Structural degradation of tungsten sandwhiched in hafnia layers determined by in-situ XRD up to 1520 °C

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The high-temperature stability of thermal emitters is one of the critical properties of thermophotovoltaic (TPV) systems to obtain high radiative power and conversion efficiencies. W and HfO2 are ideal due to their high melting points and low vapor pressures. At high temperatures and given vacuum conditions, W is prone to oxidation resulting in instantaneous sublimation of volatile W oxides. Herein, we present a detailed in-situ XRD analysis of the morphological changes of a 3-layer-system: HfO2/W/HfO2 layers, in a high-temperature environment, up to 1520 °C. These samples were annealed between 300 °C and 1520 °C for 6 h, 20 h, and 40 h at a vacuum pressure below 3 × 10–6 mbar using an in-situ high-temperature X-ray diffractometer, which allows investigation of crucial alterations in HfO2 and W layers. HfO2 exhibits polymorphic behavior, phase transformations and anisotropy of thermal expansion leads to formation of voids above 800 °C. These voids serve as transport channels for the residual O2 present in the annealing chamber to access W, react with it and form volatile tungsten oxides. An activation energy of 1.2 eV is calculated. This study clarifies the limits for the operation of W-HfO2 spectrally selective emitters for TPV in high-temperature applications.

A major portion of industries globally, still rely on fossil fuels as their primary resource of energy1. A significant amount of this energy is wasted as unexploited heat, named waste heat. One means of efficiently utilizing the wasted heat to harness electricity can be realized by using a thermophotovoltaic (TPV) system2. A TPV system can as well produce electric power from fossil fuels1, solar radiation4–8 and in radioisotope thermoelectric generators (RITEGs)9–11. In TPV, an emitter is heated to high-temperatures, typically above 1000 °C, and its radiated thermal energy is directly converted into electricity using photovoltaic (PV) cells12–15. The efficiency of the TPV system can be enhanced by using spectrally selective emitters16–23. Such selective emitters are designed to precisely emit spectrally tailored radiation that matches the band gap of the PV cell. One of the critical parameters for obtaining high power and efficiency is operating the emitters at high-temperature, as the radiated power of a black body is proportional to the Stefan Boltzmann law25 and the spectral maximum shift to shorter wavelengths proportional to 1/T which is known as Wien's displacement law and can be derived from Planck's law of radiation. This implies that the emitter should work at 1400 °C temperature, to match the bandgap Eg of high efficiency GaSb PV cell at 0.72 eV26. It is worth mentioning that besides increasing the total radiated power by almost a factor of 4, a raise in temperature from 1000 to 1400 °C will also increase the TPV efficiency of a metamaterial emitter from 33 to 48%27. Therefore, high-temperature stability and long lifetime at these temperatures are required for the effective performance of a TPV system.
In this perspective, appealing choices of metals with high melting point are Mo (2620 °C), Ta (3020 °C) and W (3410 °C), while Al₂O₃ (2054 °C), ZrO₂ (2715 °C), and HfO₂ (2800 °C) are suitable alternatives among the dielectrics²⁸–³⁰. Photonic crystals³¹, gratings³², and multilayered metamaterials³³ comprising of alternate layers of metal and dielectric are examples of selective emitters. Arpin et al.³⁴ demonstrated a stability of 1000 °C in their W based 3D photonic crystal prepared by conformal deposition of W using atomic layer deposition (ALD). Yokoyama et al.³⁵ used a combination of Mo and Al₂O₃ to fabricate a metal–insulator–metal thermal emitter that reached an operational temperature of 1000 °C. Thermal stability of a W/Al₂O₃ based metasurface solar absorber up to 1200 °C was demonstrated by Chang et al.³⁶ using in-situ high-temperature reflectance experiments. Kim et al.³⁷ reported a multilayer metamaterial design consisting of W/Al₂O₃ layers, which showed thermal stability up to 1200 °C for 3 h. The degradation in optical properties was observed due to the agglomeration of W within the layers with further increase in the temperature. Kohiyama et al.³⁸ and Shimizu et al.³⁹ carried out high-temperature experiments up to 1360 °C on selective absorber/emitters, which were a combination of layered multilayers of Mo/HfO₂ and W/Yttria stabilized zirconia (YSZ). Often the cause of structural failure at high-temperatures is due to oxidation³³,⁴⁰–⁴² and surface diffusion⁴³–⁴⁵ in the layered stacks. The other notable mechanisms that are observed at high-temperatures are stress induced in the multilayers leading to delamination and cracking⁴⁶.

In a more general perspective for high-temperature stable metamaterials, not only the melting point but as well the vapor pressure⁴⁷ of both layer materials is important for a good thermal stability. It is known that materials which attain a high vapor pressure below their melting point on heating, tend to sublime directly into the gaseous phase at a temperature below their melting point⁴⁸. Therefore, the materials selected for a high-temperature application should possess a low vapor pressure: W and HfO₂ are exceptional in their classes of materials. The vapor pressure of W is 8.15 × 10⁻¹⁰ mbar at 2000 °C⁴⁹ is the lowest among metals and that of HfO₂ is 1 × 10⁻¹³ mbar at 1732 °C⁵⁰, shows exceptional low vapor pressure among dielectrics.

Herein, we focus on in-situ X-ray diffraction (XRD) experiments of a simplified 3-layer-system of 100 nm HfO₂/20 nm W/100 nm HfO₂ shown (Fig. 1a–c) in order to investigate time-dependence of the structural and morphological changes of the layer materials at high-temperatures. The repetition of a 3-layer-system can be used to construct a multilayered selective emitter. The 3-layer-system is prepared by magnetron sputtering, a method well established for depositing refractory materials, that facilitates the growth of highly uniform and precise multilayers³¹–³³. The primary goal of using a 3-layer-system is to understand the mechanism of oxidation in a single W layer that leads to initial structural changes, followed by degradation at various annealing temperatures and durations. Chemical composition and morphology of the 3-layer-systems after preparation and after different annealing experiments were characterized by scanning transmission electron microscopy—energy dispersive X-ray spectroscopy (STEM-EDS). Further, we discuss the changes observed in the 3-layer-system due to high operational temperatures such as phase formation, changes in interplanar spacings and grain sizes, and alterations of the volume fractions of the observed phases. Structural degradation mechanisms of W films at high vacuum conditions and above 1000 °C operational temperatures are demonstrated, which reveals the current limitations in the life-time of W-based metamaterials. These measurements allow a better understanding of the underlying mechanisms of the failure of W/HfO₂-multilayered metamaterials at high-temperatures.

Results and discussion
Ex-situ X-ray diffraction measurements of two identical 3-layer-systems after 6 h annealing experiments allow us to investigate the changes of crystal structure and phase formation under high vacuum conditions at two different temperatures. Slight variations are noticeable after the moderate annealing at 650 °C (Fig. 2a) and distinct changes are visible after 1520 °C (Fig. 2b). Before annealing, it is always important to control that the as-prepared samples are comparable, which can be confirmed by nearly identical diffraction patterns, which contain only a few peaks that corresponds to three phases in the recorded range. The peaks (110) at 39.9° and (200) at 58.3° correspond to the body centered cubic (bcc) phase of α-W. The α-phase is the thermodynamically stable phase of W and the metastable β-phase of W is not present in our films⁵⁴. In the case of HfO₂, a coexistence of a nanocrystalline monoclinic phase and an amorphous phase is observed after preparation. The (−111) peaks at 28.3° can be attributed to the monoclinic phase of HfO₂, whereas the broad bump around 30.4° corresponds to the amorphous phase of HfO₂. In the case of the monoclinic HfO₂, the full width at half maximum
of the (−111) peak is only 0.48°, whereas the amorphous phase of HfO₂ at 30.4° possesses a broad peak with a FWHM value of 5.2°. From the peak shape of the diffraction pattern, it is estimated (using Bruker EVA software) that the as-prepared HfO₂ layers have volume fraction of 56% and 44% for the crystalline and amorphous phase after preparation. Due to annealing, already after 650 °C, the crystallinity of HfO₂ is increased, since the broad peak of the amorphous phase has vanished, and some new crystalline peaks of the monoclinic phase appeared. Furthermore, after 1520 °C, the peaks become sharper and a number of new peaks are identified, all of them attributed to the monoclinic phase of HfO₂. A slight change in the (110) W peak position to 40.1° is observed, since the bcc W structure relaxes towards the thermodynamic equilibrium position of bulk W, which is 40.265° (PDF card 00-004-0806) at room temperature. These changes are discussed later in more detail. It is most likely that new peaks from the monoclinic phase of HfO₂ occur during annealing as the existing grains grow, in order to attain a new position due to relaxation. In summary, after annealing to either 650 °C or 1520 °C only two phases are evident, the bcc phase from W and the monoclinic phase from HfO₂. It is worthwhile to mention that there is no evidence of any solid-state reactions between the two-layer materials in the 3-layer-system such as WOₓ, Hf-W-O alloys, and pure Hf. The two obtained phases only present small shifts in the peak positions for two different annealing temperatures, but there is no hint for any phase or composition change during annealing.

In an in-situ annealing experiment, one can clearly visualize transient phase formation and their changes as a function of time and temperature. Diffraction patterns were recorded sequentially during ramp-up of the temperature. Using controlled heating rates enables us to pursue reaction kinetics and material transport processes undergone by the sample.

The diffraction patterns of a 3-layer-system recorded at different temperatures during the ramp part of the annealing experiments are shown in Fig. 3, from room temperature up to 1520 °C. In the first instance, it is clearly noticeable that the peaks get sharper as the temperature increases and move to higher angles due to relaxation approaching the equilibrium value at the adjusted temperature. The broad, amorphous bump of HfO₂ at room temperature slowly starts to vanish. It is well known from the literature that HfO₂ exhibits polymorphic...
behavior. A new peak (101) from the tetragonal phase of HfO$_2$ (Supplementary Figs. 1, 2) emerges from the amorphous region at 30.3°, as the temperature rises to 200 °C. This peak continues to grow in intensity and tends to become narrower until the ramp temperature reaches 750 °C, after which there is a decline in (101) peak intensity. The d-values of this tetragonal phase are nearly constant with $d_{101} = 0.2951 \text{ nm}$ at 200 °C and $d_{101} = 0.2968 \text{ nm}$ at 750 °C, and this small change is associated to thermal expansion. On the other hand, the intensity of the (−111) peak, the strongest X-ray spot from the monoclinic HfO$_2$ phase starts to grow higher in intensity, where it is doubled at 500 °C and tripled at 1250 °C. This is a clear indication that the tetragonal phase formed at 200 °C is transformed into the highly stable monoclinic phase at high temperatures. These results are in good agreement to the experiments carried out by Ushakov et al. and Zhao et al., as they observed amorphous HfO$_2$ phases that crystallize into tetragonal phase first and later transform into the stable monoclinic phase in thin films, which were prepared by sol–gel, pulsed laser deposition (PLD) and atomic layer chemical vapor deposition (ALCVD). As the temperature increases above 1000 °C, we do not see any evidence for a tetragonal phase, but more new peaks corresponding to the monoclinic phase of HfO$_2$ start to appear such as (111) and (−201). We do not observe any phase change in the W layer except the sharpening of the (110) W peak during the heating ramp.

The second and most important stage in the temperature excursion of the 3-layer-system is the isothermal process. Here, diffraction patterns are recorded as a function of time under constant temperature. These measurements yield significant information on the kinetics of degradation observed in the 3-layer-system. 2θ scans in the range of 26°–43° at 1520 °C spanning over 6 h are displayed in Fig. 4. There is a slow increase in 2θ values of both the (−111) monoclinic HfO$_2$ and (110) bcc W peaks. The peaks get sharper over time and the FWHM of both the peaks decrease due to grain coarsening. There is no indication of any phase or composition change appearing in the diffraction patterns, which is ideal for the application as high-temperature stable material. The most striking feature to observe is the intensity reduction of the (110) bcc W peak over time, while the (−111) monoclinic HfO$_2$ remains constant during the 6 h annealing period. After cooling down the 3-layer-system to room temperature, the diffractograms do not show any additional phase of HfO$_2$ and it persists in the monoclinic phase. This is represented in the last scan (brown color, Fig. 4) of the sequence of diffraction patterns, in which a shift in 2θ value to higher angle is observed due to thermal contraction. It is important to mention that the final properties of the 3-layer system after in-situ and ex-situ experiments are similar, but it also indicates that the in-situ scans give us the tools to reveal information about internal and as well irreversible changes of structure and microstructure.

The time-dependent change of the interplanar spacing $d_{110}$ of the bcc W phase allow us a first insight of the degradation process. Therefore, the influence at two annealing temperatures 650 °C (red markers) and 1520 °C (blue markers) was measured (Fig. 5a). Before annealing, the d-values measured at room temperature for the two samples are nearly the same, since the values are $d_{110} = 0.2255 \text{ nm}$ for the sample 650 °C and $d_{110} = 0.2256 \text{ nm}$ for sample 1520 °C. The small vertical offset (of less than 0.0001 nm) between the room temperature values is a systematic error caused by the initial height adjustments of the high-temperature stage before the start of the experiment. In general, the interplanar spacing $d_{110}$ without any stress relaxation or other irreversible process increases with higher temperature during annealing. However, it is essential to compare steady- state d-values at two different fixed temperatures to understand the significance of grain growth, stress relaxation and recrystallization with respective to the high-temperature stability. Initially, there is a small increase of d-value until 200 °C the first five points (highlighted in green ellipse, Fig. 5a), this is due to the thermal expansion as the atomic mobility is still too small for stress relaxation. The observed thermal expansion of W film is $4 \times 10^{-6} \text{ K}^{-1}$ and it is in good agreement with bulk W, i.e. $4.12 \times 10^{-6} \text{ K}^{-1}$. Using magnetron sputtering at low Ar pressure of...
The grain size changes of monoclinic HfO$_2$ during heating ramp and isothermal stages at 650 °C and 1520 °C various annealing temperatures are exploited to determine an activation energy for this structural degradation, which is described later.

The calculated volume fraction of the W disappearing in the 20 nm thick W layer sandwiched between two HfO$_2$ layers is estimated to be 79% at the end of the 6 h isothermal stage at 1520 °C. Further experimental results at various annealing temperatures are exploited to determine an activation energy for this structural degradation, which is described later.

Microstructural information like grain size is as well contained in the broadening of X-ray diffraction peaks. The grain size changes of monoclinic HfO$_2$ during heating ramp and isothermal stages at 650 °C and 1520 °C are shown in Supplementary Fig. 4. The grain size value is calculated from the (−111) monoclinic HfO$_2$ peak area under the diffraction peak) values of (110) bcc W peak annealed at 650 °C (blue markers) and 1520 °C (red markers) shown in Fig. 5b. From quantitative analysis of powder mixtures, it is known that the integrated intensity of a diffracted peak for a particular phase depends on the concentration of the phase. This indicates that a relative change of integrated intensity of the (110) bcc W peak is proportional to a change of the volume fraction of the W present in the 3-layer-system. The integrated intensity of the sample annealed at 650 °C shows a constant behavior during the temperature ramp-up, and it remains relatively stable during the whole isothermal stage. Whereas in the sample annealed at 1520 °C, the integrated intensity remains relatively constant during the temperature ramp-up and then drastically decreases during the isothermal part of the annealing experiment. Any loss in the integrated intensity of (110) W peaks directly indicates a loss in volume fraction of W in the 3-layer-system. A change in roughness could also explain a reduction in peak intensity, but under such circumstances, the intensities of all occurred phases for HfO$_2$ and W should be reduced. Clearly, this is not observed for the intensity of the monoclinic HfO$_2$ phase (Fig. 4), that appears fairly constant over the whole isothermal stage. The growth process of the W grains is in principle similar (Supplementary Fig. 5), however in the isothermal part at 1520 °C, the atomic loss in the W grains is predominant due to sublimation of W oxide, which is discussed in detail later.

To achieve a deeper understanding of the structural changes in W and HfO$_2$ layers due to in-situ annealing experiments, TEM-EDS characterizations on the as prepared and annealed samples were performed. In the case of the as prepared sample, the STEM-high-angle annular dark-field (STEM-HAADF) image and the elemental

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Figure 5. (a) Interplanar spacing $d_{110}$ of bcc W in the 3-layer-system as a function of time during annealing up to 650 °C (blue markers) and 1520 °C (red markers). The duration of the isothermal part was about 6 h. The green ellipse highlight the initial increase in d-values due to thermal expansion below 200 °C. Integrated intensity of the monoclinic HfO$_2$ phase (Fig. 4), that appears fairly constant over the whole isothermal stage. The intensity of the (110) bcc W in 3-layer-systems annealed for 6 h at temperatures 650 °C (blue markers) and 1520 °C (red markers).
mapping (Fig. 6a,d,g) show a distinct 3-layer-system. After annealing at 650 °C for 6 h, there are no modifications in the layer stack (Fig. 6b,e,h), the interface between W and HfO₂ layers appears to be unchanged. No grain boundaries were seen in the studied TEM lamella. Figure 6a,b were captured at different camera lengths, this results in different diffraction contrast, leading to bright and dark lines above and below the W layer. No additional phase of W is observed in the EDS mapping. On the contrary, after 1520 °C and 6 h, significant changes in morphology occur (Fig. 6c). An increase in the interface roughness of the HfO₂ layers is clearly visible and the interface becomes wavy in contrast to the sharp interfaces prior to annealing. Individual grains are clearly seen with distinct grain boundaries due to the increase in grain size. The EDS mapping of W (Fig. 6f) shows a very weak presence of W in the layer, which corroborates the XRD results indicating the reduction of W volume fraction (Fig. 5b) after annealing at 1520 °C. In addition, the EDS mapping of Hf (Fig. 6i) after annealing at 1520 °C for 6 h confirms the presence of Hf in the HfO₂ layer and no diffusion of Hf into the W layer.

In order to understand the root cause for the disappearance of the W from the sandwiched layer, further heating experiments were conducted to estimate the total time required for the entire W layer to disappear from the 3-layer-system and to investigate the remaining morphology. A 3-layer-system was annealed for 40 h at 1520 °C for this purpose. The top surface and cross-section SEM images of the 3-layer-system annealed at 1520 °C for 40 h are shown in (Fig. 6j,k). The presence of voids distributed on the top HfO₂ layer is visible, which is in good agreement to early studies on crystallization of HfO₂ single films56. In our system, they extend through the entire thickness of the first HfO₂ layer, illustrated by the cross section view through the same void shown in Fig. 6k. It is also noteworthy to observe the complete absence of W in the 3-layer-system sample. The voids present in the HfO₂ layer act as a transport channel for the residual O₂ molecules in the heating chamber to reach the W layer, resulting in oxidation and the formation of volatile WOₓ₆⁶. As seen previously for the 6 h annealed 3-layer-system, a similar trend in the loss of intensity from the (110) W peak occurred, and after 20 h (see below in Fig. 8a), the intensity of the W peak was reduced considerably and was almost in-line with the background in the measured diffraction pattern. The volatile WOₓ sublimate rapidly at 1520 °C, leaving the layer through the same transport channels. Consequently, no WOₓ peaks are visible in the diffractograms. A plain sapphire “witness” substrate was therefore placed on the wall of the heating chamber close to the outlet of the vacuum pump to collect any sublimated material via re-deposition. The redeposited layer on the witness sample can be identified as WOₓ from the diffraction pattern (Supplementary Fig. 6). The possibility of W obtaining O₂ from HfO₂ can be excluded, as this reduction reaction would either result in the formation of pure Hf or result in sub-stoichiometric HfO₂. No Hf peaks are observed in any of our in-situ X-ray diffraction experiments. In addition, and complementarily, characterization by Rutherford backscattering spectrometry (RBS) (Supplementary Fig. 7) confirms no presence of pure Hf but the appearance of slightly over stoichiometric HfO₂ₓₓ and especially the loss of W.

The complex oxidation mechanism of W is extensively discussed in literature65–74. It is possible to split the oxidation mechanism of W into four different zones up to a temperature of 1500 °C depending on the partial

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**Figure 6.** STEM-EDS analysis of the 3-layer-system, as-prepared, annealed at 650 °C and 1520 °C for 6 h. (a–c) STEM-HAADF images, (d–f) W-Mα elemental mapping, (g–i) Hf-Mα elemental mapping. These images have the same scale as indicated in image (c). SEM images of the 3-layer-system annealed at 1520 °C for 40 h: (j) morphology of the top HfO₂ layer with a void in the middle and (k) cross section view along the interfaces across the same void (FIB section).
pressure of O₂. The first zone is comprised of heating experiments conducted until 500 °C in the partial pressure of O₂ higher than 100 mbar⁶⁵, an increase in the mass of the bulk W samples is observed during heating experiments. The increase in mass is attributed to the formation of an oxide layer on top of the sample. The oxidation of W is an exothermic process, where heat is liberated when W and O₂ react to form WOₓ having a negative enthalpy of formation⁷⁵ and the oxidation process is diffusion-controlled, requiring an activation energy of 1.92 eV. The second zone comprises the temperature range between 500 °C until 1100 °C in an O₂ partial pressure of 100 mbar⁶⁶, where complete oxidation of the W sample occurs. Initially, the oxidation process follows a parabolic rate law due to the diffusion of O₂ through the already formed WO₃ top layer. Later, the formed oxide cracks due to stress exposing the underlying metal and then the oxidation process follows a linear rate law. The activation energy for oxidation in this temperature range is 1.4 eV. The third zone of oxidation can be distinguished by increasing the temperature to 1250 °C at 50 mbar O₂ pressure, where all the O₂ reacts to form volatile WO₃ and no formation of any oxide layers were visible on the sample surface. Thus, the rate of WO₃ sublimation was quicker than the rate for the formation of a solid oxide layer above 1250 °C. In our previous annealing experiment²⁷ performed at 1200 °C and in a vacuum of 2 × 10⁻² mbar, the W present in the multilayered metamaterial was oxidized and the presence of an oxide layer of WO₃ on top of the first W layer was confirmed by XRD and HRTEM characterization. The fourth zone of oxidation mechanism can be distinguished for temperatures between 1250 and 2300 °C in a vacuum less than or equal to 1 × 10⁻⁵ mbar. Only weight loss is observed in all the W samples without the formation of a solid oxide layer, as the concentration of O₂ atoms is relatively small. The activation energy in this zone can vary from 0.6 to 0.86 eV depending on the temperature.

In our current in-situ annealing experiments conducted at 2 × 10⁻⁶ mbar, the diffractograms do not show any peaks of WOₓ during the entire heat treatment cycle (Figs. 3 and 4). Based on the results of the 40 h annealing experiment, further annealing experiments, for a duration of 20 h at different temperatures between 650 and 1520 °C, were performed. The objective of the latter experiments was to determine the temperature range where the voids start to originate in the HfO₂ layers. It is observed from the SEM images (Supplementary Fig. 8) that voids appear on all samples annealed above 800 °C. This temperature range can be related to the transformation of the tetragonal phase to the stable monoclinic phase discussed previously. A change in structure from tetragonal to monoclinic results in a volume increase accompanied by shear strain. It is also to be noted that the symmetry of the structure is reduced from tetragonal (a = b ≠ c, ß = 90°) to monoclinic (a ≠ b ≠ c, ß ≠ 90°). This results in anisotropic behavior of thermal expansion, where the minimum and maximum thermal expansion values do not lie in the lattice directions for a monoclinic structure. The coefficient of thermal expansion values of the monoclinic phase range between 8 × 10⁻⁶ K⁻¹ to 32 × 10⁻⁶ K⁻¹ at 1750 °C⁷⁷. Therefore, the tetragonal-to-monoclinic phase transformation in HfO₂ and the anisotropic nature of the monoclinic phase results in the formation of voids between the HfO₂ grains in the top HfO₂ layer. The mechanism of void formation and the oxidation of the W layer are summarized in Fig. 7.

The integrated intensity of the (110) bcc W peak of the sample annealed at 800 °C decrease only slightly after 20 h (Fig. 8a). On increasing the temperature, the loss of intensity is observed for the sample annealed at 1000 °C and 1240 °C. Moreover, the sample annealed at 1520 °C shows a sharp decrease in intensity during the initial 3 h
of the isothermal stage and later the process slows down to a reduced rate. The change in the volume fraction is estimated by calculating the slopes for the drop in integrated intensity of the (110) bcc W peak (filled circles in the plots were used for the calculation of slope, Fig. 8a). This thermally-activated time-dependent process is represented by an Arrhenius plot (Fig. 8b). The change in the integrated intensity of the (110) bcc W peak over time is expressed as a function of the inverse of the annealing temperature. An activation energy of 1.2 eV is calculated for O₂ to penetrate through the transport channels, oxidize the W and sublime as volatile WOₓ. Becker et al. calculated an activation energy of 1.1 eV, which is in good agreement with our experiments. They performed an estimation for the conversion of WO₃ at a pressure of 1 × 10⁻⁶ mbar and in the temperature range between 920 to 1320 °C, when O₂ molecules strike the surface of a W ribbon. In our experiments, the rate of W loss can also depend on the number and the size of pores in HfO₂. The HfO₂ heated to higher temperatures can provide more or larger channels for O₂ attack and thus increased rate of W loss. Thus, the obtained activation energy can be alternatively linked to the surface diffusion of HfO₂ that leads to grain growth and void formation.

At a temperature below the phase transformation in HfO₂, no sign of degradation is observed in the W layer. The tetragonal phase was retained on cooling the 3-layer-system annealed at 300 °C for 20 h, back to room temperature. It is worthwhile to mention that an annealing experiment at 700 °C and over a time of 100 h do not cause any remarkable degradation to the sandwiched W layer, only above this temperature, which coincides with the tetragonal to monoclinic phase transformation in HfO₂, we observe the degradation of W in our 3-layer-system.

Conclusion

The thermal stability and durability of the selective emitters at high-temperature are the essential prerequisites for the effective functioning of the TPV system. In order to meet these requirements, a 3-layer-system comprising of HfO₂/W/HfO₂ was fabricated using magnetron sputtering. The high-temperature stability of the 3-layer-system was evaluated by performing in-situ XRD annealing experiments at various annealing temperatures between 300 and 1520 °C at three different periods 6 h, 20 h, and 40 h and at vacuum pressures of 3 × 10⁻⁶ mbar.

The disappearance of W in the 3-layer-system is validated by the drop in the integrated intensity of the (110) bcc W peak during the isothermal part of all the annealing experiment performed over 1000 °C. The mechanism behind the degradation of the W layer begins with the formation of voids in the HfO₂ layers that acts later as transport channels. The tetragonal-to-monoclinic phase transformation results in volume increase, which is accompanied by the anisotropic nature of the monoclinic HfO₂ phase, and culminates in the formation of voids. Residual O₂ present in the heating chamber reaches the W layer through the transport channels and forms volatile oxides that sublimate rapidly. An activation energy of 1.2 eV is determined for the oxidation of W in the 3-layer-system through the transport channels, by performing additional annealing experiments between 800 and 1520 °C for 20 h.

Although the combination of W/HfO₂ excel exceptionally to meet the optical properties of the selective emitter, HfO₂ fails to act as a good protective layer above 1000 °C. The thermal stability and durability at high-temperature of the 3-layer-systems as well as multilayered metamaterials could be further enhanced by reducing grain growth and avoiding a phase transformation like the tetragonal-to-monoclinic transition observed in HfO₂, thereby preventing the formation of voids in the oxidic dielectric layers. We will investigate various combinations of metals for instance suitable W alloys and dielectrics such as Y-stabilized HfO₂ in order to achieve a higher working temperature and longer lifetime of selective emitters for TPV systems.

Methods

Thin-film deposition using magnetron sputtering. All 3-layer-systems used for annealing experiments were prepared by magnetron sputtering, using our in-house designed sputtering facility at the Helmholtz-Zentrum Geesthacht. The chamber was evacuated to a base pressure below 1 × 10⁻⁷ mbar before the start of the
thin film deposition process. The W and HfO₂ sputtering targets (Sindlhauser Materials GmbH) had a purity of 99.95% and were 3" in diameter. The 3-layer-system HfO₂/W/HfO₂ was deposited on a sapphire substrate 10 mm × 10 mm × 0.5 mm (purchased from CrysTec GmbH) having a (1–102) orientation and thermal expansion coefficients of 5.3 × 10⁻⁶ K⁻¹ (parallel to c-axis) and 4.5 × 10⁻⁶ K⁻¹ (perpendicular to c-axis). The substrates are mounted on a custom-made holder equipped with circulating cooling water system and are rotated at a rate of 1 Hz. The temperature of the cooling water is maintained below 20 °C throughout the deposition of the 3-layer-system. The distance between the target and the substrate is 14 cm. The calibration of the individual deposition rate of W and HfO₂ was done separately by deposition of single layers on silicon substrates 20 mm × 60 mm × 0.7 mm having a (100) orientation parallel to the substrate normal. Argon gas of purity 99.99999% at a flow rate of 28 sccm was used as sputtering gas and the sputtering pressure was maintained at 2 × 10⁻⁶ mbar. The thin-film preparation of the W films was done using a DC power of 120 W, with a deposition rate of 0.07 nm s⁻¹, whereas HfO₂ films were deposited using RF power of 400 W, which yielded in a deposition rate of 0.15 nm s⁻¹.

**In-situ X-ray diffraction experiments.** A diffractometer, equipped with a specific high-temperature chamber for annealing up to 2000 °C (Anton Paar—HTK 2000N), and a state-of-the-art position sensitive detector (Lynxeye XE-T, Bruker AXS) with an energy resolution of less than 380 eV was used to perform all the annealing experiments. Thermal expansion calibration experiments were performed with sapphire substrates up to 1900 °C, in temperature steps of 100 °C, and using a heating rate of 25 °C min⁻¹. A full pattern fit was performed for a 2θ range from 50° to 125° and the lattice parameters of the sapphire substrate at every 100 °C interval were refined using the Rietveld method (TOPAS)⁷⁹. The temperatures were later validated from the thermal expansion coefficients determined by Touloukian et al.⁵⁸. The heating chamber was evacuated to a pressure less than 3 × 10⁻⁶ mbar before every annealing experiment. The x-height of the heating stage was adjusted better than 10 µm by using the sapphire substrate peak positions according to Powder Diffraction File (PDF)³⁹ card 01-082-1468. The XRD 2θ scans were measured from 20° to 120°, with an increment of 0.02° and a time rate of 0.15 nm s⁻¹.

The layer thickness of W and HfO₂ were determined precisely by means of X-ray reflectometry (XRR), using Cu Kα₁ radiation (λ = 0.15406 nm). The layer properties were evaluated by fitting simulations to experimental reflectivity scans with the Leptos R software package (Bruker AXS).

**TEM and EDS characterization.** The cross-sectional transmission electron microscopy (TEM) samples, and cross-sections for scanning electron microscopy (SEM) were prepared using a FEI Helios G3 UC focused ion beam scanning electron microscope (FIB-SEM). Both the cross-sectional TEM lamellae and cross-sections were prepared using 30 keV Ga⁺ ion beam and different beam currents (9 nA to 40 pA). Secondary electron (SE) images of the cross-sections were taken under low keV (2 keV) electron beam and high-resolution immersion mode using a through lens detector (TLD). To prevent charging during the FIB-SEM study, the sample was coated with a thin layer of carbon by means of a carbon thread evaporator. A FEI Talos F200X TEM equipped with a four-quadrant silicon drift detector system for energy dispersive X-Ray spectroscopy [Super-X(FeI)] system was used for taking high angle annular dark field (HAADF) images and to perform EDS analysis on 10 µm × 100 nm lamellae. Spectrum images (SI) were acquired using a 1 nA beam current and dwell time of 10 µs per pixel. Resolution was 512 × 512 pixels with a pixel size of 766 pm. To get a better signal-to-noise ratio integrated intensities of over 1000 drift corrected SI were used.

Received: 27 October 2020; Accepted: 25 January 2021
Published online: 08 February 2021

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Author contributions
G.V.K. and M.S. designed the experiments, fabricated the thin films, performed in-situ annealing measurements, performed XRD measurements and analyzed the diffraction patterns regarding phase formation and structural changes in the films. S.S.R., T.K., and M.R. performed the SEM–FIB, STEM-imaging and elemental mapping. H.W.B. performed the RBS experiments and analyzed the data. G.V.K., M.C., K.P.F., A.Y.P. and M.E., interpreted the experimental results and performed oxidation and thermal analysis and M.S. supervised the project. All authors discussed the results of the manuscript and G.V.K. wrote the manuscript with inputs contributed from all the authors.

Funding
Open Access funding enabled and organized by Project DEAL. The authors gratefully acknowledge financial support from the German Research Foundation (DFG) via SFB 986 ‘Tailor-Made Multi-Scale Materials Systems: M3’, Project-ID 192346071, and projects C1, C7 and Z3.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-021-82821-0
