Ball-on-disc, dry, Si3N4, RT, 1.36 × 10⁻⁸–6.96 × 10⁻⁶ mm²·N⁻¹·m⁻¹, 0.7 × 10⁻⁸–6 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Yu et al. [433]     | AlCoCrFeNiTi0.35 | BCC1 + BCC2                | Pin-on-disc, H2O2, SiC & ZrO2, RT                             |
| Lobel et al. [434]  | AlCoCrFeNiTi0.5 | BCC (A2 + B2)              | SRV-Tribometer, dry, Al2O3, 22–900 °C                         |
| Chen et al. [435]   | Al1-xCoCrFeNiTi | BCC                        | Pin-on-disc, Dry, Al2O3, RT-500 °C                             |
| Yu et al. [436]     | AlCoCrFeNiTi0.5 | AlCoCrFeNiCu                | Pin-on-disc, dry, Si3N4                                      |
| Yu et al. [437]     | AlCoCrFeNiCu | AlCoCrFeNiTi0.5 | FCC + BCC1 | BCC1 + BCC2         | Pin-on-disc, H₂O₂, Cr18Ni9Ti steel & ZrO₂/SiC ceramic, RT |
| Jin et al. [438]    | AlCoCrFeNiCu | FCC + BCC                    | Ball-on-disc, dry, WC, 200–800 °C                             |
| Zhu et al. [439]    | AlCoFeNiCu + TiC (10–30 wt%) | FCC + BCC                   | Ball-on-disc, dry, Si3N4, 20–600 °C, -0.1 × 10⁻⁵–6.5 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Wu et al. [440]     | Al1-xCoCrFeNiCu | Al1-xCoCrFeNiCu | FCC | FCC + BCC | FCC + BCC | Pin-on-disc, dry, SKH-51 steel, RT |
| Yan et al. [441]    | AlCoCrFeNiSi + Ti (C, N) | BCC + FCC                    | Ball-on-disc, dry, GCr15, RT, -                                |
| Li et al. [442]     | AlCoCrFeNi + Ti (C, N) + TiB₂ | FCC                        | Ball-on-disc, dry, WC-6Co, 200–800 °C, 2.69 × 10⁻⁴–8.66 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Kumar et al. [443]  | AlCoCrCuFeNiSi0.3 | AlCoCrCuFeNiSi0.3 | FCC + BCC | FCC + BCC + σ | Pin-on-disc, dry, -, RT, - |
| Xiong et al. [444]  | Al0.2Co1.8CrFeNi1.3Ti0.5 | FCC                        | Pin-on-disc, dry, Si3N4, 25–800 °C, 1.21 × 10⁻⁵–6.7 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Karakaş et al. [445]| Al0.2Co1.2CrFeNi1.2Mn1.25Ni1.1 | FCC + BCC | FCC + BCC | FCC + BCC | FCC + BCC | Pin-on-disc, dry, Si3N4, 25–800 °C, 3.12 × 10⁻⁶–12.59 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Xin et al. [446]    | Al0.2Co1.5CrFeNi1.5Ti0.5(Si0.125 + Cx) (x = 0) | FCC                        | Pin-on-disc, dry, Si3N4, 25–800 °C, 1.67 × 10⁻⁶–2.22 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Zhao et al. [447]   | AlCrCoFeNiTa₂, (x = 0–1) | BCC | Pin-on-disc, 3.5% NaCl & air, Si3N4, RT, 1.67 × 10⁻⁶–2.22 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Ghanbariha et al. [448]| AlCoCrFeNi + ZrO₂ | FCC + BCC | FCC + BCC | FCC + BCC | FCC + BCC | Pin-on-disc, dry, WC, RT, 1.11 × 10⁻³–2.52 × 10⁻⁵ mm²·N⁻¹·m⁻¹ |
| Li et al. [449]     | AlCrFeNiCu | AlCrFeNiCu (x = 0–0.5) | FCC | FCC | FCC | FCC | Pin-on-disc, dry, GCr15, RT, 6.64 × 10⁻⁷–2.26 × 10⁻⁵ mm³·N⁻¹·m⁻¹ |
| Cai et al. [450]    | AlCrTiV, AlCrTiVSi | FCC                        | Nanoindenter G200, dry, GCr15 &Al2O3, RT, -                  |
| Chandrakar et al. [451]| AlCoCrCuFeNiSi | (x = 0–0.9) | FCC | FCC + BCC | FCC + BCC | FCC + BCC | Pin-on-disc, dry, -, RT, - |
| Source               | Composition                        | Microstructure           | Method, Medium, Antagonist Material, Temperature, Wear Rate                               |
|----------------------|-----------------------------------|--------------------------|-----------------------------------------------------------------------------------------|
| Erdogan et al. [452] | AlCrFeNi5Si                      | BCC                      | Ball-on-disc, dry, WC, RT, -                                                            |
| Duan et al. [453]    | AlCoCrFeNiCu                     | -                        | Pin-on-disc, H2O2, Si3N4, RT                                                            |
| Chen et al. [454]    | Al0.5CoCrFeNiCuVx (x = 0–0.2)    | FCC                      | Pin-on-disc, dry, Al2O3, RT, $1 \times 10^{-4}$–$2.7 \times 10^{-4}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Gu et al. [455]      | Al$_3$Mo$_{0.5}$NbFeTiMoN$_2$ (x = 1–2) | BCC                      | Pin-on-disc, dry, Al$_2$O$_3$, RT                                                      |
| Hsu et al. [456]     | AlCoCrFeNi$_5$Mo$_{0.5}$ (x = 0.6–2) | BCC + $\sigma$          | Pin-on-disc, dry, SKH51 steel, RT                                                      |
| Liang et al. [457]   | AlCrFe$_2$Ni$_2$W$_{0.5}$Mo$_{0.75}$ | BCC                      | Ball-on-disc, deionized water, Al$_2$O$_3$, RT, $-5 \times 10^{-8}$–$22 \times 10^{-8}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Qui et al. [458]     | Al$_2$CoCrFeCuTiNi$_4$ (x = 0–2)  | FCC + BCC                | Tribometer, -, RT                                                                      |
| Kanyane et al. [459] | AlTiSiMoW                         | BCC + TiSi$_2$ (ordered FCC) | Ball-on-disc, dry, stainless steel, RT                                                  |
| Huang et al. [460]   | AlTiSiVCr                         | BCC + Ti(V)$_3$Si$_2$ precipitates | Ball-on-disc, dry, Cr$_{15}$ steel, RT, $2 \times 10^{-5}$–$2.5 \times 10^{-4}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Zhang et al. [461]   | AlTiSiVNi                         | B2 (NiAl) + (Ti,V)$_3$Si + TiN | Ball-on-disc, dry, Si$_3$N$_4$, RT & 800 °C                                             |
| Lin et al. [462]     | AlCoCrNiW                         | W + AlNi + Cr$_{13.5}$Fe$_{42.5}$C$_6$ BCC | Pin-on-disc, dry, Al$_5$I$_{2100}$, RT                                                 |
| Yadav et al. [463]   | AlCrFeMnV$_{10}$ (AlCrFeMn)$_3$V$_3$B$_{10}$ | BCC                      | Ball-on-disc, dry, SAE 52,100 steel, RT, $1.02 \times 10^{-7}$–$7.02 \times 10^{-8}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Bhardwaj et al. [464] | AlTiZrNbHfF | BCC                      | Pin-on-disc, dry, Cr$_{15}$ steel, RT                                                  |
| Zhao et al. [465]    | AlNbTaZr$_x$ (x = 0.2–1)          | BCC + HCP                | Ball-on-disc, dry, Si$_3$N$_4$, RT, $1.85 \times 10^{-4}$–$2.41 \times 10^{-4}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Tuten et al. [466]   | TiZrHfNbTa                        | Amorphous                | Ball-on-disc, dry, Al$_2$O$_3$, RT                                                     |
| Pole et al. [467]    | TiZrHfTaV, TiZrTaVV               | BCC                      | Ball-on-disc, dry, Si$_3$N$_4$, RT-500 °C, $-1 \times 10^{-8}$–$8 \times 10^{-8}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Ye et al. [468]      | TiZrHfNb                          | BCC                      | Nano-scratch, dry, diamond indenter, RT                                                |
| Pogrebnjak et al. [469] | (TiZrHfV)$_3$N | FCC                      | Ball-on-disc, dry, Cr$_{15}$ steel, 20 °C                                             |
| Gong et al. [470]    | TiZrHfBeCu, TiZrHfBeNi, Ti$_{20}$Zr$_{20}$Hf$_{30}$Be$_{20}$Cu$_{10}$Ni$_{10}$, Ti$_{13.5}$Zr$_{11.25}$Ni$_{10}$Be$_{22.5}$Cu$_{12.5}$ | Amorphous                | Nano-scratch, dry, diamond indenter, RT                                                |
| Zhao et al. [471]    | TiZrNbBeCu                       | Amorphous                | Nano-scratch, dry, diamond indenter, RT                                                |
| Jhong et al. [472]   | (TiZrNbCrSi)$_3$C$_3$ (x = 36.7–87.8 at.%) | FCC                      | Ball-on-disc, dry, 100Cr$_3$ steel, RT, $0.2 \times 10^{-3.3}$–$3.3 \times 10^{-6}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Mathiou et al. [473] | TiZrNbMoTa                        | BCC + HCP                | Ball-on-disc, dry, 100Cr$_3$ steel, Al$_2$O$_3$, RT, $0.154 \times 10^{-1}$–$0.199 \times 10^{-1}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Petroglou et al. [474] | MoTa$_{0.25}$NbVTi (x = 0.25–1) | BCC                      | Ball-on-disc, dry, 100Cr$_3$ steel, Al$_2$O$_3$, RT, $0.19 \times 10^{-6}$–$0.38 \times 10^{-6}$ g N$^{-1}$ m$^{-1}$ |
| Poula et al. [475]   | MoTaNbVW                         | BCC                      | Ball-on-disc, dry, 100Cr$_3$ steel & Al$_2$O$_3$, RT, $1.05 \times 10^{-4}$–$4.89 \times 10^{-4}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Poula et al. [476]   | MoTaNbVW                         | BCC                      | Ball-on-disc, dry, 100Cr$_3$ steel & Al$_2$O$_3$, RT, $1.05 \times 10^{-4}$–$4.89 \times 10^{-4}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Poula et al. [477]   | MoTaNbVTi                        | BCC + hexagonal C14 Laves + cubic C15 laves | Ball-on-disc, dry, 100Cr$_3$ steel, Al$_2$O$_3$, RT                                    |
| Alvi et al. [478]    | MoTaWVCu                        | BCC                      | Ball-on-disc, dry, ES2100 steel & Si$_3$N$_4$, RT, 600 °C, $2.3 \times 10^{-7}$–$2.5 \times 10^{-3}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Hua et al. [479]     | Ti$_x$ZrNbTaMo (x = 0.5–2)       | BCC                      | HSR-2M tester, dry, Si$_3$N$_4$, RT, $2.22 \times 10^{-7}$–$2.42 \times 10^{-7}$ mm$^3$ N$^{-1}$ m$^{-1}$ |
| Gu et al. [480]      | Ni$_{1.5}$CrFeTi$_{2.5}$Mo$_x$ (x = 0–0.25) | BCC + FCC                | Ball-on-disc, dry, Al$_2$O$_3$, RT, $7.99 \times 10^{-2}$–$2.7 \times 10^{-7}$ mm$^3$ |

Table 6. Cont.
3.2.1. Content Variation

Tribological studies on HEAs are mostly conducted with variations in content of one element and finding the optimum concentration of the element for minimum wear rate [360,363–365,371,377–380,383,384,386,390,394,395,397,398,407,408,412,421,422,424,427,428,431,440,443,454,455,463,470,472,474]. Furthermore, the effect of Al content on wear properties and hardness has been studied more than any other metal. For instance, the wear resistance of CoCrFeNiCu [440], CoCrFeNiTi0.5 [386], CoCrFeNiSi [421] and CoCrFeNiMn [364] was improved with Al addition. This was primarily ascribed to the oxide formation and hardness increase due to phase transformation, grain refinement and precipitation. Similarly, Cui et al. [398] studied the effect of Al, CoCrFeNiMn (x = 0–0.75) coatings on 4Cr5MoSiV alloys. This alloy experienced FCC to FCC + BCC transition and grain refinement when Al content increased, which in turn raised the hardness from 224 HV to 344 HV. The wear weight loss of 4Cr5MoSiV coated with Al,CoCrFeNiMn with x = 0, 0.25, 0.5 and 0.75 was measured to be 6.0 mg, 4.1 mg, 3.2 mg and 1.1 mg, respectively. Under the similar wear condition, the amount of wear weight loss of the uncoated 4Cr5MoSiV was around 10 mg. Meanwhile, Gu et al. [455] analyzed Al,Mo0.5NbFeTiMn2 (x = 1–2) coating on Q235 steel and found that the microhardness and wear were positively correlated to the Al content. As the Al content increased, the grain size reduced and the hardness and wear resistance increased. Kumar et al. [412] studied the wear behavior of Al1xFeCrNiCo, (x = 0–1) under oil lubricated conditions. The hardness of this alloy reduced with the increasing Ti content on the wear performance of the Al alloy. This resulted from the enhanced plastic deformation of the FCC matrix. Additionally, Al and Ti formed some beneficial intermetallic phases such as AlNi and AlNiTi, which improved the hardness and wear resistance. With a Ni content of x = 0.8–1.2, the CoCr-like phases and abrasive wear due to thermal softening at higher temperatures. In contrast, the wear rate of CoCrFeNiCu (x = 1) decreased at higher temperatures. This was attributed to the self-lubricating effect of Cu. Cu also tended to form protective oxide layer which prevented direct metal to metal contact [481]. Hsu et al. [423] showed the improved wear resistance of Al0.5CoCrFeNiCuB. 

Increasing the Cu content from x = 0 to x = 1 reduced the wear rate of CoCrFeNiCu, (x = 0–1) alloy from 2.3 × 10^-5 to 1.7 × 10^-5 mm^3.N^-1.m^-1 at RT and from 2.5 × 10^-5 to 1.3 × 10^-5 mm^3.N^-1.m^-1 at 600 °C [377]. The wear rate of CoCrFeNi (x = 0) increased with the increasing temperature. This resulted from the enhanced plastic deformation and abrasive wear due to thermal softening at higher temperatures. In contrast, the wear rate of CoCrFeNiCu (x = 1) decreased at higher temperatures. This was attributed to the self-lubricating effect of Cu. Cu also tended to form protective oxide layer which prevented direct metal to metal contact [481].
showed the hardness of 736 HV and the maximum wear resistance of 1.76 mm−3N−1m−1 which was higher than that of SKD61 cold-work mold (1.39 m−3m−3) and SUJ2 bearing (1.52 m−3m−3). The improved wear resistance was attributed to the boride formation. Similarly, Liu et al. [378] analyzed the wear resistance of CoCrFeNiBx (x = 0.5–1.5). They noticed that the increased boron reduced the wear volume from 19 down to 4.5 mm3. The hardness increased from 550 to 1025 HV0.2 as well with increasing boron, which was attributed to the increase in the fraction of the hard boride phase with B content. The maximum hardness was measured to be 1025 HV0.2 for FeCoCrNiB0.5 which was higher than that of Q245R steels. Kumar et al. [408] reported that the wear rate of Al0.4FeCrNiCo0.5 (x = 0–1 mol) increased with the increasing Co content from 0.81 × 10−4 mm3N−1m−1 (x = 0) to 1.86 × 10−4 mm3N−1m−1 (x = 1). This was attributed to the hardness decrease form 377 HV (x = 0) to 199 HV (x = 1). As hardness decreases the wear resistance also decreases according to the Archard’s law [482]. Chen et al. [454] reported how the variation of V in Al0.5CoCrFeNiCuVx (x = 0–1) affected the wear performance. Here, increasing V from x = 0.6 to x = 1.2 improved the wear resistance by 20%. However, increasing V beyond x = 1.2 did not show a considerable effect. This was in line with the hardness results of this HEA. The hardness increased while x increased from x = 0.4 to x = 1 exhibiting its maximum value at x = 1. The optimal V addition was suggested to be in the range of x = 1–1.2 to improve the wear resistance.

Furthermore, improvements in the wear resistance of CoCrFeNiNb (x = 0–1.2) [379], (CuCrFeTiZn)1−xPbx (x = 0.05–0.2) [395], CoCrFeNiCu (x = 0–1) [377], MoTa5NbVTi (x = 0.25–1) [474] were reported. Wu et al. [410] analyzed the wear resistance of the AlCoCrFeNiTi5 (x = 0, 0.5) alloy. They noticed that AlCoCrFeNi had the plate-like disordered Fe-Cr rich BCC phase embedded in the ordered Al-Ni rich BCC phase. AlCoCrFeNiTi0.5 had the similar microstructure, but the Ti introduced honeycomb-like interdendrites which could play an important role in hindering dislocation motion. As a result, the hardness of this alloy increased from 887 HV0.5 to 1147 HV0.5. Additionally, the atomic radius of Ti was 1.76 Å which was larger than those of Ni (1.49 Å) and Al (1.18 Å). Hence, adding Ti to the Al-Ni-Ti rich phase might have resulted in lattice expansion or distortion.

Liu et al. [422] varied the concentration of Si in AlCoCrFeNiSi (x = 0–0.5) coatings and found out that the microhardness of the coatings was linearly proportional to the Si concentration. The wear rate reduced significantly from ~5.2 × 10−4 to ~1.3 × 10−4 mm2N−1m−1 with the increasing Si content. This was attributed to the formation of wear-resistant SiO and SiO2 layers on the wear surface. Huang et al. [373] studied FeCoCrNiSi (x = 0–1) and found out that increasing Si content improved the hardness and wear resistance by promoting the transformation of FCC to BCC. Its microhardness increased significantly from 89.52 HV (x = 0) to 653.71 HV (x = 1) and the wear track depth decreased from 22.14 µm (x = 0) to 5.29 µm (x = 1).

Hsu et al. [456] reported that in AlCoCrFeNiMo0.5 (x = 0.6–2), the wear resistance (at 500 °C) of x = 2 was much lower than that of x = 1.5 although both had similar hardness values. This was attributed to the fact that the x = 2 alloy formed a thicker oxide layer than x = 1.5 and more loose oxides abraded away from the surface. By this reasoning, the Fe content of x = 0.6–1 was recommended as over x = 1.5–2 for optimum wear resistance. Qiu et al. [458] described that the wear resistance of Al2CoCrFeCuTiNi (x = 0–2) first increased with the Ni content (i.e., x = 0–1) and then decreased for x = 1–2. However, the hardness monotonically increased with the increasing Ni content from ~900 HV (x = 0) to ~1100 HV (x = 2). They said that toughness and brittleness also affect the wear resistance of this alloy apart from the hardness. In this alloy, when Ni is added from x = 0–1 at% then the wear resistance increases because hardness increases. For Ni content x = 1–2 at%, the wear resistance starts to reduce because the material becomes brittle and wear happens through small brittle fracture on the surface. Beng et al. [360] analyzed the CoCrFeNiMo (x = 0–0.3) alloy and noticed that CoCrFeNi had the FCC phase. The FCC phase was maintained even after Mo addition, but the lattice parameter (denoted by a) altered (aCoCrFeNi = 3.5733 Å, aCoCrFeNiMo0.1 = 3.6016 Å, aCoCrFeNiMo0.3 = 3.5854 Å) due to atomic size misfit between Mo
and other constituents. The atomic size of Mo (1.363 Å) is distinctively larger than those of Co (1.251 Å), Cr (1.249 Å), Fe (1.241 Å) and Ni (1.246 Å) [93]. This Mo-induced lattice distortion increased the hardness from 415.2 to 465.9 HV and reduced the specific wear rate from $0.59 \times 10^{-3}$ to $0.33 \times 10^{-3}$ mm$^3$N$^{-1}$m$^{-1}$.

Yadav et al. [397] reported the influence of Pb and Bi content in (CuCrFeTiZn)$_{100-x}$Pb$_x$ (CuCrFeTiZn)$_{100-x}$Bi$_x$ on their wear properties. Here, reductions in wear rate achieved by adding Pb and Bi were 21% and 25%, respectively. This was attributed to the fact that the soft dispersoids acted as lubricants [483]. Li et al. [413] studied Al$_{0.8}$CoCrFeNi$_{0.5}$Si$_x$ ($x = 0$–0.5) coatings. They found the specific wear rate in the range of $1.19 \times 10^{-6}$–$8.99 \times 10^{-6}$ mm$^3$N$^{-1}$m$^{-1}$, the hardness in the range of 479–592 HV which was almost 5 to 7 times higher than the hardness of substrate 5083 Al alloy (79 HV$^{0.2}$).

Cheng et al. [394] noticed that the wear resistance and hardness of (Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$ (B$_{0.2}$Si$_{0.3}$,3,0,25)$_{100-x}$Nb$_x$ ($x = 0$–2) coating (on Q235 steel) increased with the Nb content from $x = 0$ to $x = 2$. The measured hardness values were 587.1 HV$_{0.1}$ and 821.5 HV$_{0.1}$ for Nb content of $x = 0$ and $x = 1$, respectively. This HEA had the FCC microstructure and the grain size was refined from 3.5 µm ($x = 0$) to 1 µm ($x = 2$). This grain refinement helped to increase the hardness and wear resistance. The wear rates were measured to be $3.64 \times 10^{-6}$ and $1.42 \times 10^{-6}$ mm$^3$N$^{-1}$m$^{-1}$ for Nb content of $x = 0$ and $x = 2$, respectively. Jones et al. [358] reported that CoCrFeNiMn showed a remarkably low wear rate of $10^{-6}$ mm$^3$N$^{-1}$m$^{-1}$ with the hardness of 1.6 GPa. Zhu et al. [359] added V and Nb in equiatomic ratio into the CoCrFeNiMn (FCC phase) coating and studied the wear resistance. The measured wear rate ranged $1.85 \times 10^{-5}–6.39 \times 10^{-5}$ mm$^3$N$^{-1}$m$^{-1}$ and the hardness varied in the range of 145–948.5 HV. The measured hardness values of CoCrFeNiMn, CoCrFeNiMnNb, CoCrFeNiMnV and CoCrFeNiMnNbV was 145 HV, 609 HV, 621 HV and 948 HV, respectively. Additionally, the wear rate of CoCrFeNiMn reduced from $6.39 \times 10^{-5}$ to $1.85 \times 10^{-5}$ mm$^3$N$^{-1}$m$^{-1}$ (for CoCrFeNiMnNbV) when both Nb and V were added. These results could be attributed to the fact that Nb and V promoted the precipitation of the Laves phase (FeNb intermetallic) and the sigma phase (CoFeV intermetallics) into the FCC matrix, which increased the hardness. Moreover, Nb and V promoted the formation of protective oxide layer on the worn surface and reduced the amount of adhesive and abrasive wear.

3.2.2. Particle Reinforcement

Hard particle reinforcement into HEAs matrix is another approach employed to improve the tribological properties. For example, adding WC into Co$_{10}$Cr$_{10}$Fe$_{0.9}$Mn$_{0.1}$ [382], adding TiN-Al$_2$O$_3$ into CoCrFeNiMn [277], adding NbC into AlCoCrFeNi [272], adding TiC into CoCrCuFeNi$_{0.2}$ [384] and AlCoCrFeNi$_{1.1}$ [416] improved the tribological performances. Such improved wear resistance was largely attributed to the increased hardness, the restrained effect of adherence abrasion, solution strengthening or microstructure refinement. Adding 10 wt% TiC particles into AlCoFeNiCu reduced the wear rate by the factor of 8 [439]. Adding TiB$_2$ into (AlCrFeMnV)$_{90}$Bi$_{10}$ resulted in $-95\%$ decrease in wear rate due to the cumulative effect of matrix, reinforcement, refined grains and improved hardness [463]. Gou et al. [396] used CoCrFeNi (reinforced with NbC) as a metal binder for Ti (C,N)-based cermet. The hardness and fracture toughness were measured to be 1853 HV and 9.93 MPa·m$^{1/2}$, respectively. The minimum wear rate was as low as $4.14 \times 10^{-6}$ mm$^3$N$^{-1}$m$^{-1}$, which was attributed to the reduced grain size and improved hardness due to the solid solution strengthening effect of the HEA binder.

Ji et al. [376] improved the tribological behavior of CoCrFeNiCu by embedding MoS$_2$ (2–5 wt%) and WC (20–80 wt%) particles. Here, adding 2% MoS$_2$ improved the wear resistance and decreased the friction coefficient but adding 5% MoS$_2$ increased the wear rate. Adding WC continuously improved the wear resistance and 80% WC yielded the lowest wear rate. Likewise, the wear resistance of CrMnFeCoNi was improved considerably by adding 25 wt% Y$_2$O$_3$ thanks to grain boundary strengthening, Orowan looping and load transfer effect [366]. Zhou et al. [189] reported that (FeCoCrNi)$_{1-x}$ (WC)$_x$ ($x = 3$–11 wt%)
alloy consisted of a FCC matrix and W/Cr rich carbides. The hardness increased with the WC content from 603 HV (for (FeCoCrNi)_{0.97} (WC)_{0.03}) to 768 HV (for (FeCoCrNi)_{0.89} (WC)_{0.11}). This was attributed to the hard WC particles and carbides precipitation into the FCC matrix. This material showed adhesive and abrasive wear. Most of the wear was caused by the debonding of WC particles.

Liu et al. [381] introduced graphene nanoplatelets into Fe_{50}Mn_{30}Co_{10}Cr_{10}. Adding 0.2% Graphene nanoplatelets promoted self-lubricating properties because the coefficient of friction (COF) decreased by 62%, which in turn increased the wear resistance. Further addition of graphene did not have much impact on COF or self-lubricating properties. Similarly, Zhang et al. [370] added self-lubricating particles (i.e., nickel coated graphite and MoS_{2} powder) into the CoCrFeNi matrix using SPS. This composite had four phases (i.e., FCC, graphite, MoS_{2} and Nickel) and showed very low wear rates (~10^{-5} mm^{2}·N^{-1}·m^{-1}) for the temperature range of RT-800 °C. After wear testing the worn surfaces were analyzed by Raman and EDS. They observed that graphite and MoS_{2} particles were accumulated on the worn surface for temperatures RT-800 °C. They indicated that these particles helped reduce abrasive/adhesive wear. Moreover, at 600 and 800 °C, the significantly increased oxygen concentration on the worn surface facilitated the formation of a smooth glaze layer of oxide (i.e., Cr_{2}O_{3}, Fe_{2}O_{4} etc.) and reduced the wear rate. Likewise, Zhang et al. [368] added solid lubricants (i.e., Ag, BaF_{2}/CaF_{2} eutectic) into CoCrFeNi. This composite showed the better wear resistance (wear rates~10^{-5} mm^{2}·N^{-1}·m^{-1}) than PM304 (wear rate~10^{-4} mm^{2}·N^{-1}·m^{-1}) [484] for the temperature range of RT-800 °C. The improved wear resistance was attributed to the solid lubricants (help to minimize friction and wear on the surface) and oxide layer (Cr_{2}O_{3}, Fe_{2}O_{4}, etc.) formation on the worn surface.

3.2.3. Use of Media and Heat Treatment

Media used during wear experiments have been proved to have considerable impact on the mechanism and rate of material removal. For example, Duan et al. [453] reported that AlCoCrFeNiCu showed more wear weight loss of ~1.2 mg when in H_{2}O_{2} lubricant and ~0.8 mg when oil was used as a media. This was because friction coefficient was lower when oil was used as a media. FeCoCrAlNiTi_{2} showed the best wear resistance in distilled water and the worst in NaCl solutions [427]. Xiao et al. [421] noticed that the wear rates of Al_{x}CoCrFeNiSi (x = 0.5–1.5 mol) reduced significantly from 5.5 × 10^{-5} to 1.6 × 10^{-6} mm^{3}·N^{-1}·m^{-1} with the increasing Al content under dry conditions. However, when water was used as a lubricant, the addition of Al content did not have a considerable impact on the wear rate. Geng et al. [369] studied the wear behavior of CoCrFeNiMn in vacuum and air in the temperature range of RT-800 °C. They found the wear rate varied in the range of 1.3 × 10^{-4}–8 × 10^{-4} mm^{3}·N^{-1}·m^{-1}. The elements of this HEA oxidized as temperature increased. At elevated temperatures, oxides (i.e., CoO, CoO, CoCrO_{3}, Co_{2}O_{3}, Fe_{2}O_{3}, Cr_{2}O_{3} and MoO_{3}) formed on the worn surface. From RT to 400 °C, the wear rate was higher in air than in vacuum due to peeling of the formed loose oxides. However, at 600 °C and 800 °C, robust oxides were formed, and these oxides served as a protective layer and improved the wear resistance resulting in the lower wear rate both in air than in vacuum.

The effect of heat treatment on the wear behavior of Al_{0.5}CoCrFeNi was studied by Gwalani et al. [399]. They hot rolled (at 700 °C), annealed (at 1150 °C) and isothermally aged their alloy at 700 °C for 1, 4, 20, 40, 80 h. As the aging time increased, the B2 phase precipitated in the FCC matrix. These precipitates increased the hardness from ~250HV to ~302 HV, the flow stress from 668 MPa to 1352 MPa, the UTS from 1157 MPa to 1503 MPa. The wear rate of this alloy was found to be sensitive to aging. The wear rate for the sample without aging was 11 × 10^{-5} mm^{3}·N^{-1}·m^{-1}. After 80 h of aging, the wear rate reduced to 1.8 × 10^{-5} mm^{3}·N^{-1}·m^{-1}. Meanwhile, Kong et al. [419] studied the Al_{1.5}CrCuFeNi_{2} alloy with different arc melting parameters. They varied welding current during arc melting as 410A, 455 A, 480 A, 505 A and 550 A to get different superheating. As a result, the hardness of the alloy increased from ~555 to ~625 HV when current increased from 410A to 480A.
Consequently, the wear rate reduced from ~0.95 to ~0.65 mg·mm⁻². However, when the current was further increased up to 550 A, the hardness reduced to ~580 HV and the wear rate increased to ~0.72 mg·mm⁻². Here, the wear resistance of the HEA was found to be directly proportional to the hardness. Malati et al. [263] improved the wear resistance of AlCrCuFeNi by heat treating at temperatures 800, 950 and 1100 °C. The microhardness increased from 310 to 381 HV when heat treated at 800 °C but decreased to 365 HV when heat treated at 1100 °C. Heat treatment at 1100 °C homogenized the microstructure and promoted grain coarsening. This alloy initially had the FCC phase alone. Upon heat treatment at 800 and 950 °C, the B2 phase precipitated and the hardness increased. When the alloy was heat treated at 1100 °C, some of the B2 precipitates decomposed and grain coarsened, which decreased the hardness. This explains why the wear resistant first increased and later reduced with the heat-treating temperatures. The observed wear weight was 0.002 g without heat treatment, and 0.001 g, 0.0007 g and 0.0016 g with heat treatment at 800 °C, 950 °C and 1100 °C respectively.

3.2.4. Nitriding/Carburizing/Boronizing/Sulfurization

Nitriding, carburizing, boronizing and sulfurizing have been used to improve the tribological behavior of HEAs. Pogrebniak et al. [469] measured the wear rate of nitrided TiZrHfVNb (i.e., (TiZrHfVNb)N) and compared with the wear resistance of (AlCoCrFeNiCu)N and steel 45. The measured wear rates for (TiZrHfVNb)N and steel 45 were 6.5 × 10⁻⁴ and 3.0 × 10⁻⁴ mm³·N⁻¹·m⁻¹, respectively. Wu et al. [406] used the pack-boronizing method to improve the wear resistance of Al₁₀CoCrFeNi alloy. A 17.3–57.9 μm boronized layer, composed of (Co, Fe, Ni)B, CrB, C₂B, formed on the surface. The boronizing time varied from 2 h to 8 h. The hardness of the alloy increased gradually with the increasing boronizing time from 201 HV (0 h) to 1398 HV (8 h) and the wear rate reduced from 1.89 × 10⁻⁴ to 0.22 × 10⁻⁴ mm³·N⁻¹·m⁻¹.

Xiao et al. [363] investigated the effects of carbon content in CoCrFeNiMnCₓ (x = 0–1.2). The hardness was increased monotonically from 327.8 HV (x = 0) to 566.4 HV (x = 1.2) with carbon addition. They found out that the wear rates of CoCrFeNiMn and CoCrFeNiMnC₀.₆ were 6.5 × 10⁻³ mm³·N⁻¹·m⁻¹ and 0.47 × 10⁻³ mm³·N⁻¹·m⁻¹, respectively. This alloy was comprised of the FCC (Co, Ni rich) phase and the Cr- and C- rich M7C3 carbide phases. The porosity increased with the increasing carbon content. Initially, the hardness increased until the volume fraction reached x = 0.6 due to introduction of M7C3 carbide and solid solutions of C atoms in the FCC phase. As the amount of carbon further increased, the hardness started to fall due to a significant increase in porosity. Meanwhile, Zhang et al. [385] found out that the wear rate and hardness of (CoCrFeNi₃Tiₐₓ)Cₓ (x = 3–12 wt%) coating were in the range of 12–24 mg·N⁻¹·m⁻¹ and 300–950 HV, respectively. This coating microstructure was made of BCC solid solution, C₂₃C₆ and TiC phases. Increase in carbon content raised the hardness due to carbides precipitates, but the wear resistance with x = 12 wt% was inferior to that with x = 6 wt% due to reduced ductility.
3.2.5. Comparison with Conventional Materials

Various HEAs have shown superior wear resistance than the commercial wear resistant materials in the following parentheses: CuCoNiCrAl0.5FeB (SUJ2 wear-resistant steel) [423], MoTaWNbV (Inconel-718) [475,476], TiZrHfNb (Nb and C103 (Nb54H2Ti)) [468], CoCrFeNiMo (HF XP-58) [372], Al0.8CrCoFeNi (AZ31) [411] and Al0.2Co1.8CrFeNi1.5Ti (AISI 52,100 steel and Inconel 713) [430]. Al3CoCrFeNi exhibited better wear resistance than 17–7 PH stainless steel [403], which was attributed to the precipitation of Cr2Ni2, CuNiSiTiZr coating improved the wear resistance of TC11 by a factor of 2.5 [393]. This improvement in hardness (and wear resistance) was resulted from the solid solutioning, precipitation and nanocomposite strengthening effects. Similarly, Al1CoCrFeNi (x = 0.1–3) showed 3 to 23 times higher wear resistance than SS316L steel [407]. This higher wear resistance was attributed to high hardness of Al1CoCrFeNi (~436–624 HV) than SS316L steel (~227 HV).

Many researchers formed coatings of HEAs on several commercial materials to improve their wear resistance. Zhang et al. [461] coated Ti-6Al-4V with AlTiSiVNi. They discovered that the wear resistance of AlTiSiVNi was 4 to 5 times higher than that of Ti-6Al-4V at RT and 800 °C. The hardness of AlTiSiVNi was also found to be in the range of 1151–1357 HV which was 4–6 times than that of Ti-6Al-4V. The increase in hardness and wear resistance of AlTiSiVNi was attributed to the dispersion strengthening with (Ti,V)5Si3 and solution strengthening with NiAl. Huang et al. [460] coated Ti-6Al-4V with AlTiSiVCr. The microstructure of AlTiSiVCr had hard silicides (Ti,V)5Si3 dispersed into the BCC matrix. The dispersed silicides improved the wear resistance of this HEA by lowering abrasive/adhesive wear. The hard BCC phase of the silicides also resisted crack propagation. While the specific wear rate for Ti-6Al-4V was ~6.5 × 10⁻⁵ to 9 × 10⁻⁵ mm³·N⁻¹·m⁻¹, that for TiVCrAlSi ranged 2 × 10⁻⁵ to 2.6 × 10⁻⁵ mm³·N⁻¹·m⁻¹. This showed that TiVCrAlSi could become a promising anti-wear coating material for Ti-6Al-4V.

Islak et al. [391] improved the wear resistance of AISI 1040 steel with CrFeNiMoTi coating. They found that the hardness and wear rate of CrFeNiMoTi coating were ~450 HV on 0.3 and ~2.732 × 10⁻³–3.952 × 10⁻³ mm³·N⁻¹·m⁻¹, respectively. Meanwhile, the hardness and wear rate for AISI 1040 were ~200 HV on 0.3 and ~8.125 × 10⁻³–9.455 × 10⁻³ mm³·N⁻¹·m⁻¹, respectively. Gu et al. [455] analyzed Al1Mo0.5NbFeTiMn2 (x = 1–2) coating and found that the microhardness and wear resistance were positively related to the Al content. The increasing Al content reduced the grain size and increased both the hardness and wear resistance. The hardness of Al2Mo0.5NbFeTiMn2 was measured to be 1098.5 HV on 0.2 which was 5 times higher than that of Q235 steel (~200 HV on 0.2). The high hardness was attributed to the microstructure composed of the BCC solid solution and (Nb,Ti)C carbides. The wear rates of Al2Mo0.5NbFeTiMn2 and Q235 steel were measured to be ~0.3 mg·m⁻¹ and ~0.9 mg·m⁻¹, respectively. Liang et al. [457] deposited AlCrFe2Ni2W0.2Mo0.25 coating on Q235 steel used for ocean engineering equipment. This coating had the hardness of 630 HV and the specific wear rate in the range of 10.13 × 10⁻⁶–23.89 × 10⁻⁶ mm³·N⁻¹·m⁻¹. The coating showed superior wear resistance to Q235 and SUS304 in deionized water, 3.5 wt% NaCl solution and artificial seawater media. They also reported that the electrochemical corrosion of AlCrFe2Ni2W0.2Mo0.25 (corrosion rate of 0.173 mm/yr) was less than both SUS304 (corrosion rate of 0.333 mm/yr) and Q235 (corrosion rate of 0.44 mm/yr).

3.2.6. Higher Temperatures Wear Resistance

HEAs have also exhibited promising tribological behaviors and thermal stability at elevated temperatures. Jin et al. [438] studied the characteristics of AlCoCuFeNi coating at temperatures up to 800 °C. This coating was mainly composed of Fe rich FCC and Cu rich BCC phases and showed good thermal stability without any phase transformation until 780 °C. Mainly Al2O3, Cr2O3, Fe2O3 and CuO were present in the oxide layer. Meanwhile, the weight losses of NiCrCoTiV at RT and 600 °C were measured to be 3.7 ± 0.1 mg and 3.5 ± 0.1 mg respectively [392]. The wear weight loss of 304L stainless steel was 7.7 ± 0.2 mg at RT. Fang et al. [405] used Al0.5CoCrFeNi as a metal binder for Ti (C,N)-
The hardness, fracture toughness and flexural strength of this cermet were 1137 HV, 6.46 MPa m^{1/2} and 761 MPa at 1000 °C respectively. This superior high-temperature performance was attributed to the hindrance of the slip system and higher oxidation resistance of the HEA binder. The wear resistance of this Al_{0.3}CoCrFeNiTi (wear groove width = 49 µm) was also better than that of the conventional Ni-Co (wear groove width = 172 µm) metal binder.

Yadav et al. [395] concluded that the wear resistance of (CuCrFeTiZn)_{100-x}Pb_x improved due to oxide formation at the surface at elevated temperatures (800–1000 °C). Because these oxides made a layer on the surfaces of the HEA which helped to avoid metal to metal contact, hence reduced the material removal rate. Moreover, Chen et al. [400] said that annealed Al_{0.6}CoCrFeNi showed the wear resistance higher than GCr15 by a factor of three at 600 °C. They also attributed this improved wear resistance to the formation of oxides on the surface. In this case, Fe_2O_3, Cr_2O_3, Al_2O_3 and Al(OH)_3 formed on the worn surfaces of Al_{x}FeCrNiCo_x and reduced the wear rate. Alvi et al. [478] studied wear behavior of AlCoCrFeNi in the temperature range of RT-600 °C. They noticed that the oxidation of Cu (into CuO) causes wear rate decrease at 400 °C. Joseph et al. [365] analyzed wear behavior of Al_{x}CoCrFeNi in the temperature range of RT-900 °C. The wear resistance of this HEA surpassed that of Inconel 718 at 900 °C. Here again, the wear resistance was enhanced at the higher temperatures thanks to the oxide layer formed at the contact interface.

Researchers also showed that the wear rate of HEAs increased initially and then decreased with temperature [380, 401, 434, 467]. It was claimed that initially loose oxides formed on the surface and lower the wear rate up to moderately high temperatures. As temperature increased higher, such loose oxides were damaged by thermal softening and therefore, the wear rate increased. In contrast, other studies reported the opposite behaviors. The wear resistance initially decreased up to moderately high temperatures and then increased at higher temperatures. This behavior was attributed to the formation of a thicker and more robust oxide layer on the worn surface which reduced the area of direct metal to metal contact. For instance, Pole et al. [467] studied the wear resistance of refractory HEAs, TiZrHfTaV and TiZrTaVW in the temperature range of RT-500 °C. The hardness of these HEAs (6–8.1 GPa) was found to be larger than two times that of SS304. The measured wear rate of these alloy was in the range of 0.5 × 10^{-5}–8 × 10^{-4} mm^{3}·N^{-1}·m^{-3}. Various oxides, such as ZrO_2, TiO_2, Ta_2O_5, V_2O_5, HfO_2 and WO_3, formed on the worn surface. The wear rate increased until 150 °C due to the formation of a delicate oxide layer at the worn surface. However, as the temperature increased further from 150 to 500 °C, a strong protective oxide layer formed on the worn surface and the wear rate reduced for both refractory HEAs. Similar findings were reported by Lobel et al. [434] for the wear resistance of AlCoCrFeNiTi_{0.5} coating in the temperature range of RT-900 °C. The depth of wear increased from ~62 µm at RT to ~82 µm at 500 °C, and afterwards it decreased down to ~50 µm at 900 °C. As the temperature increased, a loose oxides layer formed on the worn surface. The measured hardness, fracture toughness and flexural strength of this cermet were 1137 HV, 6.46 MPa m^{1/2} and 761 MPa at 1000 °C respectively. This superior high-temperature performance was attributed to the hindrance of the slip system and higher oxidation resistance of the HEA binder. The wear resistance of this Al_{0.3}CoCrFeNi (wear groove width = 49 µm) was also better than that of the conventional Ni-Co (wear groove width = 172 µm) metal binder.

Some HEAs showed fluctuation in wear resistance due to microstructural transformations at elevated temperatures. The wear rate of Al_{0.6}TiCrFeCoNi increased from RT to 300 °C and reduced from 300 °C to 500 °C, mainly due to phase transformation (i.e., the formation of sigma-CrFe) at higher temperatures [435]. Such phase transformation at higher temperatures could also significantly reduce fracture toughness. The wear resistance of (CoCrFeMnNi)$_{85}$Ti$_{15}$ increased from RT to 400 °C and then decreased...
thereafter [367]. Miao et al. [415] studied the wear behavior of AlCoCrFeNi\textsubscript{2.1} with different antagonist materials (i.e., Al\textsubscript{2}O\textsubscript{3}, Si\textsubscript{3}N\textsubscript{4} and GCr15). They also analyzed wear rate of the same composition for a temperature range of RT-900 °C against SiC. The wear rate of AlCoCrFeNi\textsubscript{2.1} against Al\textsubscript{2}O\textsubscript{3}, Si\textsubscript{3}N\textsubscript{4} and GCr15 is \(\approx 42 \times 10^{-5}\), \(\approx 37 \times 10^{-5}\), \(\approx 32.5 \times 10^{-5}\) mm\textsuperscript{3}\cdot N\textsuperscript{-1}\cdot m\textsuperscript{-1}. The hardness of Al\textsubscript{2}O\textsubscript{3}, Si\textsubscript{3}N\textsubscript{4} and GCr15 is 2300, 1500 and 680 HV, and the wear rate of AlCoCrFeNi\textsubscript{2.1} increased with the hardness of the antagonist material. Moreover, the wear rate of AlCoCrFeNi\textsubscript{2.1} (against SiC) increased from \(\approx 75 \times 10^{-6}\) to \(\approx 140 \times 10^{-6}\) mm\textsuperscript{3}\cdot N\textsuperscript{-1}\cdot m\textsuperscript{-1} when temperature increased from RT to 900 °C. This increased wear rate with temperature was attributed to the thermal softening.

Joseph et al. [355] noticed that when the wear resistance of CoCrFeNiMn was examined at high temperatures, ultrafine grains and the sigma phase formed at the contact surface, which resulted in the improved wear resistance.

4. Summery and Future Direction

This review covers recent advances in the development and manufacturing of HEAs and their performances under extreme environments such as nuclear and wear applications. The HEAs were tabulated based on manufacturing methods, irradiation responses and wear performances.

The most widely used method for HEAs manufacturing was arc melting due to its simplicity, when the idea of HEAs was conceptualized. However, recently AM processes (SLM, EBM and DED) have gained interest since they may potentially provide more freedom in shape and in properties by changing process parameters. However, there are some issues that need to be resolved. For instance, low productivity, formation of micro-level defects such as pores or unfused particle boundaries, residual stresses, composition shift due to selective evaporation of constituents with lower vapor pressure, lack of standards for quality evaluation, high initial investment cost and more. With these issues being gradually resolved, AM could make a powerful and versatile manufacturing method to fabricate application-specific HEAs with desired properties for some compositions. Most of the AM HEAs are studied in the as-cast state (after AM). In terms of HEAs characterization, most studies focused on the microstructure using SEM, tensile behavior and hardness. Structural characterization at a smaller length scale, using transmission electron microscope, would be helpful to better understand the structural evolution under various loadings. In order to identify the effect of manufacturing methods and explore more applications for HEAs, more research is needed on creep properties, dislocation behaviors, deformation microstructures, compressive strength, fatigue and more.

Structural materials for next generation nuclear reactors must survive high energy irradiation at high temperatures with reasonable service life. Similar to other structural metals, high energy irradiation on HEAs often induces microstructural changes which in turn deteriorated their mechanical properties including hardness, swelling or embrittlement. For a number of cases of HEAs, their compositional complexity hinders such microstructural degradation and results in superior irradiation resistance compared to other conventional alloys. This makes them promising candidate materials for nuclear applications. However, more studies must be conducted on the irradiation behaviors of HEAs to better understand their applicability to the next generation nuclear reactors. To date, very little is known about HEAs phase diagrams and equilibrium phases. In addition, the defect generation and movement mechanism as a result of irradiation are not clearly understood.

HEAs have also demonstrated superior tribological performances over a wide range of temperatures from RT to high temperatures in comparison to the commercial materials (i.e., Steel, Inconel, Ti-6Al-4V, Q235, SUS304 etc.). Moreover, the wear behavior of HEAs is affected by composition, particle reinforcement, media, nitriding/carburizing/sulfurizing treatments, temperature and oxides formation. Most of the wear studies are on cantor alloy or its derivatives; therefore, more elements and combinations are needed to be explored to further understand potential candidates for wear applications. For the wear
resistant applications where the weight is not a critical factor, refractory high entropy alloys would make a good option thanks to their high hardness [466–480]. Most of these alloys are equimolar. The equimolar ratios are probably a good point to start but they might not be the best to get the highest potential out of the particular element composition. Non-equimolar refractory HEAs are worth more exploration.

HEAs are being researched for more than a decade and are not yet commercially available. One reason could be because HEAs could not be manufactured with most widely used processes suitable for mass production (i.e., casting, molding etc.). Arc melting (most popular manufacturing method for HEAs) is limited to manufacture laboratory-scale samples for testing. Recently, a number of attempts have been made to fabricate HEAs using AM techniques. Some AM HEAs showed the improved mechanical properties. However, there are still many issues that need to be addressed for AM to be used for mass production of HEAs.

Overall, there are still a great number of possible HEAs compositions that are to be studied. Apart from cantor alloy, the characteristics of the majority of other HEAs have not been investigated well enough for safe practical applications. These studies cannot be used to generalize characteristics for HEAs but we can take them as screening efforts. Moreover, it is not practical to perform all the characterization studies on all these compositions. Therefore, it would be more reasonable to use simulations and material informatics to screen compositions before experimental studies instead of using trial and error.

Author Contributions: Conceptualization, S.S. and J.L.; methodology, S.S.; software, S.S.; validation, J.L. and S.S.; formal analysis, S.S.; investigation, S.S.; resources, S.S.; data curation, S.S.; writing—original draft preparation, S.S.; writing—review and editing, J.L.; visualization, S.S.; supervision, J.L.; project administration, J.L.; funding acquisition, J.L. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: The authors declare that the data supporting the findings of this study are available within the article.

Conflicts of Interest: The authors declare no conflict of interest.

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