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# Experimental and theoretical evidence of charge transfer in multi-component alloys – how chemical interactions reduce atomic size mismatch†

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*Ab initio* simulations of a multi-component alloy using density functional theory (DFT) were combined with experiments on thin films of the same material using X-ray photoelectron spectroscopy (XPS) to study the connection between the electronic and atomic structures of multi-component alloys. The DFT simulations were performed on an equimolar HfNbTiVZr multi-component alloy. Structure and charge transfer were evaluated using relaxed, non-relaxed, as well as elemental reference structures. The use of a fixed sphere size model allowed quantification of charge transfer, and separation into different contributions. The charge transfer was generally found to follow electronegativity trends and results in a reduced size mismatch between the elements, and thus causes a considerable reduction of the lattice distortions compared to a traditional assumption based on tabulated atomic radii. A calculation of the average deviation from the average radius (*i.e.* the so-called  $\delta$ -parameter) based on the atomic Voronoi volumes gave a reduction of  $\delta$  from *ca.* 6% (using the volumes in elemental reference phases) to *ca.* 2% (using the volumes in the relaxed multi-component alloy phase). The reliability of the theoretical results was confirmed by XPS measurements of a Hf<sub>22</sub>Nb<sub>19</sub>Ti<sub>18</sub>V<sub>19</sub>Zr<sub>21</sub> thin film deposited by sputter deposition. The experimentally observed core level binding energy shifts (CLS), as well as peak broadening due to a range of chemical surroundings, for each element showed good agreement with the calculated DFT values. The single solid solution phase of the sample was confirmed by X-ray diffraction (XRD) and transmission electron microscopy (TEM) including energy dispersive spectroscopy (EDS) with nm-resolution. These observations show that the HfNbTiVZr solid solution phase is non-ideal, and that chemical bonding plays an important part in the structure formation, and presumably also in the properties. Our conclusions should be transferable to other multi-component alloy systems, as well as some other multi-component material systems, and open up interesting possibilities for the design of material properties *via* the electronic structure and controlled charge transfer between selected metallic elements in the materials.

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## Introduction

Multi-component materials, and in particular alloys, are a group of materials that have generated a huge amount of interest over the past 15 years. They are generally understood to be materials that have at least five major components. The interest was sparked by the concept of high entropy alloys (HEAs), which was introduced in 2004 in two separate papers by Cantor *et al.*<sup>1</sup> and Yeh *et al.*<sup>2</sup> Cantor and co-workers studied cast samples of equiatomic alloys, and identified a single-phase material with a close-packed cubic (ccp) structure; Yeh and co-workers published a study on several arc-melted and annealed multi-component alloys, where they observed single-phase ccp

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or bcc (body-centred cubic) alloys. The formation of a single-phase alloy was explained by a stabilising effect by the entropy of mixing and the concept of high entropy alloys was introduced as a design route in metallurgy. The term high entropy alloys (and materials) is under some debate, concerning definitions, as well as the mechanisms behind the formation of solid solution phases where both entropy and kinetics are important. These points are well summarised in the review by Miracle and Senkov.<sup>3</sup> In the following, the more general term multi-component alloys will be used for alloys with five or more principal components.

One reason for the continued interest in multi-component materials is that many of them have attractive properties such as excellent high-temperature strength, high oxidation resistance, good corrosion properties, high radiation resistance *etc.*, as well as new possibilities to combine functional properties; and that they offer a seemingly endless compositional space for materials design. There are several recent and substantial reviews published on the general concepts of high-entropy materials, and their properties, see *e.g.* ref. 3–7.

Several empirical design rules have been formulated to predict the formation of multi-component solid solution phases. One factor which has been shown to be important is the differences in atom radii between the elements in the multi-component phase. Too large size differences cannot be accommodated in a single lattice. The size difference is commonly quantified by the average deviation from the average atomic radius of the included metals, denoted as the  $\delta$ -parameter and calculated as:

$$\delta = \sqrt{\sum_{i=1}^N c_i \left(1 - \frac{r_i}{\bar{r}}\right)^2}. \quad (1)$$

where  $c_i$  and  $r_i$  are the concentration and radius of element  $i$ , and  $\bar{r}$  is the concentration weighted average radius of all included elements. Empirically, it has been found that  $\delta$  needs to be smaller than *ca.* 6.4% for a solid solution phase to form.<sup>8</sup> Multi-component alloys with a higher  $\delta$ -value tend to be amorphous. This rule is clearly related to both the traditional Hume-Rothery's rules for substitutional solid solutions in alloys,<sup>9</sup> and the Inoue rules for the formation of metallic glasses.<sup>10</sup> For the well-known so-called Cantor alloy the  $\delta$ -value is 3.42%, while the here studied HfNbTiVZr with equimolar metal ratios yields a  $\delta$ -value of 5.72%,<sup>‡</sup> indicating a large mismatch of metallic radii.

In the literature of multi-component alloys, there are many references to a local lattice distortion, caused by differences in atomic radii of the included elements, and its suggested effect on mechanical and other properties. The value of the  $\delta$ -parameter is commonly also used as a measure of the assumed local lattice distortion. Direct evidence of a local lattice distortion is, however, sparse, and it is technically difficult to attain, see *e.g.* the recent review by Owen.<sup>11</sup> Actually, applying a hard sphere model is an over-simplification of the structure of alloys that ignores the chemistry: also in metallic alloys there is bonding and charge transfer between the elements. Nevertheless, there are local distortions where atoms are not located at the ideal lattice points

of the underlying crystal structure (*i.e.* displacive disorder). Such local lattice distortions are clearly observed in *ab initio* simulations when structural relaxation is allowed. For example, it has recently been demonstrated that the distortions could be particularly large in bcc multi-component alloys and contribute to their stabilisation.<sup>12</sup> Also, in a recent series of publications, Tong *et al.* investigated the local lattice distortions of multi-component alloys using experimental scattering methods, yielding pair distribution functions (PDF), and *ab initio* materials simulations.<sup>13–15</sup> Their two main results are that there is a local lattice distortion, but it is significantly smaller than predicted from a hard sphere model; and that charge transfer between the atoms is responsible for reducing the size differences between the different elements and thus yielding a lattice distortion that is smaller than predicted from a hard sphere model. It is also noted that among the studied multi-component alloys, alloys containing Hf and Zr have larger distortions than the other alloys. Tong *et al.* however do not present any direct experimental evidence of the suggested charge transfer; their conclusions are based on the *ab initio* simulations.

An experimental method which is commonly used to evaluate chemical bonding and charge transfer in solid materials is X-ray photoelectron spectroscopy (XPS). The exact core level binding energy will be affected by the chemical surrounding of the photo-ionised atom, and thus the charge transfer between neighbouring atoms. However, this dependency is not simple and although a rule of thumb based on electronegativity can provide insights in many cases a more thorough approach is needed to attain reliable data. This must include simulation of the electronic state of the material, both in the initial and final state (*i.e.* before and after photo ionisation). Hence, *ab initio* simulations are required to predict XPS core level shifts.<sup>16</sup>

Interestingly, there is not much literature on multi-component alloys reporting XPS core levels. In a recent publication Xu *et al.* studied XPS core level shifts systematically for multi-component alloy nanoparticles based on group 6 to 11 transition metals.<sup>17</sup> The main conclusion was that for the studied alloys there were core level shifts in the multicomponent alloys, compared to elemental literature references, thus indicating a charge transfer. The observed shifts were found to be consistent with the electronegativity rule of thumb. These conclusions were also supported by *ab initio* materials modelling of binary alloys, and the results were discussed in the view of fundamental understanding of bonding in the multi-component alloys, and possibilities to design the electronic structure. Some interesting data have also been published in studies focussing on the reactive sputter deposition from metallic targets in a nitrogen containing process gas to attain multi-component nitride coatings. In this literature XPS core level binding energies of multi-component alloys that differ from elemental reference positions have been reported.<sup>18,19</sup> There is thus some experimental evidence in the literature for a charge transfer between different elements in multi-component alloys. This has however not been connected to the structure of the material, and in particular not to the concept of local lattice distortion.

The present study has investigated the matter of charge transfer, connected to the concept of local lattice distortion and

<sup>‡</sup> Using the metallic radii for 8-coordination, see Table 1.



XPS core level shifts in more detail, employing a combination of theory and experiment: detailed DFT simulations of relaxed and non-relaxed multi-component alloys and high-resolution XPS to experimentally validate the suggested charge transfer and its effect on the atomic structure. The HfNbTiVZr system was chosen as a demonstrator system, as it is a known entropy-stabilised phase,<sup>20</sup> which can be produced as thin films.<sup>21</sup> Furthermore, the presence of Zr and Hf suggests that a (comparably) large distortion and charge transfer is expected,<sup>15</sup> making the system suitable for further studies.

## Methods

### Theoretical methods

**Density functional theory (DFT) calculations.** The DFT calculations were performed using the projector augmented wave method<sup>22</sup> as implemented in the Vienna Ab initio Simulation Package (VASP)<sup>23,24</sup> with the Perdew–Burke–Ernzerhof generalized gradient approximation<sup>25</sup> to model the exchange correlation effects. All calculations are performed using a 420 eV kinetic-energy cut-off, and with a convergence criterion on forces of 0.01 eV Å<sup>-1</sup> for structural relaxations.

The HfNbTiVZr multi-component alloy was modelled using a bcc underlying crystal lattice, equimolar concentrations, and a 3 × 4 × 5 supercell with 120 atoms, *i.e.* 24 atoms of each element. The supercell was created using the special quasirandom structure method (SQS).<sup>26</sup> The supercell was first optimised with respect to volume using a fixed atomic position. An equilibrium structure was obtained by relaxing the supercell, allowing atomic displacements and supercell volume change. Both relaxed and non-relaxed supercells were used to determine the displacements of the individual atoms from their ideal positions in the bcc structure. This was quantified by determining the pair distribution function (PDF) which gives the observed atomic distances; and the angular distribution function (ADF) which provides the distribution of observed angles ( $\theta$ ) between three atoms. The calculations were performed using the same system as in a previous study on HfNbTiVZr, where theoretical simulations were confirmed by experimental scattering data from bulk samples.<sup>27</sup> In addition, the pure elements were modelled for comparison, simulating both room temperature stable phases (bcc for Nb and V, and hcp for Hf, Ti and Zr), as well as bcc structure for all the metals.

**Estimation of charge transfer.** The charge on a given ion is not uniquely defined in *ab initio* simulations and depends on the choice of model to partition the electron density between different atomic nuclei. In the simplest approach, one can take the charge enclosed in the volume of a sphere defined by the atomic radius, henceforth denoted as the fixed sphere model. In addition, two more advanced methods were used: in the framework of the atoms-in-molecules theory proposed by Bader<sup>28</sup> atoms are defined through a partitioning of real space using the gradient of the electron density to define where one atom basin ends and the next begins. The Voronoi tessellation method offers a geometrical approach to partition space, where the atomic volumes take the form of irregular polyhedra, constructed

with the planes, which are perpendicular bisectors between an atom and its neighbours. The Bader charge integration was performed using the Henkelman Bader Charge Analysis code,<sup>29,30</sup> and the Voronoi volumes were calculated with the Voro++ library.<sup>31</sup>

**Core level shifts by *ab initio* calculations.** To allow direct comparison between experiments and *ab initio* modelling, the XPS process was simulated using the so-called  $Z + 1$  approximation.<sup>32</sup> The core ionized atom  $Z^*$  is replaced by the next element in the periodic table:  $Z + 1$ . In this way, the core-electron excitations and the effect of a core-level hole screened by an extra valence electron are approximated with an extra proton and an extra electron. The photoelectron binding energy shifts (relative elemental references) can thus be calculated for all atoms in the modelled multi-component alloy supercell, as described, for instance in ref. 33.

### Experimental methods

A multi-component thin film sample was synthesised by non-reactive DC-magnetron sputtering using an ultra-high vacuum system, base pressure  $< 1 \times 10^{-7}$  Pa ( $< 1 \times 10^{-9}$  mbar). The system was equipped with four 2" magnetrons in a confocal sputter-down configuration. Elemental targets of Hf, Ti, and V were used on three of the magnetrons. The fourth magnetron was equipped with a segmented target, composed of one-half Zr and one-half Nb. An Ar<sup>+</sup> plasma was ignited at 0.4 Pa ( $4 \times 10^{-3}$  mbar), using a 42 sccm Ar gas flow. Prior to the deposition, the single-crystal Si(001) substrates were cleaned in an ultrasonic bath, first in acetone followed by ethanol, each time for 5 minutes, and then dried in a N<sub>2</sub>(g) flow. The substrates were pre-heated to 275 °C for at least 60 min before the deposition, to minimise the temperature gradient. Deposition was carried out at this substrate temperature, and a dc bias of –50 V was applied.

The deposited thin film was analysed with a number of techniques to ensure the presence of a single solid solution phase. The crystal structure was studied by X-ray diffraction (XRD) using Cu K $\alpha$  radiation. A Bruker D8 advance diffractometer was used for symmetrical ( $\theta/2\theta$ ) scans using Bragg–Brentano geometry and a Johansson monochromator to attain pure Cu K $\alpha_1$  radiation. A Philips MRD X'Pert was used for grazing incidence (GI) scans and determination of in- and out-of-plane lattice parameters. GI scans were performed with an incidence angle of 2°, using a Göbel mirror as primary optics and parallel plate collimator of 0.27° divergence as secondary optics. For determination of the lattice parameters in different spatial directions point focus with capillary lens optics on the primary side; and a parallel plate collimator of 0.18° divergence with a flat graphite monochromator on the secondary side were used.

Transmission electron cross-section samples were prepared using the *in situ* lift out technique in a FEI Strata DB235 focused ion beam (FIB) instrument. The transmission electron microscopy (TEM) analysis was conducted on a FEI Titan Themis 200 kV, equipped with a C<sub>s</sub> probe corrector and SuperX Energy dispersive spectroscopy (EDS) system. Analysis was mainly performed in scanning mode, using a high angle annular dark field (HAADF) detector. EDS analysis was performed to attain spatially resolved compositional data, and evaluated using the metal K-lines, and Bruker Esprit software.



Bulk metal reference samples were attained from the back-side of used elemental sputter targets. These all had a claimed purity of 99.9% or higher, and were supplied by Plasmateials (Hf and V) and Kurt J. Lesker (Zr, Nb and Ti).

The chemical environment of both elemental references and the multi-component thin film was studied with X-ray photo-electron spectroscopy (XPS) using a PHI Quantera II XPS scanning microprobe with Al  $K\alpha$  radiation ( $h\nu = 1486.7$  eV) and an electron take off angle of  $45^\circ$ . The spectrometer was calibrated against reference samples of Ag, Au and Cu according to ISO 15472.<sup>34</sup> For the present samples the Fermi levels were also measured, and all align to zero within an uncertainty of  $\pm 0.05$  eV. Compositional analysis was performed using sputter depth-profiles with 1 keV Ar<sup>+</sup> etching, and custom sensitivity factors calibrated using data from Rutherford backscattering (RBS) on similar samples.

The core levels were investigated by high-resolution XPS which was obtained after sputter etching with 4 keV Ar<sup>+</sup> for 6 minutes to remove the surface oxide layer. For these analyses an X-ray spot size with a diameter of 50  $\mu\text{m}$  and a low pass energy (6.5 eV) was used to attain the highest possible spectral resolution. This resulted in a full width half maximum (fwhm) for the Hf 4f<sub>7/2</sub> of 0.29 eV, but also very long measurement times: about 2 h for each elemental reference, and 9 h for the multi-component alloy sample. These long measurement times, combined with a vacuum level of  $2 \times 10^{-6}$  Pa ( $2 \times 10^{-8}$  mbar), unfortunately resulted in some *in vacuo* oxidation of the samples during analysis. In all cases, however, the contribution from oxide could be clearly separated from the pure metallic contribution by peak fitting.

XPS core level spectra were analysed by means of peak fitting to attain peak positions and widths of the metallic contributions of elemental references and the alloy sample. Igor Pro 6.37 software was used, and Shirley-type backgrounds were implemented. Doniach-Šunjić type asymmetric peak functions were used for metallic contributions, and Voigt type symmetric peak functions were used for the oxide contributions. For the metallic contributions, the Lorentzian (connected to the life time, and assumed constant) was set from ref. 35, and the Gaussian was allowed to vary. The Gaussian will be connected to both the instrumental broadening (also assumed constant), and any variation in the local chemical surrounding which can be expected in a multi-component alloy. By comparing the total fwhm of the fitted peak for reference metal samples and the multi-component alloy sample, the additional alloy broadening of the core levels was assessed.

## Results and discussion

### Simulation results

DFT simulations were performed on the alloy, using the equimolar Hf<sub>20</sub>Nb<sub>20</sub>Ti<sub>20</sub>V<sub>20</sub>Zr<sub>20</sub> composition. The non-relaxed multi-component alloy with an (undistorted) bcc structure has a lattice constant of 3.40  $\text{\AA}$ , whereas the supercell relaxation yielded a slightly smaller lattice constant of 3.36  $\text{\AA}$ . Previous theoretical calculations on the ideal (undistorted) alloy have reported a 3.40  $\text{\AA}$  lattice constant,<sup>36</sup> in excellent agreement with our calculations, whereas previous

experimental results have reported values in the range of 3.37–3.46  $\text{\AA}$ .<sup>20,21,36–38</sup> The values are plotted in Fig. 1 together with data for the pure elements.<sup>‡</sup> Note that Vegard's law for a disordered bcc structure yields a lattice parameter of 3.33  $\text{\AA}$  for the equimolar alloy. It is noted that the present value and literature values are slightly larger than what Vegard's law predicts. This difference is usually explained with a non-ideal solid solution and indicates that a simple hard-sphere model is insufficient to explain the structure.

The results shown in Fig. 2 give further insight into the structural changes connected to the relaxation of the DFT supercell: (a) shows the pair distribution function (PDF); and (b) the angular distribution function for an undistorted perfect bcc structure (black) and the DFT relaxed multi-component alloy (red). Note that the reliability of the theoretical PDF for the relaxed alloy has been confirmed in explicit comparison with experiment.<sup>27</sup> Inspecting the PDFs in Fig. 2, it is possible to observe that after relaxation the alloy still has a bcc structure, yet the distortions caused by different local chemical environments produce a range of neighbour distances. The first and second nearest neighbour distances are no longer easily distinguishable, and appear as a shoulder in the first peak. Likewise, the angular distribution also presents a range of values, but also confirms that the bcc structure is preserved.

To further investigate the structural change, the distribution of the displacements from the ideal bcc positions for each element type in the multi-component alloy is plotted in Fig. 3. As can be seen, all atoms are displaced in the relaxation, and the average atomic displacement ranges from 0.25 to 0.31  $\text{\AA}$ , with standard deviations in the range of 0.09 to 0.15  $\text{\AA}$ . These values are in good agreement with what Samolyuk *et al.* attained for the similar HfNbTiZr alloy.<sup>12</sup> This means that the average displacements of all the different elements are within the standard deviations, and therefore the magnitudes of the displacements seem more dependent on the local chemical environment, than on a particular element. The large spread in displacement also suggests that this

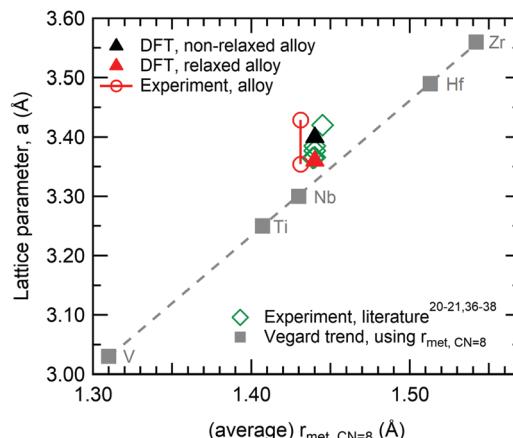


Fig. 1 Illustration of the different lattice parameters obtained for the HfNbTiVZr multi-component alloy, as a function of (average) metallic radii, using the metallic radii for 8-coordination, see Table 1. The Vegard trend (square marker and dashed line) shows calculated lattice parameters, based on the ideal bcc structure; other series show observed lattice parameters.



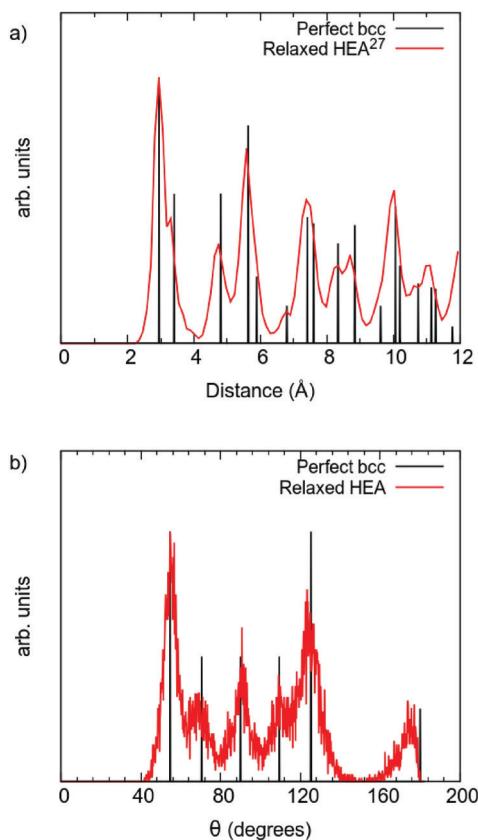


Fig. 2 Comparison between perfect bcc structure multi-component alloy (black), and DFT relaxed multi-component alloy (red): (a) pair distribution function, and (b) angular distribution function.

dependence goes beyond the first nearest neighbour shell. Still, it may be worth noting that Nb and Hf are the elements with the smallest average displacement from ideal positions, and V is the element with the largest average displacement. The latter may be rationalised by V being the smallest in size. For all elements, except Ti, the distribution is 'bottom heavy' with the most frequent displacement smaller than the average displacement and generally decreasing frequencies for larger displacements. Due to the limited statistical base (24 atoms of each element), it is however difficult to draw any conclusions based on this.

This displacive disorder of the atoms clearly shows that there is a local lattice distortion. It is however important to note that this is not simply caused by atoms of different sizes, leading to stretched bonds and strain. DFT simulations do not apply a hard sphere model, but instead determine the total electron density (and the position of the nuclei). This means that also the chemical interaction of the elements is incorporated, and there is more knowledge to be gained from studying the electron density and atomic positions carefully.

To visualise how the electronic structure is affected by the presence of different neighbouring elements (*i.e.* local chemical environments) and distortions, electronic density map slices are presented in Fig. 4, for both the HfNbTiVZr multi-component alloy (a) and a bcc metal (Nb) as a reference (b). From these maps one can clearly see that there is a large difference between

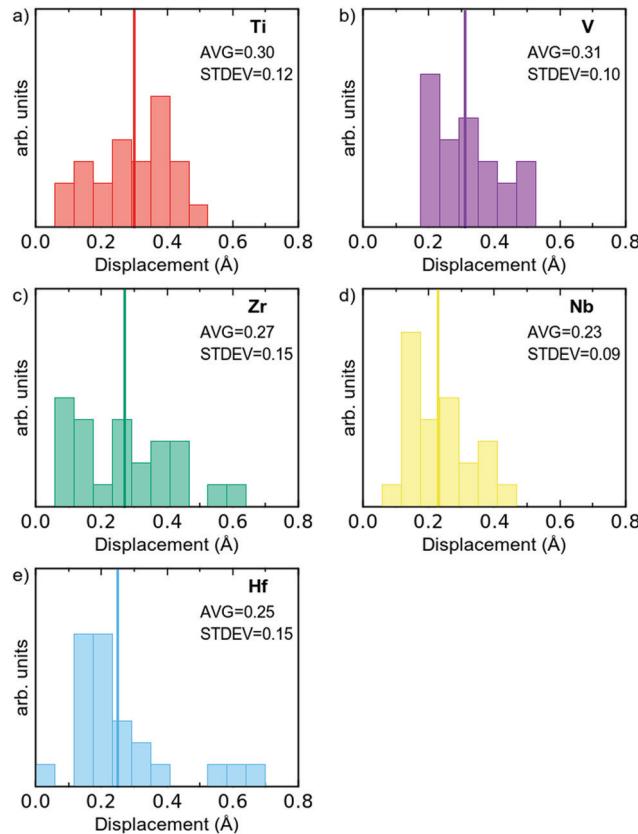


Fig. 3 Distribution of the displacement from ideal bcc positions, per element type for all 120 atoms in the simulated supercell. Average displacement indicated by a solid line.

the electronic structure of the multi-component alloy and an element with the same structure. The electronic density is delocalised to a much larger degree in the case of the alloy, indicating charge transfer between the elements. Also the displacive disorder is illustrated in Fig. 4, as the balls representing the nuclei are shifted in, and perpendicular to, the plane of the slice. The latter is seen by the different sizes of the balls.

The above noted delocalisation makes it hard to define clear borders between the atoms in the alloy, in contrast to the pure Nb metal where the borders are quite clear. The concept of assigning electrons (charge) to a specific atom and discussing charge transfer between atoms is thus not straightforward, as this is a grave simplification of reality. One may however approach this simplification by assigning the volume over which to integrate the charge that belongs to a specific atom. There is not one unique way to do this, but several approaches exist, see the Experimental section above.

The more complex Voronoi and Bader approaches are attractive as they go beyond the non-physical model of hard spherical atoms. However, in the present case where each atom has a unique local environment, each atom will also have a unique volume and the larger the volume, the larger the enclosed charge. The actual charge transfer between atoms depends thus on both the chemical environment and atomic displacements. An issue with both the Bader and Voronoi methods, which occurs in the studied system,

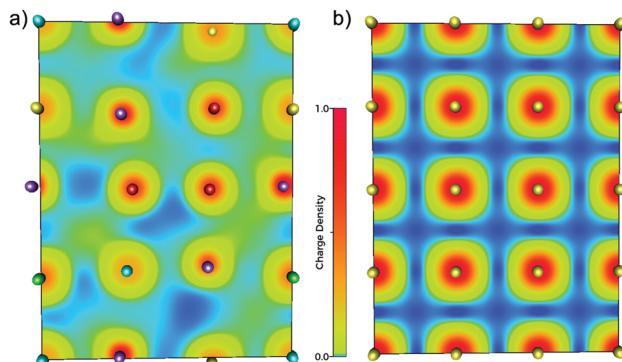


Fig. 4 Electronic density map slices from DFT simulations, for a (100) plane for the  $\text{Hf}_{20}\text{Nb}_{20}\text{Ti}_{20}\text{V}_{20}\text{Zr}_{20}$  multi-component alloy (a), and a reference bcc metal (b). Different elements indicated by different colour of the spheres: Ti – red, V – purple, Zr – green, Nb – yellow, and Hf – light blue. The colour scale shows the normalized charge density for the respective panel.

is that it is difficult to separate between these two contributions. Additionally, charge transfer strongly depends on the reference state, the unrelaxed multi-component alloy super cell could be used as the reference state but carries the trouble of comparing the charge enclosed in different volumes.

To assess the redistribution of charge due to different local chemical environments and lattice distortions, we choose to count the number of electrons present in a fixed spherical volume around each atomic nucleus. Comparing the electrons enclosed in this volume for a given atom in the alloy and in its elemental reference gives a picture of the change in charge distribution in the same region of space due to the different chemical environments. We believe this approach is a good descriptor to assess charge transfer and core level shifts. The used “charge transfer” ( $\Delta q$ ) is defined in eqn (2), where  $q$  denotes the elemental charge enclosed in the fixed spherical volume. One must note that this is not a space filling approach and therefore not all the electrons in the system are encompassed, and the summation of charge differences does not achieve charge neutrality. We have included a comparison between the Bader and fixed sphere model in the ESI.†

$$\Delta q = q_{\text{alloy}} - q_{\text{elemental ref.}} \quad (2)$$

However, this approach is also challenging, since there is a distribution of enclosed charges over the 24 atoms of each

element in the simulation supercell, and there are several contributing factors when comparing directly with the elemental references, such as change of lattice parameter and local chemical environment that influence the enclosed charge of each atom. For the latter, a step-wise analysis can be helpful; and in a first evaluation, the distributions of charge transfer for the different atoms can be assessed with the help of averages. In Table 1 the average charge transfer for each element is analysed in steps: from the elemental reference to the relaxed alloy. The analysis was made using the fixed sphere model and the Wigner–Seitz radii provided in the VASP pseudopotentials to probe charge transfer. The charge transfer ( $\Delta q$ ) has been defined as a difference in elemental charges, and an atom gaining electrons will thus show a negative  $\Delta q$  value. As Hf, Ti and Zr have no stable low temperature bcc phases, DFT simulations were used to determine how charge would be distributed in the bcc phase, with a relaxed lattice parameters denoted  $a_0$ . Thus, the first column showing charge transfer (average  $\Delta q$ ) gives the change in enclosed charge due to the difference in structure, from hcp to bcc (elemental ref. – bcc  $a_0$ ). This only leads to minor charge transfer, 0.02 to 0.08 elemental charges.

In the second column for charge transfer (average  $\Delta q$ ), the elements are compared in the bcc structure but with the same lattice constant as in the multi-component alloy (bcc  $a_{\text{non-relax}}$  – bcc  $a_0$ ), thus probing the result of different lattice constants. This step creates the largest contributions to the charge transfer, with absolute values of 0.08 to 0.43 elemental charges. This can be rationalised in the following: Hf and Zr possess larger lattice constants than the alloy, one could imagine the atoms are squeezed to fit into a smaller structure, therefore having more electrons in the enclosed volume, while the opposite happens for the smaller Nb, Ti, and V that will have less enclosed electrons in the fixed volume.

In the third column for charge transfer (average  $\Delta q$ ), the unrelaxed alloy phase is compared with the elemental bcc phases with the lattice constant of the alloy (alloy  $a_{\text{non-relax}}$  – bcc  $a_{\text{non-relax}}$ ). This can give an estimate of the charge transfer due to the local chemical environment. From the data in Table 1 it is clear that the transfer due to local chemical environment is in line with electronegativity, with the larger and less electronegative atoms (Hf and Zr) losing electrons to the smaller and more electronegative atoms (Ti, V, Nb), thus yielding a positive  $\Delta q$  for Hf and Zr, see Table 1.

Table 1 Atomic literature data and result of charge analysis in the DFT simulated phases. Pauling electro negativities (E.N.),<sup>39</sup> metallic radii from the literature ( $r_{\text{met,tab}}$ )<sup>39</sup> metallic radii for 8-coordination ( $r_{\text{met,CN=8}}$ ) implementing a coordination number compensation factor<sup>40</sup> for the elements not found in the bcc structure. Average charge transfer ( $\Delta q$ ) in elemental charge for the fixed sphere model, comparing different DFT simulated phases. a with different subscripts denotes the lattice parameter

E.N.	$r_{\text{met,tab}}$ (Å)	$r_{\text{met,CN=8}}$ (Å)	Element ref. – bcc $a_0$	Average $\Delta q$ (elemental charge)				Sum Alloy $a_{\text{relax}}$ – elemental ref.
				Structure bcc $a_{\text{non-relax}}$ – bcc $a_0$	Lattice const. Alloy $a_{\text{non-relax}}$ – bcc $a_{\text{non-relax}}$	Chem. env. Alloy $a_{\text{relax}}$ – Alloy $a_{\text{non-relax}}$	Atomic displac. Alloy $a_{\text{relax}}$ – Alloy $a_{\text{non-relax}}$	
Ti	1.5	1.45	1.41	-0.03	0.14	-0.06	-0.05	0.00
V	1.6	1.31	1.31	n/a	0.43	-0.16	-0.15	0.11
Zr	1.4	1.59	1.54	-0.08	-0.28	0.15	0.10	-0.11
Nb	1.6	1.43	1.43	n/a	0.08	-0.03	-0.05	-0.01
Hf	1.3	1.56	1.51	-0.02	-0.32	0.14	0.11	-0.10



In the fourth column of charge transfer (average  $\Delta q$ ) the unrelaxed and relaxed alloy phases are compared (alloy  $a_{\text{relax}}$  – alloy  $a_{\text{non-relax}}$ ), thus giving the charge transfer contribution due to the atomic displacement. This contribution is very similar to the previous one, and can be interpreted as that the relaxation allows for an increased charge transfer that is driven by the chemical environment.

The final column shows the sum of the previous columns, which thus becomes the comparison between the relaxed alloy phase and the elemental references in their room temperature phase. As can be seen, this average  $\Delta q$  does not follow any clear trend as it contains contributions from several factors.

In summary, the DFT simulations show that there is a re-arrangement of the electron density of a multi-component alloy compared to the pure elements, *i.e.* a charge transfer between the elements. These results are also consistent with the previous results of Tong *et al.*<sup>15</sup>

From the discussion above, it is also clear that the volume required for each atom is unique in the alloy, where a distribution exists, compared to the elemental references where the elements have a well-defined volume. This is illustrated in Fig. 5, where the Voronoi volumes are plotted for each element, and compared to Voronoi volumes for the elemental reference phases (dashed vertical lines). As can be seen the large Hf and Zr atoms

shrink (Fig. 5c and d), and the smaller V and Ti atoms expand (Fig. 5a and b). Nb, which has a metallic radius in-between the four other elements, exhibits a small spread of values around that of the elemental reference (Fig. 5d). This observation is also consistent with the suggestion from Tong *et al.*<sup>15</sup> that charge transfer in multi-component alloys reduces the size mismatch between the elements and thus reduces the proposed local lattice distortions.

As stated in the Introduction the  $\delta$ -value of HfNbTiVZr based on eqn (1) is 5.72% if one uses the metallic radii for 8-coordination (see Table 1). If the Voronoi volumes in Fig. 5 are equated to a spherical volume, a sort of atomic radii can be attained. If the Voronoi volumes from the elemental reference phases are used, a slightly different value of 6.52% is obtained for the HfNbTiVZr alloy. The slightly higher value comes from the space filling nature of the Voronoi approach when compared to the hard sphere approach. If, however, the average volume for each element in the multi-component alloy is used, a  $\delta$ -value of only 1.76% is attained for the alloy. One could also refrain from using the average radii, and let the sum in eqn (1) be over all 120 individual atoms in the supercell, which gives a  $\delta$ -value of 2.07%. Both these values are significantly lower than when employing the hard sphere model with tabulated metallic radii. This shows that the concept of local lattice distortion based on tabulated radii is not generally valid to estimate actual local lattice distortions, as chemical interaction between the elements reduces the size mismatch of the atoms.

The above-described conclusions are purely based on *ab initio* materials simulations using DFT. There are however experimental possibilities to probe the electronic structure of alloys, and compare them to the electronic structure from modelling. Using XPS one can probe the binding energy of different core electrons, the value of which is sensitive to the local chemical surrounding. If the electronic structure of the alloy has changed, compared to the pure elements, one would expect a change in the core level binding energies, *i.e.* a core level shift (CLS).

Based on DFT models it is possible to simulate the XPS process, and attain binding energies for core electrons.<sup>33</sup> The absolute values of these binding energies will not be directly comparable to experimental values, due to the work function, which is not considered, and other systematic errors. However, one can successfully compare differences in binding energy between different environments, to calculate the CLS. Here the room temperature stable elemental phases of the respective element were used as the reference, and the CLS of the multi-component alloy was calculated for direct comparison to experimental data. The attained CLS for all the 120 atoms in the multi-component alloy supercell are presented in Fig. 6 as histograms; and the averages and fwhm of the distribution are given in Table 2. As can be seen, practically all atoms show a shift (CLS values different from 0), consistent with the changed electronic structure. Furthermore, there is for all elements a spread in the observed CLS, which is the result of the varying chemical surroundings. Electrons from the Hf 4f and Zr 3d orbitals show the largest negative CLS, with average values of  $-0.26$  and  $-0.53$  eV, respectively. V 2p shows the largest

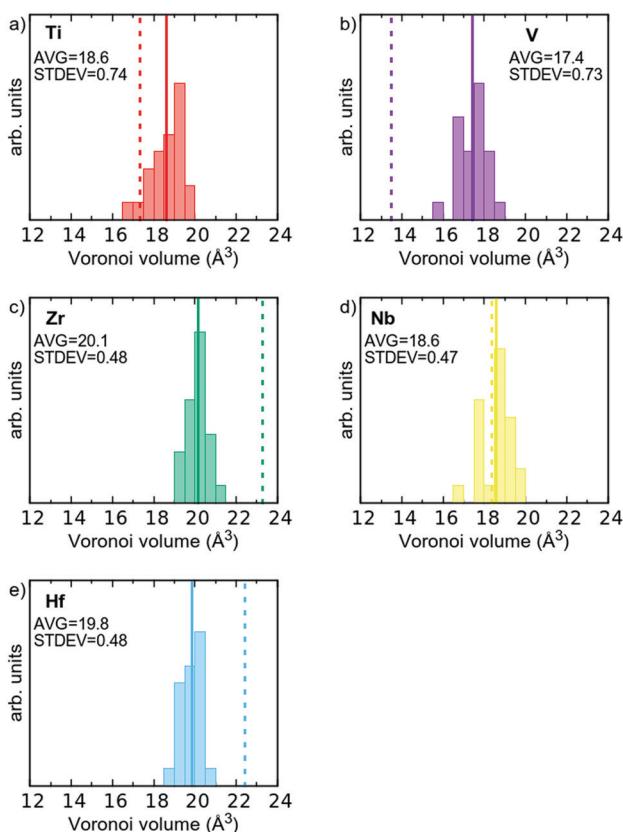


Fig. 5 Histograms of atomic volumes using the Voronoi approach in the relaxed multi-component alloy, per element type for all 120 atoms in the simulated supercell. Average volume indicated by the solid lines. Atomic volume of respective elemental reference phase given as dashed vertical lines.



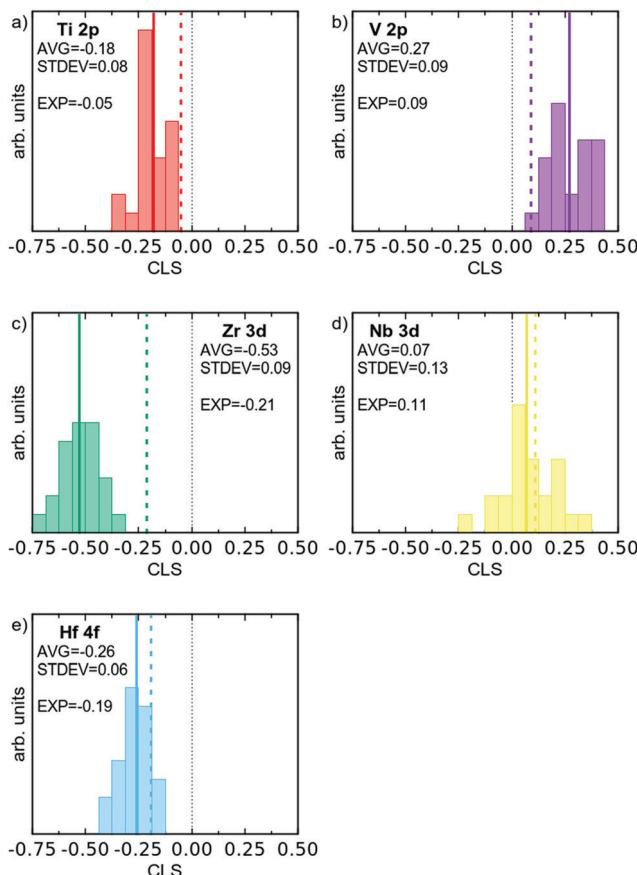


Fig. 6 Histograms of simulated XPS CLS for the 120 atoms of the DFT supercell, divided according to element. Average CLS indicated by a solid line, and experimental value by a dashed coloured line.

positive CLS with an average of 0.27 eV. Nb 3d shows a slight positive shift, with an average of 0.07 eV. The CLS of Ti 2p is in the middle of the group.

The spread of CLS values for each element can be expected to correlate to an experimentally observed alloy broadening, and the theoretical CLS data gives broadenings of 0.15 to 0.30 eV. In the theoretical values we have included results from relaxed and non-relaxed structures in order to assess the importance of structural relaxation (see below).

**Table 2** Summary of numerical values attained from XPS peak fitting of metal core levels, and from *ab initio* simulations.  $E_B$  denotes the core level binding energies and fwhm the peak broadening as full width at half maximum. CLS is the core level shift of the alloy, compared to the elemental reference. Theoretical values within (parentheses) are for the non-relaxed structure

Core level	Method	Elemental reference		Multi-component alloy		CLS (eV)	Broadening (eV)
		$E_B$ (eV)	fwhm (eV)	$E_B$ (eV)	fwhm (eV)		
Ti 2p <sub>3/2</sub>	Experiment	453.87	0.54	453.82	0.92	-0.05	0.38
	Theory	0		0.18 (0.23)	-0.18 (0.17)	0.18 (0.23)	
V 2p <sub>3/2</sub>	Experiment	512.08	0.60	512.17	0.94	0.09	0.34
	Theory	0		0.22 (0.26)	0.27 (0.78)	0.22 (0.26)	
Zr 3d <sub>5/2</sub>	Experiment	178.72	0.39	178.51	0.60	-0.21	0.21
	Theory	0		0.21 (0.39)	-0.53 (-0.54)	0.21 (0.39)	
Nb 3d <sub>5/2</sub>	Experiment	202.27	0.48	202.38	0.65	0.11	0.17
	Theory	0		0.30 (0.38)	0.07 (0.44)	0.30 (0.38)	
Hf 4f <sub>7/2</sub>	Experiment	14.15	0.29	13.96	0.43	-0.19	0.14
	Theory	0		0.15 (0.35)	-0.26 (-0.73)	0.15 (0.35)	



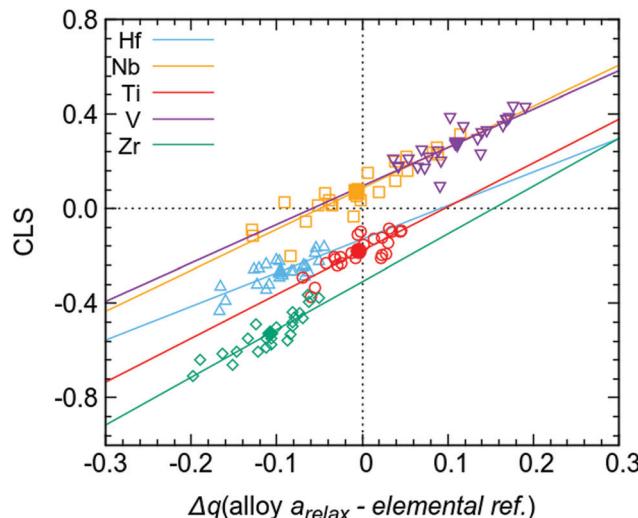


Fig. 7 Calculated core level shifts for all 120 atoms (open markers) in the supercell versus charge transfer compared to elemental references, calculated using the fixed sphere model, i.e. all individual values giving the average  $\Delta q$  (alloy  $a_{\text{relax}}$  – elemental ref.) in the last column of Table 1 (also shown here as filled markers).

to vary between 3.35 Å (in-plane) and 3.43 Å (out-of-plane) due to compressive residual stresses. This range is indicated in Fig. 1 (red circular markers), and is in agreement with the value attained by DFT for the relaxed structure (3.36 Å). Experimental values are offset in  $X$  due to the slightly different composition, and thus different average metallic radius.

The multi-component alloy sample was further studied with TEM, and a scanning transmission electron microscope (STEM) bright field micrograph is shown in Fig. 9, giving an overview of the prepared TEM lamella. A fine-grained columnar and dense microstructure is clearly observed. This is in good agreement with the strong (110) texture, observed by  $\theta/2\theta$ -XRD. In general, the sample is found to be homogeneous with no indications of

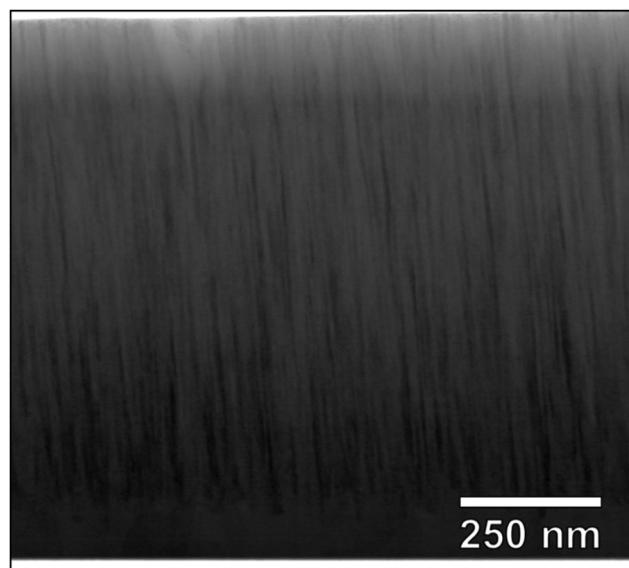


Fig. 9 STEM bright field cross section overview micrograph. The top region of the lamella is thinner than the bottom region, which leads to the observed contrast.

secondary or minority phases, the difference in brightness between the top and the bottom of the film is due to a variation in thickness of the TEM lamella. Selected area electron diffraction (not shown) gave rise to spot patterns, consistent with the highly textured bcc phase observed in XRD. To confirm the assumed random solid solution of the alloy coating, STEM EDS was performed on a representative part of the sample. Fig. 10 shows a HAADF STEM image (a) of the columnar microstructure at higher magnification, together with the corresponding EDS maps (b). As can be seen, Hf, Zr and Nb appear homogeneously and randomly distributed on the nanometre scale. However, for V and Ti the outlines of the columnar microstructure can be observed just above the noise level. This indicates a partial segregation of these two elements. The result of a line-scan across several columns is shown in Fig. 10(b), where one can see that the segregation is indeed partial with a variation of about  $\pm 5$  at% (15–22 at% for Ti, and 10–21 at% for V).<sup>§</sup> This could be connected to results reported by Pacheco *et al.*,<sup>20</sup> where Calphad calculations show that a multi-component HfNbTiVZr alloy is enriched in Ti and depleted in V in the lower temperature range as secondary phases start to appear. It is thus concluded that a five-component solid solution phase with a bcc structure exists throughout the sample. There is however in some regions a partial segregation of Ti and V to the extent of a few at%. The sample is hence relevant to study the chemical bonding in a multi-component alloy, and for direct comparison with the above presented DFT simulations.

High-resolution core level XPS spectra of the multi-component alloy (red spectra) and the corresponding reference metals (black spectra) are presented in Fig. 11(a)–(e). As can be seen, there are

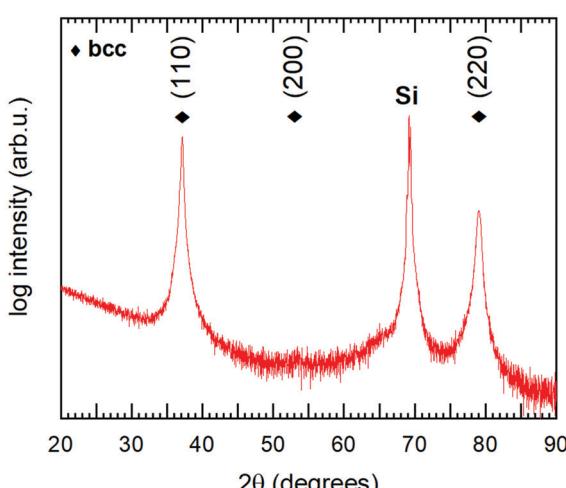


Fig. 8  $\theta/2\theta$  diffractogram of the studied  $\text{Hf}_{22}\text{Nb}_{19}\text{Ti}_{18}\text{V}_{19}\text{Zr}_{21}$  coating. Diamond markers show positions for the observed bcc phase. The peak at  $2\theta \sim 69^\circ$  originates from the Si substrate.

<sup>§</sup> N.B. these concentration values are directly from the EDS analysis, not calibrated XPS values.



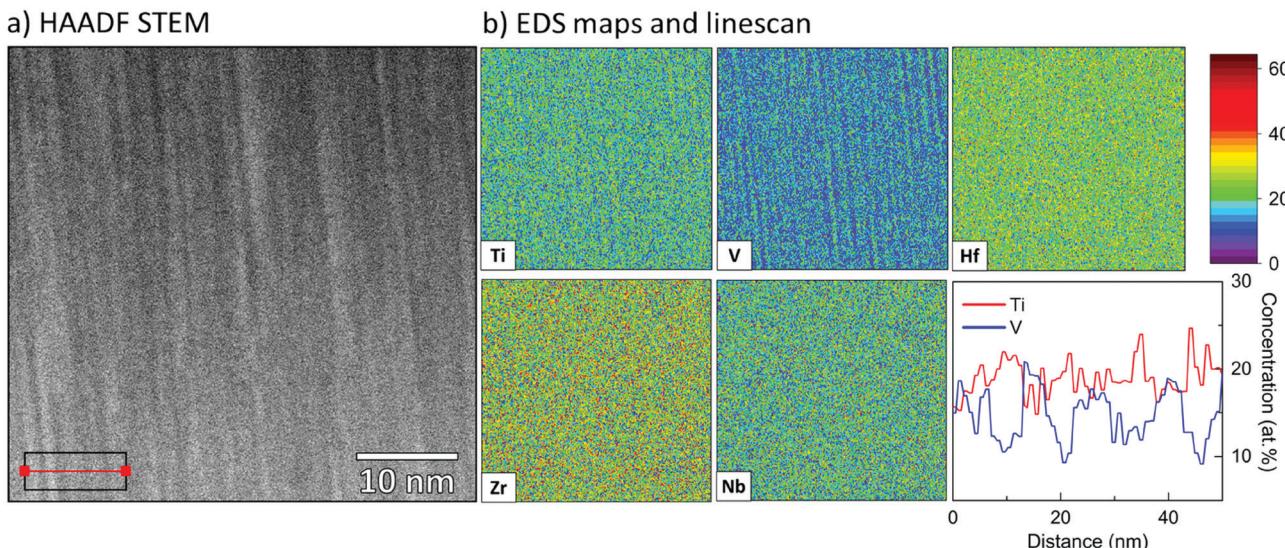


Fig. 10 HAADF image of EDS area and the corresponding maps, and an EDS line-scan over several columns for Ti and V, indicated by the black box and red line in (a). EDS analysis was performed using the metal K-lines (b).

differences between the spectra of the alloy and the elemental references, which is the main interest of this paper. Apart from the metallic contributions, there are for Nb, Zr and Hf clearly also contributions at higher binding energies from metal atoms in an oxygen environment (positions marked with dashed lines in Fig. 11). These originate from the *in vacuo* oxidation which occurred in the analysis chamber of the electron spectrometer, and the effect is larger for the alloy sample as the measurement time for this sample was about four times longer. The oxide contributions can however easily be separated from the metallic contributions by peak-fitting procedures, which have been employed. An example of the peak fitting (for Hf 4f of the multi-component alloy) is shown in Fig. 11(f), and all fits are found in the ESI.† Peak positions and widths of the main, metallic, contributions are summarised in Table 2. The lowest energy spin-orbit component ( $2p_{3/2}$ ,  $3d_{5/2}$  or  $4f_{7/2}$ ) was used for fittings determining the CLS as this component is more well-defined due to a higher intensity, smaller Lorenzian broadening and lesser overlap with oxide components. We also note small changes in the spin-orbit splitting in the alloy compared to the metal references. Such a change is indicative of a small change in chemical environment.<sup>44</sup>

Coming back to the differences between the spectra of the multi-component alloy and the elemental references, the following can be observed: a significant shift of  $-0.2$  eV for the Hf  $4f_{7/2}$  and Zr  $3d_{5/2}$  peaks of the alloy, compared with the elemental references; whereas Nb  $3d_{5/2}$  and V  $2p_{3/2}$  are shifted to  $0.1$  eV higher binding energies. The binding energy of Ti  $2p_{3/2}$  remains unchanged, within the uncertainty of the determination. These core level shifts clearly show that the electronic structure of the multi-component alloy is different from the pure elements, thus indicating charge transfer between the elements and a non-ideal solid solution, in agreement with the theoretical modelling presented above.

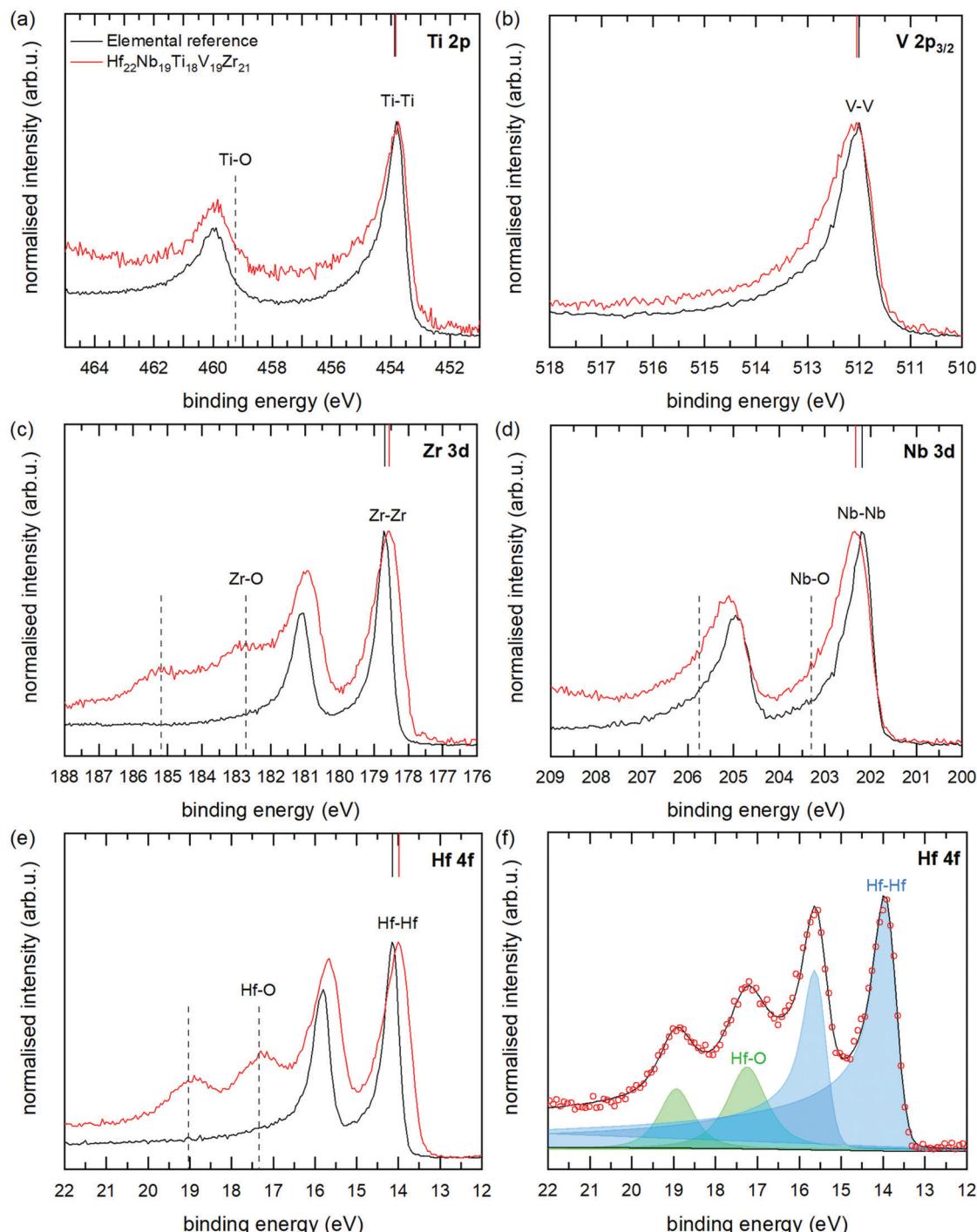
The sign and magnitude of the CLS of Ti 2p, V 2p, Nb 3d, and Hf 4f all agree well with the CLS values from theory and the relaxed alloy simulation. As can be seen from Table 2 and Fig. 6 (experimental values indicated by dashed coloured lines) V 2p and Nb 3d exhibit a positive CLS with experimental values of  $0.09$  and  $0.11$  eV, respectively; while the Ti 2p, Zr 3d and Hf 4f all exhibit a negative CLS with experimental values in the range of  $-0.05$  to  $-0.21$  eV. The largest deviation from theoretical values is observed for Zr 3d, which exhibits a smaller CLS experimentally ( $-0.21$  eV) than predicted from theory ( $-0.53$  eV), although both values put Zr in the group with distinct negative CLS.

All the core levels of the alloy also exhibit an increased broadening compared to that of the elemental references, with  $0.14$  to  $0.38$  eV larger fwhm, see Table 2. The observed broadening is in excellent agreement with the CLS variation from the relaxed alloy simulation ( $0.15$  to  $0.30$  eV), see Table 2. This indicates that the range of chemical environments modelled in the relaxed DFT simulations reflect the variation in the experimental sample.

This agreement between experimental and theoretical values, for both CLS and disorder broadening, is experimental confirmation that the electronic structure in the DFT simulation is a reasonable representation of reality. It thus confirms that there exists a re-distribution of electrons (*i.e.* charge transfer between the atoms) that will reduce the size mismatch of the atoms, reducing the suggested local lattice distortion, and that there is a range of chemical environments for each element.

It is interesting to note that there is a substantial difference between theoretical CLS from non-relaxed and relaxed simulations, with the latter values distinctly closer to the experimental results. This stresses the importance of performing structure relaxation in simulations; as well as the strong effect the displacive disorder has





**Fig. 11** XPS core level spectra of the studied HfNbTiVZr coating (red) and elemental reference samples (black), is shown in (a) to (e). Solid vertical lines mark the observed binding energies for the metallic contributions of the lowest energy spin-orbit component:  $2p_{3/2}$ ,  $3d_{5/3}$  or  $4f_{7/2}$ , respectively. Dashed vertical lines mark oxide peaks which formed *in vacuo* during the XPS measurement. An example of the peak fitting of alloy spectra, for Hf 4f, is shown in (f) with experimental data points as circles; individual contributions as blue (metallic) and green (oxide) semi-transparent areas; and the Shirley background as a thin black line. The sum of the different contributions is shown as the solid black line following the experimental data points.

on the CLS values for this alloy, in contrast to alloys with similar sized elements.<sup>33,45</sup>

The observed discrepancies between theory and experiment however also indicate that the DFT-based modelling has not

fully captured the structure of the multi-component alloy. One part of this could be due to the limited size of the supercell, resulting in an incomplete sampling of local order. It should however be noted that part of the discrepancies may be due to a

mismatch between the experimental sample and the theoretical model, as TEM analysis shows a non-homogeneous distribution of Ti and V in the analysed sample.

## Concluding remarks

The here presented results combine theoretical (DFT-based *ab initio* simulations) and experimental (thin film synthesis and characterisation using XPS) results to show that multi-component alloys form non-ideal solid solutions where each atom has a unique chemical environment, and that the chemical interactions (*i.e.* charge transfer) cause a reduction of the atomic size differences assumed based on the metallic radii of the elements. The results show that there exists local lattice distortion (displacive disorder), but to a smaller extent than assumed from the metallic radii.

This has been demonstrated for a system based on early transition metals (HfNbTiVZr), and the same phenomena should be present in other multi-component alloy systems. In cases where the differences between the elements in the multi-component alloy are smaller, the effects will of course be smaller; and conversely one can also envision systems with larger differences and even larger effects. The latter can be an interesting approach for the design of multi-component alloys, by using the here observed charge transfer to influence *e.g.* the bonding character or other functional properties connected to the electronic structure.

A general validity of the here observed charge transfer, and reduction of atomic size differences, will have further implications. This size compensation mechanism may be the reason that the suggested lattice distortion effect has been difficult to confirm experimentally – the real mismatch of atomic radii in the multi-component alloy is, due to the chemical bonding, smaller than assumed from elemental radii. Thus, the often used  $\delta$ -parameter, which is based on tabulated radii, may not be relevant to assess the amount of local lattice distortion in multi-component alloys. One could ask the question if some of the properties of multi-component alloys that have been attributed to the so-called lattice distortion effect, *via* the  $\delta$ -parameter, may instead be the effect of the chemical bonding within the non-ideal solid solution, which will be different than just an average of the included metals.

An additional interesting point is that the presence of charge transfer between the atoms in multi-component alloys may also be transferred to some other multi-component materials. In, *e.g.*, transition metal carbides and nitrides there is metallic bonding between the metal atoms, and it is probable that a charge transfer similar to what has been studied here is present also in the related multi-component carbides and nitrides. Thus, using the here observed charge transfer to design material properties could also be an applicable design route for transition metal carbides and nitrides.

In conclusion, the confirmation of the charge transfer between atoms in multi-component alloys, and connection to (the smaller) lattice distortion, proposed by Tong *et al.*<sup>13–15</sup> opens up new interesting possibilities for materials design for multi-component materials, and may be part of the explanation for non-linear properties observed in some multi-component materials.

## Author contributions

The study was conceptualised by EL; investigation was performed by LCT, BO, DK, SF and KVF; formal analysis was performed by LCT, IAA, BO, RL, and AIA; writing of the original draft was performed by LCT, BO, EL, and AIA; reviewing and editing by EL, RL, BA, IAA and UJ; BA, IAA, IH, UJ, and EL were responsible for supervision; UJ, IAA, IH, and BA contributed to funding acquisition.

## Conflicts of interest

There are no conflicts of interest to declare.

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