The role of doping and microstructure on hydrogen solubility in monoclinic ZrO$_2$: Experimental validations of simulated defect chemistry

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**ABSTRACT**

The effect of chemical doping and ceramic microstructure on H solubility was measured systematically for the first time in monoclinic ZrO$_2$. Excitingly, the influence of chemical doping by Fe, Cr and Ta cations on bulk H solubility was in qualitative agreement with defect chemistry simulations based on density functional theory and statistical thermodynamics. This is the first experimental validation of the recently-developed modeling framework, which is capable of high-throughput defect chemistry computations of light elements in metal oxides. H solubility was quantified using temperature-programmed desorption and was measured for various microstructures and under varied oxygen partial pressure, enabling us to identify the microstructural origin (i.e., bulk, surfaces and grain boundaries) of sorbed H defects and infer their defect type. Doping the oxide resulted in a significant increase in the H-uptake time, indicating that solute cations slowed H defect diffusion. Ta doping offered the lowest bulk H solubility at all measurement temperatures (410° C – 60 °C), though Fe doping also lowers bulk solubility at lower temperatures and yielded the most tightly bound H defects of all oxides. This could be significant in applications where one wishes to minimize H permeation via limiting diffusion kinetics. Based on this work we emphasize the role of both doping and microstructure on the overall H uptake and highlight the importance of elucidating the relative nuances of each oxide system; we thus reiterate the value of a high-throughput computational analysis.

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1. Introduction

Metal oxide ceramics and thin films are employed broadly in the science and engineering of advanced energy materials and nanotechnology. Absorption and adsorption of hydrogen, hydroxyls, and water by these materials is critical to their performance in energy conversion [1,2], harvesting [3] and storage [4–6] applications. There has been sustained interest in solid oxide cells that employ metal oxide catalysts to perform hydrogen and/or water dissociation reactions [1], or rely on solid H electrolytes [7]. Thermochemical syngas (H$_2$ and CO) production schemes using reducible metal oxides to remove oxygen from water and CO$_2$ are also under investigation [2]. Absorption of H was shown to improve conductivity without sacrificing transparency in polycrystalline ZnO, which was proposed as an alternative to Sn-doped indium oxide in Si-based solar cells [3]. Li$^+$ charge transfer across the electrolyte-electrode interface critically impacts solid state battery performance, and studies show that protonation of the solid oxide Li$^+$ electrolyte is a key reaction in the formation of Li-blocking carbonates at this interface [4,6]. The ability to reversibly hydrate VO$_2$ thin films was also demonstrated recently, motivating their proposal as a novel H storage medium [5]. In the context of nuclear energy, water and H absorbed by UO$_2$ [8–10] has been cited as a source for H which embrittles Zr alloy cladding, whose oxide scale has also been the subject of hydration studies [11–13]. Gaining a fundamental understanding of the chemical interactions between H and metal oxides – as well as developing the ability to predict them with high-throughput computational simulations – undergirds progress in materials science and will accelerate the rational design of future materials.

The present work focuses on quantifying and elucidating H solubility in cation-doped monoclinic ZrO$_2$ (α-ZrO$_2$), which was the subject of a recent computational study by our group [13]. In that work, Youssef et al. developed a thermodynamic modeling framework based on density functional theory (DFT) [13–15] that predicted chemical and charge transport properties critical to H uptake in doped α-ZrO$_2$. For example, they showed that the predominant H point defects in doped α-ZrO$_2$ at 600 K are either interstitial H or a H-Zr vacancy complex, depending on the dopant species. Importantly, that work produced material design guidelines for mitigating H uptake based on (i) thermodynamically lowering the H solubility and (ii) kinetically accelerating H$_2$ evolution at the α-ZrO$_2$ surface in contact with H$^+$. However, a first experimental validation of this framework is absent from the literature. If validated the framework could provide high-throughput
analyses and predictions of many metal oxides, particularly those containing light-element point defects such as H and Li.

Beyond serving as a model oxide, $\alpha$-ZrO$_2$ passes Zr alloys in nuclear reactors [11,16], and has been studied for heterogeneous catalysis [17–22] and photocatalytic water splitting [23] applications. However, experimental studies quantifying H solubility in this material appear limited to undoped $\alpha$-ZrO$_2$. Yamanaka et al. measured solubility in sintered $\alpha$-ZrO$_2$ pellets hydrated between 500 – 1000 °C and showed that solubility decreases with increasing hydration temperature from $10^{-4}$ to $10^{-11}$H/ZrO$_2$ [24]. In tetragonal ZrO$_2$, Park and Olander measured similar levels of H solubility at 1300 °C – 1600 °C [25]. We hope this work can illuminate the positive and negative impacts of various cation dopants selected based on our initial computational work mentioned above.

We also aim to elucidate the impact of microstructural features on H sorption in doped $\alpha$-ZrO$_2$, where quantitative studies again appear to be limited to undoped $\alpha$-ZrO$_2$. In sintered polycrystalline $\alpha$-ZrO$_2$, Yamanaka et al. measured distinct H$_2$O desorption events around 150 °C and 500 °C, attributed to surface and bulk desorption, respectively [24]. Attribution of low-temperature desorption to surface-physorbed H$_2$O was corroborated by Kouva et al. via desorption from $\alpha$-ZrO$_2$ particles [20]; the contribution of internal surfaces and interfaces to H absorption has not been clarified. Navio et al. showed using IR spectroscopy that hydroxyl and molecular water adsorbed on Fe-doped ZrO$_2$ (< 5 wt% Fe) nanoparticles [26]. Studies of H permeation through oxide scales of Zr alloys indicate that grain boundaries, pores, and cracks are diffusion pathways facilitating oxidative corrosion. Sundell et al. investigated oxide chemistry by atom probe tomography and showed that Ni, Fe$^{2+}$ [27] and OH$^-$ [11,12] decorated grain boundaries in the corrosion scale on Zircaloy-2 and Zr-2.5Nb. However, there persists a gap in the qualitative and quantitative understanding of the effect of doping $\alpha$-ZrO$_2$ on H sorption at microstructural features like pores, surfaces and grain boundaries.

In this work, we systematically investigated the effects of chemical doping and microstructure modification on H solubility in monoclinic ZrO$_2$ doped with Fe, Cr or Ta. These dopants were selected following recent theoretical predictions of composition-dependent H solubility in this oxide by Youssef et al. [13]. Hydrogen solubility was measured via temperature-programmed desorption (TPD) for various microstructures and oxygen partial pressures, which allowed us to identify the origin of H defects in the oxides (i.e. bulk, surfaces and grain boundaries). To elucidate the type of H defects, solubility was measured under varied oxygen partial pressure, accentuating the role of oxygen in the H desorption process. Our measurements demonstrate that multiple H-containing defects and complexes are formed during bulk and surface hydration. We show that chemical doping effectively modulates H solubility in monoclinic ZrO$_2$, though we emphasize the role of microstructure control in controlling H sorption in certain compositions. Measurements of bulk H solubility presented here are in qualitative agreement with updated defect chemistry simulations based on DFT, which is the first experimental validation of the framework’s predictive capability and we reiterate its potential for accelerating the rational design of materials containing light-element point defects such as H and Li.

### 2. Methods

#### 2.1. Monoclinic ZrO$_2$ pellets were fabricated via solid state synthesis

A solid-state route was used to synthesize doped oxides pellets. ZrO$_2$ powder was mechanically mixed using a mortar and pestle with Cr$_2$O$_3$, Fe$_2$O$_3$, and Ta$_2$O$_5$ powders to yield nominally $D_xZr_{1-x}$O$_2$, where $D$ is Cr, Fe or Ta, and $x$ < 0.01.

Table 1. A binder of 5 wt% polyvinyl alcohol dispersed in distilled water was ground into the powder, which was pressed to pellets under uniaxial compression at 235 MPa in a steel piston/dye assembly. Pellets were sintered in dedicated alumina crucibles in air for 20 h with heating and cooling rates of 2.5 °C/min. Sintering temperatures varied from 1125 °C – 1700 °C, as indicated in Fig. 3, producing disc fragments with edge length of 5 mm – 1 cm, e.g. Fig. 4c and Figure S2.

Phase analysis of the sintered pellets was performed using x-ray diffraction (XRD) with Cu K$_\alpha$ radiation (Panalytical X'Pert Pro). Mean grain diameter was measured after each heat treatment from SEM images using the mean line intercept method from > 100 grains per sample.

The porosity of the pellets was also measured after each heat treatment. The total porosity, $P_{\text{Total}} = P_{\text{Closed}} + P_{\text{Open}}$, was determined from total density, $\rho_{\text{Total}}$, which was calculated from the total volume (total volume includes the solid and all porosity), and theoretical density, $\rho_{\text{Theory}}$, Eq. (1).

$$P_{\text{Total}} = 1 - \frac{\rho_{\text{Total}}}{\rho_{\text{Theory}}}$$  

The closed porosity, $P_{\text{Closed}}$, was determined from the so-called closed density, $\rho_{\text{Closed}}$, Calculated using the oxide volume measured via water displacement, Eq. (2). The closed density is calculated assuming that the measured volume includes only that of the solid oxide and any closed porosity.

$$P_{\text{Closed}} = 1 - \frac{\rho_{\text{Closed}}}{\rho_{\text{Theory}}}$$  

Finally, the open porosity was calculated as $P_{\text{Open}} = P_{\text{Total}} - P_{\text{Closed}}$.

#### 2.2. Defect chemistry was simulated using DFT and statistical thermodynamics

Equilibrium defect concentrations at various oxygen partial pressures and temperatures were predicted using a first-principles modeling framework. Formation energies of intrinsic defects and doping elements in monoclinic ZrO$_2$ are obtained from DFT calculations performed using the Vienna Ab initio Simulation Package (VASP) [28–30]. The generalized gradient approximation (GGA) with Perdew-Burke-Ernzerhof (PBE) functional is used [31,32]. We use a supercell size of $2 \times 2 \times 2$ ZrO$_2$ conventional cells (32 formula units) for the defect calculations; energy cut-off is set to 700 eV and a $2 \times 2 \times 2$ K-point grid is used. Additional details about the thermodynamic model and DFT defect calculations are provided elsewhere [13,33]. All possible interstitial sites, substitutional sites and valence states are considered for the dopant species. To compare with experiment, dopant concentrations were fixed at 1 atomic percent for all doped calculations in the defect equilibrium model.
2.3. Sintered oxides were hydrated by humid H₂ at elevated temperature

The oxides were hydrated at 410 °C – 760 °C by exposure to humidified 3% H₂:N₂ (Airgas), Fig. 4a. The sample temperature was measured with a K-type thermocouple positioned outside of the quartz tube which contained the sample and hydrating gas. The H₂: N₂ was bubbled at 20 sccm through 25 °C de-ionized water, producing a reducing gas with partial pressures of H₂O = 3.13 x 10⁻⁶ atm, H₂ = 2.91 x 10⁻² atm, and N₂ = 9.40 x 10⁻¹ atm. The pO₂ was 1 x 10⁻¹⁶ atm – 1 x 10⁻¹¹ atm at the hydration temperatures, as measured with a yttria-stabilized zirconia Nernst cell with porous Pt electrodes and one side of the cell exposed to atmosphere, Figure S1. To freeze in the equilibrium defects formed under each hydrating condition, samples were cooled to room temperature within 5 mins by removing the quartz tube from the tube furnace and fanning the tube with room-temperature air using a motorized fan.

2.4. Hydration times were determined by electrical conductivity relaxation

Electrical conductivity relaxation measurements were performed at each hydrating temperature by monitoring the current response to 20 mV applied while switching the sample environment between dry O₂ and humid H₂, e.g. Fig. 4c. The oxides were contacted with electrodes comprised of 100 nm of sputtered Pt covered with Pt ink fired at 1000 °C for 1 h. Measurements were performed using a digital source measure unit (Keithley 2400) controlled by LabVIEW software.

2.5. H₂ solubility was measured using temperature-programmed desorption

Following hydration, the H₂ uptake was quantified using temperature-programmed desorption (TPD). Hydrated samples were transferred to a second tube furnace flow reactor equipped with a mass spectrometer configured to quantify the effluent composition, Fig. 4b. This reactor is described in detail in chapter two of the doctoral dissertation of Zhao [34]. Samples were dehydrated by heating at 10 °C/min from 25 °C to 1075 °C in dry synthetic air (21% O₂:Ar; Airgas ultrahigh purity) flowing at 25 sccm. The sample was in a quartz tube inside a three-zone tube furnace, and the temperature was measured locally by a K-type thermocouple located inside the reactor adjacent to the sample. The concentrations of desorbed H₂O and H₂ in the effluent gas were monitored using a quadrupole mass spectrometer (Hiden Analytical Inc., HPR20) whose gas-probing filament was located adjacent to the sample on the gas exhaust side. To prevent water condensation, the gas-handling tubes were heated to ~130 °C. H₂ solubility reported in Fig. 6b, d, f, h, was calculated by integrating over the temperature ranges containing peaks in the steam release rate vs. temperature profiles in Fig. 6a, c, e, g. The solubility values reported in Fig. 6a-b were calculated similarly from the release rate data in Fig. 5. The integration windows are tabulated in Table S3.

Calibration gasses were used to convert the raw signal, R, in Eq. (3), to gas mole fraction, X, in Eq. (4). B₀ and η are the background signal and calibration factor for species i, respectively. The calibration factor was measured after each desorption measurement by flowing calibration gasses with identical flow rate and in concentrations comparable to those detected during experimental measurements. Calibration of H₂O was performed by bubbling the carrier gas through distilled water heated to 40 °C, and calibration of H₂ employed 5% H₂:Ar (Airgas).

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\text{Corrected signal of species } i = \frac{R_i - B_i}{\eta_i}
\]  

\[
X_i = \frac{\text{Corrected signal of species } i}{\sum_j \text{Corrected signal of species } j}
\]  

3. Results & discussion

3.1. Doped monoclinic ZrO₂ pellets were fabricated with modified microstructure

The sintered ZrO₂ was monoclinic (α-ZrO₂) as confirmed by XRD, Fig. 1. The diffraction patterns were unchanged following hydration-dehydration cycling, indicating that the sintered oxides were phase stable throughout the study. Fe- and Cr-doped ZrO₂ were both monoclinic without trace of the binary precursor oxides Fe₂O₃ or Cr₂O₃. Fe is stable in α-ZrO₂ up to 15 at% [19,35]. Ruh and Garrett also showed using XRD of vacuum-sintered Cr-doped ZrO₂ that (i) Cr addition did not stabilize the tetragonal phase, (ii) Cr was lost via vaporization during sintering, and (iii) the Cr solubility limit was ~1 at%–1.6 at% [36]. The Fe- and Cr-doped ZrO₂ were orange and blue, respectively, and the alumina crucibles in which the pellets were sintered were slightly discolored, indicating some Fe and Cr loss.

The only minor phase observed was in Ta-doped ZrO₂ (Ta₀.003Zr₀.997O₂), wherein a phase commonly attributed in the literature
to either tetragonal or orthorhombic zirconia (so-called \(\beta\)-ZrO\(_2\)) was stabilized. The Ta-doped ZrO\(_2\) XRD pattern obtained before hydration (Fig. 1) was analyzed using Rietveld refinement assuming the coexistence of Ta-containing monoclinic and tetragonal phases to estimate the \(\beta\)-ZrO\(_2\) phase fraction. This analysis indicated that the \(\beta\)-ZrO\(_2\) phase fraction was 0.5\% ± 0.2\%. This is consistent with the monoclinic phase fraction expected based on a systematic study of this oxide by Sponchia et al. [37], though they did not investigate compositions below 1\% Ta whereas we used 0.3\% Ta. Bhattacharya et al. calculated phase diagram isothermal sections indicating that Ta\(_2\)O\(_5\) in concentrations < 2 at\% is soluble in tetragonal ZrO\(_2\) above \(\sim 1250^\circ\)C [38]. If we take 2 at\% as the solubility limit of Ta in \(\beta\)-ZrO\(_2\)—whose phase fraction is 0.5\% in our case, then an estimated 0.01 at\% of Ta dopants are stabilized in \(\beta\)-ZrO\(_2\), leaving \(\sim 0.02\) at\% Ta to dope the monoclinic phase.

Microstructures were evaluated throughout the study by SEM imaging, Fig. 2. The as-prepared pellets were polycrystalline with grain size 0.21 \(\mu\)m − 1.63 \(\mu\)m, Fig. 3a. Their total porosity (the sum of closed and open porosity) was 0.32−0.42, and the open porosity (the porosity connected to the pellet surface) was 0.27−0.29, Fig. 3b. Closed density (derived from the combined volume of oxide and closed pores) was 85%–95% relative to theoretical density, indicative of a microstructure comprising regions of open porosity and a densified polycrystalline network of 85%–95% relative density. At higher sintering temperatures a decrease in closed porosity can occur if

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**Fig. 2.** Secondary electron SEM images of the as-prepared oxides with composition, sintering temperature and sintering time indicated.

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**Fig. 3.** Mean grain size (a) and open porosity (b) varied with increasing sintering temperature. Mean grain diameters are plotted with one standard deviation derived from measuring > 100 grains using the mean lineal intercept method. Open porosity is plotted with propagated uncertainty.
internal open pores join with the network of open porosity. This would lower the amount of closed porosity without affecting total porosity, thus increasing the measured open porosity. This is consistent with the dense polycrystalline microstructures shown in Fig. 2. The composition of the doped ZrO$_2$ was assumed to be uniform from grain to grain as compositional heterogeneity was not detected using SEM energy-dispersive x-ray spectroscopy or SEM backscatter imaging. The influence of grain size and open porosity on H sorption is described below.

4. H solubility was modified by doping and microstructure variation

4.1. Equilibration time under hydrating conditions was measured using electrical conductivity relaxation

To verify that the oxides reached chemical equilibrium during hydration (Fig. 4a), electrical conductivity relaxation (ECR) measurements were performed under each hydration condition, e.g. Fig. 4c. This technique detects changes in electrical conductivity during equilibration of the oxide with the hydration gas, humid 3% H$_2$:N$_2$ (see Methods). Ideally, a concentration gradient of the mobile defect drives mass transport in and out of the oxide, which occurs orthogonally to the measured electrical current flow. The equilibration (or hydration) time, therefore, is the time required for the electrical current to reach steady state after switching the gas from dry O$_2$ to humid H$_2$. Switching from dry O$_2$ to the hydrating gas changed the oxides’ electrical conductivity, as exemplified by ZrO$_2$ in Fig. 4c. The time required to transition between steady states was taken as the equilibration time for hydration. For ZrO$_2$ at 700 °C, the conductivity drop is attributed to a transition from p-type electronic conduction under dry O$_2$ (high pO$_2$) to oxygen ionic or n-type electronic conduction (low pO$_2$) in the hydration gas – whose pO$_2$ was 1 - 10$^{-16}$ atm - 1 - 10$^{-11}$ atm at the hydration temperatures. This would be consistent with experimental work by Kumar et al. [39] on the pO$_2$ dependence of electrical conductivity above 600 °C, and theoretical work of Youssef et al. [13] who computed the pO$_2$ dependent defect chemistry at 323 °C and 923 °C.

Hydration times decreased with increasing hydration temperature, attributed to slower diffusion kinetics at lower temperatures, Fig. 4d. Hydration times reported here for α-ZrO$_2$ are in good agreement with equilibration times reported for ECR experiments on α-ZrO$_2$, which range from 1 h – 4 h [39,40]. Repeated measurements of H solubility were also insensitive to hydration time above the equilibration times in Fig. 4d, which corroborated the ECR data. Interestingly, doping ZrO$_2$ increased hydration time at each temperature. One explanation for this was reported by Muta and Yamanaka, whose ab initio studies of monoclinic ZrO$_2$ showed that Cr and Fe dopants can be trap sites for diffusing H, thus lowering the mobility of H point defects [41]. Une et al. showed also that β-ZrO$_2$ had slower H diffusivity than α-ZrO$_2$, which may also contribute to the slower equilibration times of Ta-doped ZrO$_2$ relative to undoped ZrO$_2$ [42].

Reversible color change during hydration was observed in each oxide: ZrO$_2$ turned from white to light grey suggesting oxygen deficiency [23,43], Fe-doped ZrO$_2$ changed from orange to blue-grey (Fig. 4e), Cr-doped ZrO$_2$ turned from dark to light teal-blue (Figure S2), and Ta-doped ZrO$_2$ turned from white to very light yellow. Decomposition or mechanical failure of the oxides did not result from (de)hydration, consistent with previous experimental work demonstrating chemical stability of ZrO$_2$ in solution with pH > 1.5 [44].

Fig. 4. Requisite hydration times were measured using electrical conductivity relaxation. (a) Oxides were hydrated at elevated temperature by exposure to humidified H$_2$. (b) Oxides were dehydrated by heating in dry O$_2$ and the effluent composition was monitored by mass spectrometry. (c) An exemplary conductivity relaxation curve shows that ZrO$_2$ at 700 °C equilibrates in 0.4 hours; the decrease in conductivity is attributed to a transition from p-type to oxygen ionic or n-type electronic conduction. (d) Hydration times increased with decreasing hydration temperature and upon doping ZrO$_2$. (e) Hydrating the oxides caused reversible color change, exemplified by pellets fragments of Fe-doped ZrO$_2$. 
4.2. Hydrogen desorption was affected by the choice of dopant

Hydrogen desorption during dehydration was characterized using temperature-programmed desorption (TPD) measurements, where the hydrated oxides were heated in dry synthetic air (21% O2:Ar), Fig. 4b. Water desorption was detected from all samples, and in some cases H2 desorption was also detected. However, in those cases < 1% of the total hydrogen dissolved in the oxide was released as H2. Multiple desorption peaks were typically observed in each desorption curve, with the curve shape varying with oxide composition and hydration temperature. Sherman and Olander compared H release from single and polycrystalline UO2, and showed that H2 desorption from single crystals could be described quantitatively based on H chemical diffusivity, whereas desorption curves from polycrystalline samples exhibited multiple peaks and followed no simple diffusion mechanism [9]. In that and an earlier work [8] they reported desorption peaks with sharp onsets and very clear high-temperature tails/shoulders, somewhat similar to peak shapes presented here.

The shape of water desorption profiles from ZrO2 (ZrO2' in Fig. 5) was in very good agreement with data reported previously for monoclinic ZrO2 pellets by Yamanaka et al. [24]. The desorption curves exhibit a low-temperature peak at ~ 150 °C and high-temperature peak/tail at ~ 550 °C, and the relative intensity of the two peaks is also in good agreement. The low-temperature desorption was observed previously by Yamanaka et al. who attributed it to desorption of surface-adsorbed water [24]. Interestingly, those desorption measurements of sintered pellets of monoclinic ZrO2 were performed in a vacuum of 10^-7 Pa, whereas similar studies of single crystals of tetragonal ZrO2 by Park and Olander [25] did not detect low-temperature H desorption from samples measured in a vacuum furnace (no base pressure was reported). This suggests that surface defects present in polycrystalline ZrO2 plays a critical role in the adsorption of H-containing species. The high-temperature desorption increased with decreasing hydration temperature, which is again in fair agreement with the report of Yamanaka et al., as discussed below.

Kouva et al. [20] investigated water’s interaction with the surface of monoclinic ZrO2 particles (240 μm – 500 μm) using a combination of TPD and optical spectroscopy. Water desorption from their reduced/hydrated sample in N2 began around 150 °C and peaked around 250 °C – 300 °C, with a tailing decay in desorption to over 550 °C. In addition to TPD, the authors correlated sample temperature during desorption with diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) to characterize the surface-related H species during heating. When heating the reduced/hydrated ZrO2 in N2 they showed that the predominant form of chemisorbed H was in hydroxyl groups with O multi-coordinated with lattice Zr. The combination of their desorption measurement—which is in qualitative agreement with our observations of ZrO2—and their DRIFTS result assists in our attribution of low-temperature water desorption to a combination of (i) physisorbed water and (ii) chemisorbed hydroxyl groups which are multi-coordinated likely by filling O vacancies formed in the reducing environment present during hydration. Additionally, the coexistence of multiple ‘liquid-like’ and ‘ice-like’ surface-adsorbed waters on porous Y-stabilized tetragonal ZrO2 at low temperatures was established experimentally by Stub et al. [45].

For Cr-doped ZrO2 two distinct water desorption processes were observed, with the high-temperature desorption increasing with decreasing hydration temperature. The release rate from Fe-doped...
ZrO₂ increased modestly at low temperature and was sustained at that initial level until ∼ 450 °C, when desorption increased rapidly to its peak value before decaying to the baseline value at 1000 °C. Unlike the undoped case, the high-temperature desorption process decreased with decreasing hydration temperature. Ta-doped ZrO₂ performed similarly to undoped ZrO₂ in terms of water release rate, though its low-temperature desorption peaks were less intense. The subsequent sections of this paper elucidate the H-containing defects from which the desorbed H originate and quantify H solubility from desorption data in Fig. 5. The impact/role of each dopant on stabilizing sorbed H is examined and discussed throughout the subsequent sections.

4.3. Microstructure variations influenced H absorption

To assist interpretation of the desorption profiles in Fig. 5 we varied the oxides’ grain size and porosity via heat treatments, which impacts the concentration of H binding sites such as surfaces and grain boundaries without changing the number of bulk sites. Increasing the annealing temperature increased the mean grain size (measured from SEM images, Figure S3 – Figure S6) and decreased the open porosity, Fig. 3.

In ZrO₂ increasing the annealing temperature from 1125 °C to 1600 °C increased the grain size from 0.21 µm to 4.2 µm and decreased the open porosity from 0.27 to ∼ 0.1, Fig. 3. The water desorption profiles from samples annealed at higher temperatures, e. g. 1600 °C, contain two peaks, Fig. 6a. The low-temperature peak (Peak 1) generally decreased with annealing temperature, while the high higher-temperature peak (Peak 2) was insensitive to annealing temperature. In this sample Peak 1 is attributed to desorption of surface-adsorbed water (see also Fig. 7), suggesting that pore shrinkage may explain the decreased desorption with annealing. Because Peak 2 in this sample is insensitive to annealing temperature it is attributed to bulk defects, discussed below (Fig. 7). When varying sintering temperature, the hydration temperature was fixed at 760 °C.

For Fe-doped ZrO₂, increasing the annealing temperature resulted in grain growth from 1.63 µm to 11.3 µm, Fig. 3a. This grain growth was accompanied by a slight reduction of the closed density, indicating that closed porosity increased with sintering temperature. The open porosity was not measured, however, because reliable dry density measurements could not be performed; increasing the sintering temperature from 1450 °C to 1650 °C, and then to 1700 °C caused disintegration of the pellet fragments into small, ill-defined pieces < 3 mm whose dry volume could not be reliably determined.

Increasing the annealing temperature revealed three unique water desorption peaks in Fe-doped ZrO₂, whose relative intensities and positions varied considerably with sintering, Fig. 6c. Desorption in peak 1 (below 250 °C, colored blue) was found to decrease slightly with increasing sintering temperature, with H solubility dropping from 1.70 · 10⁻⁴ to 5.92 · 10⁻⁵ H/ZrO₂ attributed to desorption of physically absorbed water molecules, Fig. 6d. This was concluded based first on the expectation that open porosity decreased slightly via densification during sintering—which should reduce the surface area, and second on desorption measurements performed under O₂-poor conditions, Fig. 7.

Water desorption in peak 2 showed no obvious trend with annealing temperature, and H solubility was constant at ∼ 3 · 10⁻⁴ H/ZrO₂. Unlike peak 1, desorption in peak 2 did consume O₂ under O₂-poor conditions (Fig. 7), suggesting that the H-containing desorbing species may be surface hydroxyls or protons. It is unclear whether these are bound to the oxide surface or absorbed at grain boundaries. Though, considering the insensitivity of peak 2 to changes in grain boundary area, it may be reasonable to assume these species are not related to grain boundaries. Peak 2 also appears to shift to higher desorption temperatures, which may indicate that the desorbing species adhere more strongly to the oxide surface which has experienced higher sintering temperatures.

Water desorption in peak 3 was the most sensitive to sintering, with H solubility decreasing from 1.14 · 10⁻³ to 8.16 · 10⁻⁴ H/ZrO₂. This approximately 14-fold decrease in peak 3 H desorption corresponded to an increase in grain diameter from ∼ 2 µm to ∼ 12 µm. This peak is tentatively attributed to the desorption of H absorbed at grain boundaries. The ratio of grain boundary area (i.e. the area of a grain’s surface) to grain volume has an inverse dependence on grain diameter, so an increase in the grain diameter by six times should correspond to an equal drop in the grain boundary area. Assuming the grain boundary H coverage does not change during sintering, the decrease of 14 times is a factor of two greater than expected, so we suspect that the grain boundary areal coverage decreases during sintering. It is likely that the local defect concentrations at grain boundaries following heating to 1450 °C varies from the defect concentrations following sintering at 1650 °C. It is well known that solutes and impurities accumulate at oxide grain boundaries which may serve to decrease the H coverage, in this case by a factor of about two. It was shown experimentally that Fe³⁺ segregation as Fe₂O₃ to the surface of Fe-doped ZrO₂ lowered the oxide’s catalytic activity by reducing the capacity of the surface to accommodate OH- groups [46]. In our case, accumulated Fe₂O₃ may lower the H capacity of grain boundaries in a similar manner. The Fe-doped sample crumbled into ∼ 3 mm fragments after repeated heat treatments. This was observed previously by Beck and Kaliba, who state that Fe is incorporated into ZrO₂ in the +3 oxidation state, but at high temperature changes to +2 leading to more oxygen vacancies in the tetragonal lattice and thus greater lattice parameter mismatch with the monoclinic phase which exacerbates stress [35].

For Cr-doped ZrO₂, increasing the annealing temperature from 1450 °C to 1675 °C yielded modest grain growth and open pore shrinkage, from 1.24 µm to 3.11 µm and from 0.29 to 0.22, respectively, Fig. 3. This relatively small influence of annealing temperature on grain size compared to the Fe-doped case may be related to Cr loss from the solid solution during annealing. Following each heat treatment, the color of the aluminia sample crucible was changed to scarlet, with the intensity of scarlet coloration increasing with annealing temperature. Rub and Garrett showed using XRD of vacuum-sintered Cr-doped ZrO₂ that Cr was lost via vaporization during sintering [36]. Here, since grain growth was minimal, it seems that the high mobility of the Cr ions served primarily to deplete the oxide of Cr via diffusion into the aluminia crucible rather than to promote sintering.

The water desorption profiles from Cr-doped ZrO₂ had two distinct peaks: Peak 1 from 40 °C – 350 °C, and Peak 2 from 350 °C – 1000 °C, Fig. 6e. The amount of H desorption in low-temperature Peak 1 decreased with increasing annealing temperature (Fig. 6f), likely reflecting the drop in the surface area associated with open porosity. Like previous cases Peak 1 was attributed to desorption of water physisorbed to surfaces based both on the temperature range and O₂-poor desorption measurements presented below, Fig. 7. The amount of H in Peak 2 did not vary significantly with annealing temperature, suggesting that these H may originate from bulk defects rather than defects associated with grain boundaries. According to our simulations of formation free energies detailed in [13] and presented here, we believe H stabilizes as interstitial defects in Cr-doped ZrO₂ at the hydration temperatures use here. This is consistent with the experimental observation that the desorbing H originates from bulk defect in this oxide. From O₂-poor measurements, Peak 2 desorption was accompanied by a very aggressive O₂ consumption, indicative of desorption of H in the form of hydroxyls or protons; the latter interpretation being consistent with our belief that H exists as interstitials.

For Ta-doped ZrO₂, the surface desorption peak (Peak 1) unexpectedly increased despite the oxide sintering during serial annealing. Accumulation of Ta on the surface of Ta-doped ZrO₂ was shown to increase the catalytic activity for 2-propanol dehydration [47], so perhaps segregation of Ta as Ta₂O₅ during sintering increases surface
Fig. 6. Desorption measurements were performed as a function of annealing/sintering temperature, which affected the oxides’ porosity and grain size. (a,b) ZrO$_2$, (c,d) Fe-doped ZrO$_2$, (e,f) Cr-doped ZrO$_2$, (g,h) Ta-doped ZrO$_2$. (a,c,e,g) The steam release rate ($H_2O$ per unit cell of $ZrO_2$ per second) from each $ZrO_2$-based material after sequential heat treatments at various sintering temperatures. Samples were hydrated at 760 °C and water release rates were measured by temperature-programmed desorption. Multiple water desorption peaks (numbered 1, 2 or 3) were observed during TPD measurements. (b,d,f,h) $H$ solubility associated with each of the desorption peaks was quantified as a function of sintering temperature. The profiles are offset to ease comparison of the profile shapes.
reactivity with OH− groups, promoting hydration/hydroxylation. This could explain why surface desorption increased with annealing temperature. Perhaps Ta surface segregation during sintering gives rise to more surface adsorption via hydride or hydroxide formation, though further study is required to be conclusive. Like the other oxides, Peak 2 desorption was basically insensitive to annealing which indicates that it may be attributable to bulk H defects.

5. Desorbed H species were investigated via oxygen-poor TPD

To elucidate the type of H-containing species desorbing during TPD, measurements were performed using an inert carrier gas (Ar or N2), rather than synthetic air, Fig. 7. Use of inert carrier gas significantly lowered the pO2 during desorption, which accentuated the effect of O2 on desorption processes. A detectable amount of O2 was present in the reactor presumably due to imperfect sealing.) In all cases H2 desorption was negligible. As above, during heating of ZrO2 two primary water desorption processes occurred both without O2 consumption. It has been established above that the low-temperature desorption results from desorption of molecular water from surfaces, but it is remarkable that no O2 is consumed during high-temperature water desorption. One possibility is that during hydroxyl desorption from hydrated ZrO2, the oxide becomes oxygen-deficient (via reduction of the Zr cation) and lattice O is consumed to produce water. Surprisingly, O2 is apparently not consumed to refill anion vacancies in the ZrO2 lattice, which is known to occur during interaction of gas-phase O2 with the oxide at high temperature.

For Fe-doped ZrO2 desorption below 250 °C occurred without O2 consumption, likely due to the desorption of physisorbed water. From 250 °C–625 °C water desorption occurs in conjunction with a decrease in the O2 concentration, indicating that the desorbed H species are hydroxyls and/or protons. Interestingly, high-temperature water desorption (whose intensity decreases with increasing grain diameter) does not consume O2, indicating that H desorbing from grain boundaries may be consuming lattice O to form water leaving behind O vacancies.

In the Cr-doped case, water desorption below ~200 °C occurred without O2 consumption, consistent with desorption of physisorbed water. As the temperature increased to ~500 °C, the O2 signal drops gradually while the water signal remains above the baseline. This indicates desorption of H from hydroxyl or proton defects believed to be associated with microstructural features such as internal surfaces and/or interfaces; hydroxyls or protons bonded to the material’s internal surfaces and/or interfaces desorb and contribute to the detected steam via consumption of O2, which drops gradually in the temperature range 200–500 °C. Water desorption in this temperature range was shown to be sintering temperature dependent, Fig. 6f. At ~550 °C, there is the expected intense water desorption during which the O2 is zero, again indicating desorption of hydroxyl/protonic defects. These defects are suspected to be bulk defects since this desorption event was found to be unaffected by microstructural changes induced by annealing, Fig. 6f.

For Ta-doped ZrO2, the O2-poor desorption measurements looked much like the O2-rich measurements (Fig. 6g), and there was no indication of increased O2 consumption during water desorption. Like ZrO2, this may simply reflect the low concentration of H defects in this oxide and thus the small amount of desorbing water, whose requisite O could be derived from non-stoichiometry created in the oxide lattice rather than the surrounding O2 gas.

6. Solubility was measured from desorption profiles and compared with theory

With a better understanding of the origin of H defects producing the measured water desorption profiles, it was possible to determine heats of solution of H associated with surface and bulk desorption. By integrating the water desorption curves in Fig. 5 over the appropriate temperature ranges (determined from annealing-temperature-dependent desorption measurements, Table S3), we quantified H desorption attributed to (low-temperature) surface adsorption, and H desorbed at higher temperatures attributed to bulk and grain boundary (in Fe-doped ZrO2) defects, Fig. 8. Surface-desorbed H (Fig. 8a) and high-temperature-desorbed H (Fig. 8b) were taken as unique measures of H solubility, and were fit to Eq. (5) to determine heats of solution for these H defects in the oxides [9], Table 2. Eq. (5) describes the Arrhenius relationship between H solubility, S, and the heat of solution, $\Delta H_{\text{solution}}$, over a range of hydration temperatures, $T_{\text{hydration}}$. A is a pre-exponential factor. The measured H solubility derived from low- and high-temperature steam desorption data (Fig. 8a-b) were plotted in the Arrhenius fashion and Eq. (5) was fit to the data using a least-squares fitting routine implemented in Python. Surface desorption increased with hydration temperature for all samples and was more endothermic when the ZrO2 was doped, Fig. 8a and Figure S7. The low-temperature desorption of physisorbed water was reduced in all cases by doping (aside from Fe-doped ZrO2 hydrated at 760 °C), and the lower heats of solution indicate that dopants influence surface chemical properties of the oxide via surface segregation during sintering. Perhaps O vacancies that serve as hydroxyl binding sites may be charge-neutralized by a segregated cation making the O site inactive for H absorption.

$$S = A \cdot \exp \left( \frac{\Delta H_{\text{solution}}}{RT_{\text{hydration}}} \right)$$

The H desorbed at higher temperature was attributed to bulk and grain boundary defects, Fig. 8b and Figure S7. Like the surface, bulk and grain boundary H defects were more endothermic when the ZrO2 was doped, Table 2. Bulk H solubility in undoped ZrO2 increased with decreasing hydration temperature, which is in fair quantitative agreement with Yamanaka et al. [24] who showed that H absorption...
in ZrO₂ is exothermic. Relative to the undoped case, Cr doping increased bulk H solubility by several times. Fe-doped ZrO₂ was the only material with grain boundary solubility, which decreased rapidly with decreasing hydration temperature. Sundell et al. used atom probe tomography to show that Fe and Ni decorate grain boundaries in oxide scale on corroded Zr alloys [27], and that OH⁺/OD⁺ and Fe²⁺ also accumulated at grain boundaries in these oxides [11,12]. Based on the present work, it appears that Fe doping yields the most tightly bound H defects in the bulk and at grain boundaries, which could improve the H resistance of a coating via kinetically limiting H diffusion. Furthermore, it seems that lower-temperature applications demanding low H solubility may benefit from Fe doping, whereas the potential for H solubility minimization appears to be most significant. In Ta-doped ZrO₂ the bulk H solubility was lower than the undoped case, and solubility decreased with decreasing temperature making this dopant another candidate for minimizing H solubility at low temperatures. Moreover, the absence of grain boundary H absorption may be useful when considering future microstructural optimization.

Excitingly, bulk H solubility measurements are in qualitative agreement with our theoretical predictions in several notable cases, Fig. 8c. Using a DFT-based modeling framework developed and demonstrated recently by our group [13–15], we calculated the H point defect concentration under simulated hydration conditions closely replicating those used in our experiments, see Methods and Supporting Information. There is a considerable discrepancy between the absolute values of measured and simulated solubility, perhaps suggesting the presence of additional bulk H point defects in the materials which were not in the defect chemical computation. We provide additional details discussing possible origins of these discrepancies in the Supporting Information. Our simulations indicate that bulk H solubility increases upon doping, with the trend in isothermal solubility following: Undoped < Fe-doped < Ta-doped < Cr-doped ZrO₂. Ignoring the Ta-doped case this trend is nearly identical to that observed in our experimental measurements, although ZrO₂ did absorb more H than Fe-doped ZrO₂ hydrated at 410 °C. While further investigation is warranted to theoretically elucidate the role of Ta doping, this

**Table 2**
Heats of solution for surface-desorbed H (low-temperature peak, below ~350 °C) and H desorbed at higher temperature (high-temperature peaks, above ~350 °C). *Data from [24] was fit using Eq. (5).

<table>
<thead>
<tr>
<th></th>
<th>α ZrO₂*</th>
<th>α ZrO₂ (This work)</th>
<th>Cr-doped α ZrO₂</th>
<th>Fe-doped α ZrO₂</th>
<th>Ta-doped α/β ZrO₂</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heat of solution, surface desorbed H (kJ mol⁻¹)</td>
<td>1.3</td>
<td>19</td>
<td>52</td>
<td>3.6</td>
<td></td>
</tr>
<tr>
<td>Heat of solution, Bulk desorbed H (kJ mol⁻¹)</td>
<td>-29</td>
<td>-7.7</td>
<td>5.9</td>
<td>17</td>
<td></td>
</tr>
<tr>
<td>Heat of solution, Grain boundary desorbed H (kJ mol⁻¹)</td>
<td>35</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

![Fig. 8. Temperature programmed desorption was used to measure H desorbed from undoped and doped zirconia.](image-url)
result is encouraging and highlights the predictive capability of this defect chemical modeling framework.

7. Conclusions

We performed novel systematic measurements of the effect of chemical doping and microstructure on H solubility in monoclinic ZrO$_2$ ceramics. Temperature-programmed desorption was used to quantify H absorption, which was measured systematically for various microstructures and under varied oxygen partial pressure conditions. This helped identify the microstructural origin of the H defects (i.e., bulk, surfaces and grain boundaries), and infer the defect type. Upon doping there was a significant increase in measured hydration times, demonstrating that solute cations slowed the diffusion of H defects in the oxides. Doping with Ta offers the lowest H solubility at all hydration temperatures (410 °C–760 °C), though Fe-doping also reduces H solubility at lower temperatures and results in the strongest H binding of all oxides, which could improve H resistance based on diffusion kinetics. While it is clear from this work that doping significantly impacts H solubility in monoclinic ZrO$_2$, we emphasize the role of microstructural features in controlling H sorption in these ceramics. Excitingly, the impact of chemical doping by Fe, Cr and Ta substitutional cations on bulk H solubility was found to be in qualitative agreement with our theoretical predictions made using a thermodynamic modeling framework based on DFT. This is the first experimental validation of the framework, which offers high-throughput defect chemistry simulations of metal oxides with light element point defects (e.g. H in Al$_2$O$_3$ [15]), such as advanced energy materials.

Declaration of Competing interests

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

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Supplementary materials


References


[3] H. Iwahara, T. Esaka, H. Uchida, N. Maeda, Proton conduction in sintered oxides of all oxides, which could improve H resistance based on diffusion kinetics in Al$_2$O$_3$ [15]), such as advanced energy materials.

[4] D.R. Olander, D. Sherman, M. Balooch, Retention and release of water by sintered microstructures and under varied oxygen partial pressure conditions. This helped identify the microstructural origin of the H defects (i.e., bulk, surfaces and grain boundaries), and infer the defect type. Upon doping there was a significant increase in measured hydration times, demonstrating that solute cations slowed the diffusion of H defects in the oxides. Doping with Ta offers the lowest H solubility at all hydration temperatures (410 °C–760 °C), though Fe-doping also reduces H solubility at lower temperatures and results in the strongest H binding of all oxides, which could improve H resistance based on diffusion kinetics. While it is clear from this work that doping significantly impacts H solubility in monoclinic ZrO$_2$, we emphasize the role of microstructural features in controlling H sorption in these ceramics. Excitingly, the impact of chemical doping by Fe, Cr and Ta substitutional cations on bulk H solubility was found to be in qualitative agreement with our theoretical predictions made using a thermodynamic modeling framework based on DFT. This is the first experimental validation of the framework, which offers high-throughput defect chemistry simulations of metal oxides with light element point defects (e.g. H in Al$_2$O$_3$ [15]), such as advanced energy materials.

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