A high-entropy silicide: \((\text{Mo}_{0.2}\text{Nb}_{0.2}\text{Ta}_{0.2}\text{Ti}_{0.2}\text{W}_{0.2})\text{Si}_2\)

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**A B S T R A C T**

A high-entropy metal disilicide, \((\text{Mo}_{0.2}\text{Nb}_{0.2}\text{Ta}_{0.2}\text{Ti}_{0.2}\text{W}_{0.2})\text{Si}_2\), has been successfully synthesized. X-ray diffraction (XRD), energy dispersive X-ray spectroscopy (EDX), and electron backscatter diffraction (EBSD) collectively show the formation of a single high-entropy silicide phase. This high-entropy \((\text{Mo}_{0.2}\text{Nb}_{0.2}\text{Ta}_{0.2}\text{Ti}_{0.2}\text{W}_{0.2})\text{Si}_2\) possesses a hexagonal C\(4\)0 crystal structure with ABC stacking sequence and a space group of P\(6\)\(_2\)22. This discovery expands the known families of high-entropy materials from metals, oxides, borides, carbides, and nitrides to a silicide, for the first time to our knowledge, as well as demonstrating that a new, non-cubic, crystal structure (with lower symmetry) can be made into high-entropy phase. This \((\text{Mo}_{0.2}\text{Nb}_{0.2}\text{Ta}_{0.2}\text{Ti}_{0.2}\text{W}_{0.2})\text{Si}_2\) exhibits high nanohardness of 16.7 ± 1.9 GPa and Vickers hardness of 11.6 ± 0.5 GPa. Moreover, it has a low thermal conductivity of 6.9 ± 1.1 W m\(^{-1}\) K\(^{-1}\), which is approximately one order of magnitude lower than that of the widely-used tetragonal MoSi\(_2\) and 1/3 of those reported values for the hexagonal NbSi\(_2\) and TaSi\(_2\) with the same crystal structure.

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1. **Introduction**

Research on high-entropy alloys (HEAs), also known as multiple principal element alloys (MPEAs) or complex concentrated alloys (CCAs), has attracted considerable interest in the last ~15 years due to their unique properties and large compositional space for engineering [1–8]. A majority of the metallic HEAs adopt the simple face-centered cubic (FCC) or body-centered cubic (BCC) crystal structures, and a few hexagonal close packed (HCP) HEAs have been made [1–8].

Only in the last ~3.5 years have the ceramic counterparts to the metallic HEAs, or “high-entropy ceramics,” been successfully fabricated in bulk forms. In 2015, Rost et al. reported an entropy-stabilized oxide, \((\text{Mg}_{0.2}\text{Ni}_{0.2}\text{Co}_{0.2}\text{Cu}_{0.2}\text{Zn}_{0.2})\text{O}\), of a rocksalt structure, yet with a rather high symmetry (P\(\text{6}_3\)\(\text{mmm}\)) [9]. In 2016, high-entropy metal diborides, e.g., \((\text{Hf}_{0.2}\text{Zr}_{0.2}\text{Ta}_{0.2}\text{Nb}_{0.2}\text{Ti}_{0.2})\text{B}_2\), were reported as a new class of ultra-high temperature ceramics (UHTCs) and the first high-entropy borides (as well as the first non-oxide high-entropy ceramics made in the bulk form) [10]. Subsequently, the research on high-entropy ceramics has made rapid advances and attracted increasing attention. First, the high-entropy (entropy-stabilized) rocksalt oxides have been studied extensively due to their great potential as functional materials with low thermal conductivities [11–13] and colossal dielectric constants [14], as well as their potential applications in lithium-ion batteries [15,16]. Second, high-entropy metal diborides have also been studied by many groups as a new class of promising structural ceramics with increased hardness [17–19]; this line of work has further stimulated the subsequent development of high-entropy metal carbides as another class of UHTCs with increased hardness by various groups worldwide [20–29]. Third, several other classes of high-entropy ceramics have also been reported, including perovskite [30–32], spinel [33], defective fluoride-structured [34,35], and rare earth [32,36] oxides, as well as high-entropy nitrides [37,38]. It is worth noting that the high-entropy oxides [30–32,34–36], carbides [20–29], and nitrides [37,38] discovered to date all have cubic crystal structures with high symmetries. The only exception is the high-entropy metal diborides, which have a hexagonal (AlB\(_2\)) crystal structure, yet with a rather high symmetry (P\(\text{6}_3\)\(\text{mmm}\)) [9].

As an increasing number of high-entropy oxides [30–32,34–36], borides [10,17–19], carbides [20–29], and nitrides [37,38] have been discovered, this study first reports, to the best of

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our knowledge, the synthesis and characterization of a high-entropy silicide: \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\). Moreover, this \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) possesses a CrSi2-type hexagonal C40 structure with the ABC stacking sequence (Fig. 1); it represents a more complex crystal structure (with a lower P6_322 symmetry) in comparison with those reported in prior studies, thereby extending the state of the art for the discovery of new high-entropy materials.

In general, refractory silicides, particularly MoSi2, are of great interest for high-temperature applications [39–45]. In this study, we have also examined the properties of this new high-entropy \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) showing high hardness (16.7 ± 1.9 GPa nanohardness and 11.6 ± 0.5 GPa Vickers hardness) and much reduced thermal conductivity (6.9 ± 1.1 W m⁻¹ K⁻¹).

2. Experimental procedure

Powders of MoSi2, NbSi2, TaSi2, TiSi2, WSi2, and ZrSi2 (99% purity, ≥45 μm; Alfa Aesar) were utilized as starting materials. The raw powders were mixed via high-energy ball milling (HEBM) utilizing a SPEX 8000D mill for 6 h in a silicon nitride jar with silicon nitride media. Heptane was used to create a slurry for grinding to prevent caking of the powders and to minimize oxidation in the milling containers. The HEBM was done in 30-min intervals, interrupted by 10-min resting pauses to avoid overheating. The powders were then densified into 20-mm diameter disks via spark plasma sintering (SPS, Thermal Technologies, CA, USA) at 1650 °C for 10 min under a uniaxial pressure of 50 MPa; then the pressure was immediately reduced to 10 MPa at a rate of 40 MPa/min at 1650 °C to minimize creep. The chamber was initially pumped down to vacuum of at least 20 mTorr and backfilled with argon for three times prior to the SPS experiments to minimize oxidation and a vacuum was maintained throughout the sintering process. The graphite die was lined with 125 μm thick graphite paper to prevent reaction of the specimen with the die.

The silicide was characterized by X-ray diffraction (XRD) utilizing a Rigaku diffractometer with Cu Kα radiation. Scanning electron microscopy (SEM) was carried out, and the corresponding energy dispersive X-ray (EDX) spectroscopy compositional maps and electron backscatter diffraction (EBSD) maps were collected. The EDX measurements were performed at an e-beam voltage of 20 kV to examine the higher energy peaks of Hf, Ta, and W for minimal convolution of the peaks.

Hardness and modulus measurements were conducted via nano-indentation on a KLA-tencor G200 Nanoindenteter (KLA-tencor, CA, USA). Nanohardness measurements were performed according to ISO 14577 under a load of 100 mN. In order to produce more statistically relevant data, the KLA-tencor Express Test software module was employed to enable very large datasets to be generated. Vicker’s hardness measurements were performed with a Vickers’ diamond indenter at 500 kgf/cm² with a hold time of 30 s. The indentations were examined for conformation with the ASTM C1327. The indentations averaged 15–20 μm in width during the testing. Thirty measurements were performed at different locations of the specimen; the mean and standard deviation are reported. The Vickers indentation test was also carried out following the ASTM standard for measuring the microhardness.

Thermal conductivities were measured using time-domain thermoreflectance [46]. A thin Al transducer (84 ± 4 nm) is thermally evaporated onto the sample. Using a Ti:Sapphire laser emitting a train of <200 fs pulses at a central wavelength of 800 nm and a repetition rate of 80 MHz, the output is divided into a pump and probe path. The pump is modulated at 8.4 MHz to heat the sample, while the probe is used to measure the resulting change in temperature as a function of delay time out to 5.5 ns after pump absorption. The pump and probe 1/e² diameters are 15 and 9 μm, respectively. The volumetric heat capacity was taken to be 2.5 ± 0.3 cm⁻³ K⁻¹ based on the rule of mixtures average of constituent heat capacities [47].

3. Results and discussion

The XRD pattern shown in Fig. 2 suggests that the \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) specimen made by SPS possesses a hexagonal structure with the space group P6_322, or the CrSi2 prototype structure. All peaks, except for one very minor peak, in the XRD pattern (Fig. 2) can be indexed to the hexagonal C40 structure with the ABC stacking sequence, as schematically illustrated in Fig. 1. SEM and EDX maps (Fig. 3) further demonstrated that this fivescattering \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) specimen indeed formed a homogeneous high-entropy solid solution. This hexagonal C40 structure was further confirmed by EBSD of a polished sample surface (Fig. 4). Lattice parameters of this \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) specimen were determined from the XRD to be: \(a = 4.711 \text{ Å}\) and \(c = 6.522 \text{ Å}\).

The formation of a hexagonal C40 crystal structure (with the ABC stacking sequence, as shown in Fig. 1) for this high-entropy \((\text{Mo}_0.2\text{Nb}_0.2\text{Ta}_0.2\text{Ti}_0.2\text{W}_0.2)\text{Si}_2\) specimen is noteworthy and interesting since only two of the five constituent silicides, NbSi2 and TaSi2 [44,48], form this hexagonal structure at high temperatures. TiSi2 possesses an orthorhombic structure (with the ABCD stacking sequence) [49]. Both MoSi2 and WSi2 normally form tetragonal structures (with the AB stacking sequence), though the hexagonal
phases were observed at lower temperatures (below 900 °C and 550 °C, respectively) in thin films [48,50].

This (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 represents a new high-entropy ceramic made, with a new, and perhaps the lowest, symmetry among all high-entropy metals and ceramics reported. To date, all except for two high-entropy metals and ceramics reported have cubic symmetries (of simple FCC and BCC [1-8], rocksalt [9,20-29,37,38], fluorite [34,35], pervoskite [30-32], and spinel [33] structures). The two other classes of non-cubic high-entropy materials reported are the metallic HCP HEAs (with the space group of P63/mmc) [8] and high-entropy metal diborides (with the space group of P6/mmm) [10]. This high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 has a lower symmetry of P6222, with a more complex ABC stacking sequence (Fig. 1).

It should be noted that a secondary TiO phase is also present, producing a minor XRD peak as indicated in Fig. 2. We assume that TiO formed because TiSi2 possesses a melting point of ~1500 °C [26,27], below our SPS temperature; thus, it is likely that TiSi2 promoted the formation a (transient) liquid phase that assisted sintering but captured surface oxides. TiSi2 has been utilized for liquid assisted sintering of diborides in a similar manner [51,52]. The secondary oxide phases seen in the SEM image (the dark phase in the first panel of SEM image in Fig. 3) are likely SiO2-based glass, which did not show up in XRD (since the amount of TiO identified by XRD, as shown in Fig. 2, is small). ImageJ analysis of the SEM image was performed to estimate the high-entropy silicide phase to

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Fig. 2. XRD pattern of the (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 specimen. Except one minor peak from a secondary hexagonal TiO phase (labeled by the red solid square), all other XRD peaks are indexed to a hexagonal C40 structure (or the CrSi2 prototype structure with the P6222 space group and the D6 point group) with the lattice parameters a = 4.711 Å and c = 6.522 Å.

Fig. 3. SEM micrograph and the corresponding EDX elemental maps of the (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 specimen.
be approximately 89 vol%.

EBSD was utilized to measure the grain size and examine the texture of the sintered (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 specimen. An average grain size of 5.4 ± 3.3 μm was found from a measurement of over 5000 grains. No significant texturing was evident in the sample. Two EBSD maps at low and high magnifications, an inverse pole figure, and the measured grain size distribution are shown in Fig. 4.

Nanoindentation hardness measurements of this (hexagonal) high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 following the ISO 14577 standard using a load of 100 mN produced a value of 16.7 ± 1.9 GPa with a large number of indents. It also measured an elastic modulus of 421 ± 19 GPa, in agreement with the measurements taken by Nakamura et al. for MoSi2 and WSi2 [53]. Moreover, we have conducted Vickers indentation test and measured a microhardness value of 11.6 ± 0.5 GPa from our high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2Zr0.2)Si2 specimen. These measured hardness values are comparable to those reported for MoSi2 in literature, with Newman et al. reporting up to 17.5 ± 2.0 GPa in nanoindentation and Vickers hardness in other prior studies varying from approximately 9 to 14 GPa [53–57]. The microhardness value of our high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 specimen is compared with five individual constituent metal disilicides in Table 1. Notably, the Vickers hardness of this high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 specimen is higher than the average of the microhardness values of the five individual metal disilicides reported in the literature (which was calculated to be 9.32 GPa by taking a median value for MoSi2).

A significantly reduced thermal conductivity was measured for this (hexagonal) high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2, in comparison with other metal disilicides [58,59]. Fitting a multilayer heat diffusion model to experimental ratio data [60], the best-fit thermal conductivity was determined to be 6.9 ± 1.1 W m⁻¹ K⁻¹.

In comparison, the thermal conductivity of the (tetragonal) MoSi2 has been measured to be 65 W m⁻¹ K⁻¹ [58]. The thermal conductivities of (hexagonal) NbSi2, (hexagonal) TaSi2, (orthorhombic) TiSi2, and (tetragonal) WSi2, respectively, were measured by Neshpor [59] to be 19.1 W m⁻¹ K⁻¹, 21.9 W m⁻¹ K⁻¹, 45.9 W m⁻¹ K⁻¹, 46.6 W m⁻¹ K⁻¹, respectively; these reported values from literature are listed in Table 1 to be compared with our measured thermal conductivity of the high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2. While it is possible that the presence of oxide contamination and porosity reduces the thermal conductivity of our specimen, the measured value of 6.9 ± 1.1 W m⁻¹ K⁻¹ is significantly lower than reported values of any of the five constituent disilicides. Noting that NbSi2 and TaSi2, which have the same hexagonal crystal and lowest thermal conductivities of 19.1 W m⁻¹ K⁻¹ and 21.9 W m⁻¹ K⁻¹, respectively [59], among the five individual disilicides, are perhaps the best benchmarks for comparison. Still, the measured thermal conductivity of this high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 is substantially lower (−1/3), presumably due to the high phonon scattering from the five different cations with different masses and a highly distorted lattice. A prior modeling study has demonstrated that >10X reduction in thermal conductivity can be achieved in high-entropy ceramics [13], and similar levels of thermal conductivity reduction was indeed observed in entropy-stabilized oxides [12].

We also attempted to fabricate a (Mo0.2Nb0.2Ta0.2W0.2Zr0.2)Si2 specimen via the same procedure, but it did not form a single high-entropy phase. The measured XRD pattern and EDX elemental maps of this (Mo0.2Nb0.2Ta0.2W0.2Zr0.2)Si2 specimen are shown in Fig. 5. While a primary hexagonal C40 phase did form, additional Ta–Zr–Si and Nb–Zr–Si rich secondary phases were observed.

4. Conclusions
A high-entropy metal disilicide, (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2, was successfully synthesized. It possesses a hexagonal structure

![Fig. 4. EBSD map of >1 mm² area of the high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 surface, showing a rather uniform microstructure. No significant texture was observed. The measured grain size distribution is given, which fits a lognormal curve. The inset on the right-bottom corner is an additional EBSD map taken at a higher magnification.](image-url)
with a space group of P6222, representing a new high-entropy material family (a high-entropy silicide) and a new non-cubic high-entropy crystal structure made. Characterization by XRD, EDX, and EBSD confirm the presence of a single high-entropy solid-solution phase, albeit some oxide contaminations.

This high-entropy (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 exhibits high nanohardness of 16.7 ± 1.9 GPa and Vickers hardness of 11.6 ± 0.5 GPa. The measured thermal conductivity of (Mo0.2Nb0.2Ta0.2Ti0.2W0.2)Si2 is 6.9 ± 1.1 W m⁻¹ K⁻¹, which is approximately one order of magnitude lower than that of the widely-used tetragonal MoSi2 [58] and ~1/3 of those reported for the hexagonal NbSi2 and TaSi2 with the same crystal structure [59]. The significant reduction in the thermal conductivity can be explained from the high phonon scattering in the high-entropy ceramic.

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