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Surface chemistry of pure tetragonal ZrO₂ and gas-phase dependence of the tetragonal-to-monoclinic ZrO₂ transformation†

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The surface chemical properties of undoped tetragonal ZrO₂ and the gas-phase dependence of the tetragonal-to-monoclinic transformation are studied using a tetragonal ZrO₂ polymorph synthesized via a sol-gel method from an alkoxide precursor. The obtained phase-pure tetragonal ZrO₂ is defective and strongly hydroxylated with pronounced Lewis acidic and Brønsted basic surface sites. Combined *in situ* FT-infrared and electrochemical impedance measurements reveal effective blocking of coordinatively unsaturated sites by both CO and CO₂, as well as low conductivity. The transformation into monoclinic ZrO₂ is suppressed up to temperatures of ~723 K independent of the gas phase composition, in contrast to at higher temperatures. In inert atmospheres, the persisting structural defectivity leads to a high stability of tetragonal ZrO₂, even after a heating-cooling cycle up to 1273 K. Treatments in CO₂ and H₂ increase the amount of monoclinic ZrO₂ upon cooling (>85 wt%) and the associated formation of either Zr-surface-(oxy-)carbide or dissolved hydrogen. The transformation is strongly affected by the sintering/pressing history of the sample, due to significant agglomeration of small crystals on the surface of sintered pellets. Two factors dominate the properties of tetragonal ZrO₂: defect chemistry and hydroxylation degree. In particular, moist conditions promote the phase transformation, although at significantly higher temperatures as previously reported for doped tetragonal ZrO₂.

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1. Introduction

The physical chemistry of different ZrO₂ polymorphs has been an intensively studied research field over decades, as recent reviews prove.^{1–9} However, upon literature screening, open questions especially regarding phase stability, surface termin-

ation, surface activity, or gas-phase-dependent stability limits still remain. Despite this deficiency, the diversity of the application of ZrO₂-based materials is considerable and directly reflects its manifold specific properties. Applications in ceramic engineering,¹⁰ as biomedical implants,¹¹ in sensor technology,¹² as microelectronic devices,¹³ as solid electrolytes¹⁴ and in catalysis² have been reported. Concerning the latter, ZrO₂ is used as both catalyst material and catalyst support. It is used for conversion of hydrocarbons in biomass-related reactions⁶ or doped with Y₂O₃ as an anode support and electrolyte material in fuel cells. As such, it encompasses use in catalytic processes over a wide temperature range between few hundred K up to 1273 K.¹⁵ Obviously, the respective requirements of the intrinsic properties of the used ZrO₂ material are strongly depending on the specific field of application. Many empirical and fundamental research attempts have therefore been made over the last decades to resolve the nature of the active sites on these ZrO₂-based materials alongside their selective generation and exact control with suitable reducibility and stability. Parameters that strongly influence the final activity are preparation routine, pre-treatment, phase

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(purity), termination, surface area, particle size, doping level or stoichiometry. Furthermore, when used as catalytic support material, in combination with metals like Au, Pt, Cu (methanol synthesis, water gas shift reactions) or Pt, Ni, Co, Cu (reforming reactions), also metal–oxide interfacial zones with additional multifaceted properties have to be taken into account.¹⁶ To add to the complexity, the Zr–O phase diagram exhibits five stable ZrO₂ polymorphs. However, the most commonly used and best characterized are the phases with the monoclinic, the tetragonal and the cubic structure. Due to the fact that the monoclinic phase (baddeleyite) is the by far most stable polymorph under realistic experimental conditions, this phase is most widely studied among pure undoped ZrO₂ materials. To access the other undoped polymorphs, specific annealing treatments are necessary. The temperature-induced phase transitions from monoclinic ZrO₂ to tetragonal ZrO₂ occurs at ~1400 K, and above ~2600 K the cubic phase is present.¹⁷ At lower temperatures the cubic and tetragonal polymorphs are usually stabilized by doping with different metal cations (e.g. Y³⁺, Si⁴⁺ and Zn²⁺). However, not only the structure, but directly also other intrinsic properties such as the amount and acidity/basicity of hydroxylation, stability of surface area or electrochemical conductivity are drastically changed. Apart from cationic doping, the tetragonal phase can also be stabilized by adding organic additives, but also by simple grain size control and additionally by interfacial effects (below a critical diameter d of $d \leq 33$ nm).⁸ Furthermore, also anions such as OH[−], SO₄^{2−} are potential stabilizers for low-temperature stable tetragonal ZrO₂.¹⁷ Sulfated tetragonal ZrO₂ is a special case in terms of catalytic surface modification, since it is a solid superacid, e.g. it exhibits a unique activity in the catalytic isomerization of hydrocarbons. Aside from the sulfated surface termination, the presence of especially a crystalline tetragonal phase is usually reported to be crucial for the performance of the ZrO₂-containing catalysts.¹⁸ Having said that, the stability of undoped and unsulfated pure tetragonal ZrO₂ (in terms of thermally or gas-phase-induced transformation into monoclinic ZrO₂) is still an interesting research field, since there are many preparation and pre-treatment attempts leading to different stability, but also activity and surface reactivity.

In the present study, we aim to characterize the surface chemistry of a pure undoped tetragonal ZrO₂ sample, prepared *via* a very simple sol–gel route starting from a solution of zirconium(IV)isopropoxide in isopropanol without any further additives. Calcination of the resulting Zr hydroxide aerogel leads to essentially phase pure nanocrystalline tetragonal ZrO₂ (which is most likely stabilized *via* OH[−]/OH-groups, lattice anionic defects and crystallite size effect).¹⁹ Furthermore, the focus is on elucidating the gas-phase-dependent physical chemistry of this tetragonal ZrO₂ phase under oxidative, reductive or inert gas atmospheres of dry and moist gases He, O₂, CO, CO₂ and H₂. The deliberate addition of H₂O is also worthwhile since phase transformation of tetragonal to monoclinic ZrO₂ in H₂O at room temperature has been reported.²⁰ This includes “low temperature” studies up to 723 K to characterize the surface

termination (*i.e.* hydroxylation, acid–base properties) and the corresponding electrochemical behavior of phase pure metastable t-ZrO₂. Additionally, “high temperature” studies, to gain detailed information about the gas-phase dependent stability, up to a temperature of 1273 K were performed. For this purpose *in situ/operando* Fourier-Transform infrared spectroscopy, *in situ/operando* electrochemical impedance spectroscopy, *ex situ* X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS) and atomic force microscopy (AFM) studies were conducted. From a previous study,¹⁹ detailed information about the general kinetic stability of the synthesized tetragonal ZrO₂ material in air has already been compiled. In fact, the aerogel precursor crystallizes at 673 K to essentially chemically and structurally phase-pure tetragonal ZrO₂. Transformation to monoclinic ZrO₂ in air is pronounced only upon annealing to 873 K and prolonged isothermal treatments. Less conversion takes place at isothermal treatment at higher temperatures up to 1273 K, but the main part of the sample is transformed to monoclinic (m-) ZrO₂ upon re-cooling to room temperature, once the critical temperature of 873 K is overcome.

2. Experimental

2.1. Materials

Pure tetragonal ZrO₂ was prepared *via* a sol–gel route starting with zirconium(IV)isopropoxide (isopropanol adduct, 99.9% Zr, Strem Chemicals) dissolved in isopropanol (Finne Gatt-Koller, 99.98%) and finally quenched with water. This preparation routine has been described previously in ref. 19 and is essentially a variant of the one reported by Mazdiyasn *et al.*²¹ (decomposition of zirconium alkoxides and the subsequent hydrolysis of zirconium hydroxy aerogels). The resulting aerogel was subsequently calcined at 723 K in pure O₂ in the respective experimental setups. All samples were without exception checked by *ex situ* XRD. The surface of the calcined (at a maximum temperature of $T = 723$ K) tetragonal ZrO₂ was 50 m² g^{−1} (according to the Brunauer–Emmett–Teller (BET) method, performed using a Quantachrome Nova 2000 Surface Area and Pore Size Analyzer). To ensure chemical purity, chemical analysis of all samples was performed using a Spectro XEPOS energy dispersive X-ray fluorescence analyzer (EDXRFA) which proves purities of 99.99%. The gases were supplied by Messer (CO 4.7, CH₄ 4.5, O₂ 5.0, He 5.0, CO₂ 4.5, H₂ 5.0). For a typical experiment in the *in situ/operando* FT-IR/EIS setups the sample was annealed in dry O₂ prior to each experiment, to ensure reproducible starting conditions (heating and cooling rates 10 K min^{−1}, gas flow ~1.0 mL s^{−1}). For absolutely dry conditions, the gases were dried with a liquid N₂-ethanol cooling trap at a temperature of approximately 163 K for CO, CH₄, and O₂, 223 K for CO₂ and for He and H₂ a N₂ cooling trap at 77 K was used.

As a structural reference commercial pure monoclinic ZrO₂ (zirconium(IV) oxide, 99.99%, Alfa Aesar) subjected to the identical chemical structural and spectroscopic analysis was used (see XRD pattern in Fig. S1†).



2.2. FT-IR spectroscopy

FT-IR spectra were recorded in transmission mode on an Agilent Cary 660 spectrometer with a mid-infrared source and a DTGS detector. Transmission mode was chosen to monitor both, the surface and bulk-related infrared-detectable phenomena. The powder samples were pressed into thin pellets using a weight of ≤ 2 t (10 mm diameter, thickness ~ 0.1 mm, sample mass about 20 mg) and then subsequently placed inside a homemade *in situ/operando* reactor cell,²² providing chemically inert surroundings of the sample in the heated area. *In situ/operando* measurements up to 1273 K under flowing and static conditions can be performed, in addition to vacuum experiments at minimum pressures of 3×10^{-7} mbar. BaF₂ as window material allows to access wavelengths above 800 cm⁻¹. Experiments in flowing mode (ambient pressure) can then be directly correlated with electrochemical impedance measurements. In static mode, the gases are pre-adsorbed on a 5 Å zeolite trap binding water sufficiently strongly, before the dry gases are desorbed into the evacuated and degassed cell. All reported spectra are corrected by the spectrum of the dry pre-oxidized oxide pellet at room temperature and under vacuum prior to exposure to the gases.

2.3. Electrochemical impedance spectroscopy (EIS)

The *in situ/operando* impedance cell consists of an arrangement of an outer quartz tube and two smaller inner quartz tubes, to which the sample and the electrodes are attached to. For heating, a tubular Linn furnace was used. Temperature control was ensured by a thermocouple (K-element), located in the reactor about 5 mm downstream of the sample, and a Micromega PID control unit. The impedance was measured using an IM6e impedance spectrometer (Zahner Messsysteme), providing data on the impedance and the phase angle of the current as a function of voltage simultaneously. The powder samples were pressed into pellets with a weight of ≤ 2 t (5 mm diameter, ~ 0.1 mm thick, sample mass about 20 mg) and placed between two circular Pt electrodes acting as a plate capacitor in mechanically enforced contact with the pelletized sample.

For all temperature-programmed impedance measurements described in this work, an amplitude of 20 mV of the superimposed sinusoidal modulation voltage signal at an overall DC potential of 0 V and a frequency of 1 Hz were applied to the Pt electrodes, *i.e.* the impedance of the pellet was effectively measured in an electrochemically unpolarized state. The impedance modulus value $|Z|$ represents the “impedance” in all temperature-programmed experiments.

For moist conditions, water vapor of 24 mbar was produced by passing He (~ 1 mL s⁻¹) through a water saturator at room temperature.

Note that the used spectrometer has a detection limit near $\sim 10^{10}$ Ω. In order to ensure an appropriate discussion, this phenomenon (plateau-like impedance stagnation near the detection limit) will be referred to as an “insulating” behavior in this work.

2.4. X-ray photoelectron spectroscopy (XPS)

In order to gain information about the surface-near chemistry, all samples were analyzed by X-ray photoelectron spectroscopy (XPS) using a Thermo Scientific MultiLab 2000 spectrometer with a base pressure in the low 10^{-10} mbar range. The instrument is equipped with a monochromated Al-K_α X-ray source (1486.6 eV), an Alpha 110 hemispherical sector analyzer. A flood gun is used for charge compensation, providing electrons with a kinetic energy of 6 eV. The energy axis shift was calibrated relative to the C–C component of the C 1s peak (set to 284.8 eV). For the analysis of the chemical states, the peak shapes used to fit the respective Zr 3d and C 1s spectra were products of Gaussian and Lorentzian peak shapes (30% Lorentzian). Several constraints were applied for the fitting procedure of the zirconium region: the splitting between the 3d_{5/2} and the 3d_{3/2} components was fixed at 2.4 eV, according to literature,²³ and the peak widths were set to be the same for the spin-split components as well as for the peaks of the different oxidation states. Furthermore, to enable convergence of the fit towards a sensible minimum in the presence of multiple oxidation states, the energy shifts of the 3d_{5/2} peaks between the various chemical states have been constrained to literature values.²⁴

2.5. X-ray powder diffraction (XRD)

The X-ray powder diffraction patterns were obtained by measuring the powder patterns in transmission geometry using a STOE STADI P powder diffractometer with Ge(111)-monochromatized MoK_{α1} ($\lambda = 70.93$ pm) radiation. The data were recorded in 2θ with steps of 0.1° from 2–45.875°. As a detector, a silicon microstrip solid state detector Mythen 1K was used. The structural refinement was performed with the program TOPAS 4.2²⁵ by taking the crystal structure parameters of monoclinic²⁶ and the tetragonal ZrO₂²⁷ as starting values. The obtained crystallographic data from the associated Rietveld refinement²⁸ are listed in Table S1.†

The *in situ* high-temperatures synchrotron XRD experiments in H₂ and CO₂ flow are performed at beamline 12.2.2, Advanced Light Source, Lawrence Berkeley National Labs, California. The *in situ* diffraction patterns were collected in the angle-dispersive transmission mode with a focused 25 keV monochromatic beam ($\lambda = 0.4959$ Å/50 μm spot size). The sample powder was heated in 700 μm quartz capillaries under a continuous gas flow (CO₂ or H₂) injected through another 300 μm capillary with a cut-open end. The capillary is heated in an infrared heated SiC tube furnace we described in ref. 29. Diffraction patterns are recorded by a PerkinElmer flat panel detector (XRD 1621, dark image and strain correction) every 25 seconds during the heating and cooling cycle. As the synthesized starting material is a powdery amorphous aerogel, previous to the experiment it is *in situ* sintered at 723 K inside the capillary under air flow until crystallization to tetragonal ZrO₂ is completed.

2.6. Atomic force microscopy (AFM)

Atomic force micrographs were obtained using a Veeco Dimension 3100 microscope in tapping mode, equipped with



HI-RES cantilevers by MikroMasch (with a tip radius below 1 nm) used for high-resolution images, as well as Veeco RTESPW silicon cantilevers (about 10 nm tip radius) to obtain information on larger scales.

3. Results and discussion

3.1. Surface chemistry of tetragonal ZrO₂ between room temperature (RT) and 723 K

3.1.1. Surface acidity and basicity. The acid–base properties of an oxide are of fundamental importance when evaluating its general catalytic potential.³⁰ It is quite common to determine the acidity/basicity of Lewis sites *via* the adsorption of the probe molecules CO (Fig. 1–3) and CO₂ (Fig. 4). While

CO is used to characterize the Lewis acidity, using CO₂ mainly basic sites can be evaluated.^{1,18,30,31} ZrO₂ can provide both, acidic and basic centers, but the quality and quantity of these active sites are strongly dependent on phase or crystal facet and (pre-)treatment (including parameters such as temperature, pressure or gas phase composition).^{16,18,30,32} Concerning tetragonal ZrO₂, detailed studies about CO and CO₂ adsorption have been already conducted – theoretical calculations^{16,30} as well as experimental infrared studies.^{18,32–37} However, as it has been already discussed for monoclinic ZrO₂, adsorption studies strongly vary between different works, since only slight differences in the preparation routine and (pre-)treatment of the samples lead to essentially different surface activities.^{1,31} In addition, for tetragonal ZrO₂ the situation is further complicated by the fact that usually “tetragonal” ZrO₂ in fact is deliberately doped by different cationic or anionic species, significantly influencing its physico-chemical (surface) properties. The present study now provides adsorption data on an undoped, phase-pure, metastable tetragonal ZrO₂ that is highly hydroxylated and defective, and most importantly, is stable and phase-pure up to temperatures of several hundred degrees ($T < 773$ K).¹⁹ What can principally be expected as active binding sites are coordinatively unsaturated (*cus*) Zr⁴⁺ and O^{2–} sites, hydroxyl groups and defects (oxygen vacancies).¹⁶

CO adsorption. The used pure tetragonal ZrO₂ was pretreated in dry O₂ up to 723 K and the spectrum of the fully oxidized sample at room temperature in vacuum was taken as background for all further spectra. The exposure of the oxide to an increasing pressure of CO leads to linearly adsorbed CO molecules, as displayed in Fig. 1.

In addition to the gas phase signal of CO between ~2240 and 2020 cm^{–1} a peak at 2191 cm^{–1} is present, which can be assigned to CO molecules bonded to Lewis acid Zr⁴⁺ centers, most likely on a step, kink or corner due to the high shift in

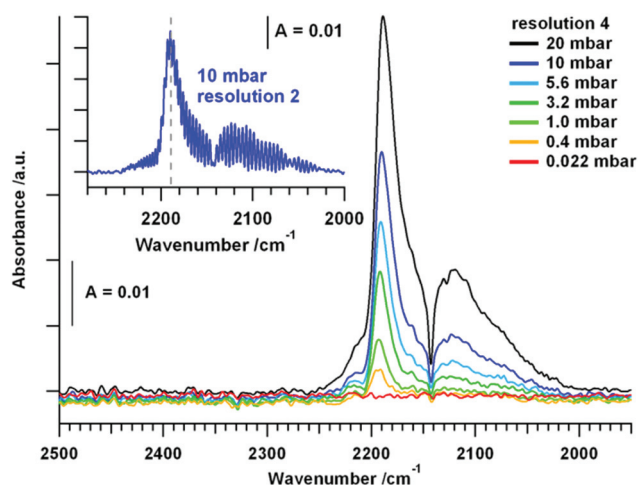


Fig. 1 FT-IR measurements of CO adsorbed on tetragonal ZrO₂ at CO pressures between 0.022 mbar and 20 mbar at room temperature. Spectra in the main panel were collected using a resolution of 4 and the inset shows a spectrum of 10 mbar CO with a resolution of 2.

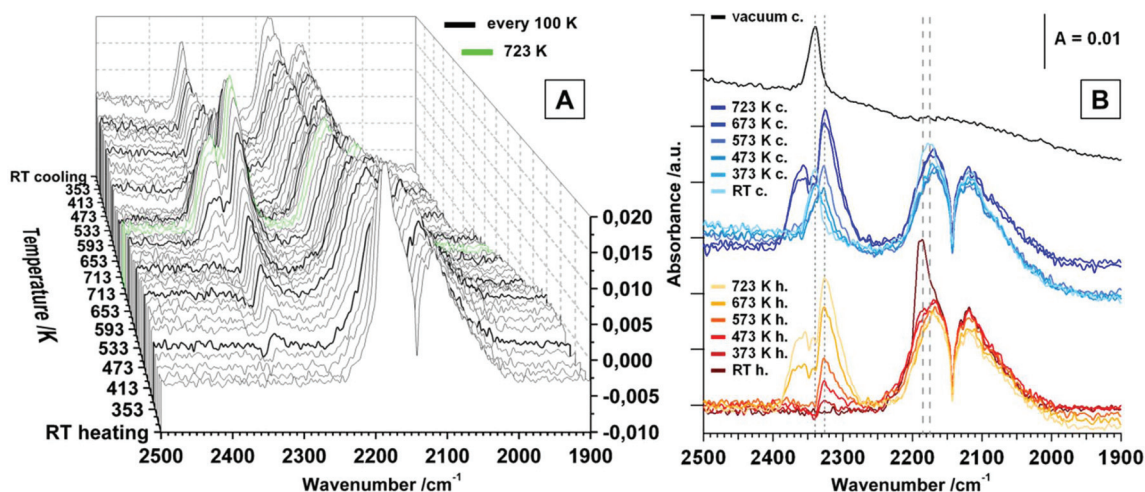


Fig. 2 FT-IR spectra of heating and cooling tetragonal ZrO₂ in a static atmosphere of 20 mbar CO; maximum temperature = 723 K, heating/cooling rates = 10 K min^{–1}. (A) Waterfall plot – spectra are collected every 20 K step. (B) Selected spectra upon heating and cooling, including vacuum spectrum after the heating/cooling cycle.



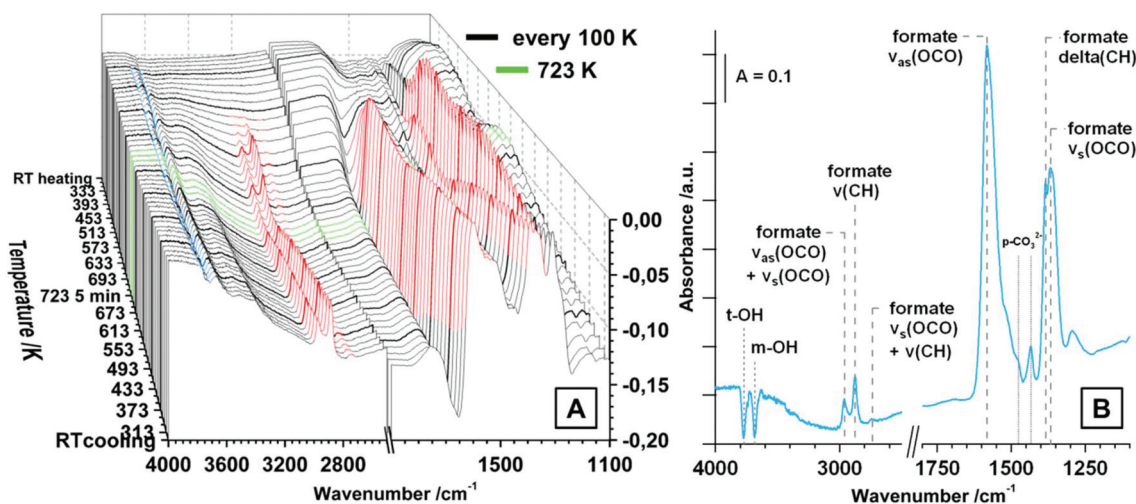


Fig. 3 (A) FT-IR spectra of heating and cooling tetragonal ZrO_2 in flowing CO; maximum temperature = 723 K, heating/cooling rates = 10 K min^{-1} , isothermal period at 723 K = 10 min, flow = $\sim 1.0 \text{ mL s}^{-1}$; (B) FT-IR spectrum of tetragonal ZrO_2 in flowing CO at 673 K.

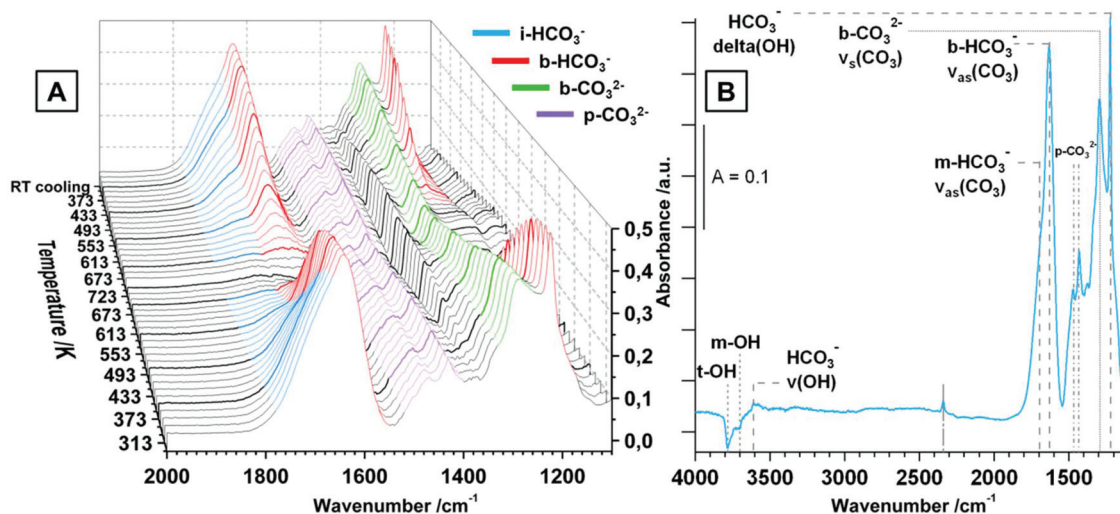


Fig. 4 (A) FT-IR spectra of heating and cooling tetragonal ZrO_2 in 20 mbar CO_2 static atmosphere; maximum temperature = 723 K, heating/cooling rates = 10 K min^{-1} , isothermal period at 723 K = 5 min; (B) FT-IR spectra of tetragonal ZrO_2 in vacuum after the heating and cooling routine in 20 mbar CO_2 .

wavenumber.¹ The evolution of the CO signal upon heating the sample in static 20 mbar CO up to 723 K is subsequently shown in Fig. 2.

The signal for physisorbed CO species decreases with increasing temperature and is no longer present above 373 K. This is correlated with the observation of a peak at 2326 cm^{-1} above 373 K, which is increasing up to 723 K. At first sight, this band could be assigned to a physisorbed CO_2 molecule, but the molecular adsorption of CO_2 on ZrO_2 is usually observed at temperatures below room temperature and the presence at higher temperatures would indicate a high amount of Lewis acidic sites on the surface.¹⁸ Since this signal is actually getting more intense with increasing temperature, this interpretation can be discarded. In studies on CeO_2 this

phenomenon is assigned to result from electronic origin, such as oxygen vacancies or defective lattice sites.^{38–40} It is a well-known fact that CO can be oxidized to CO_2 via two reaction pathways: (i) reaction with a lattice oxygen to CO_2 and an formation of oxygen vacancy and (ii) reaction with a surface hydroxyl group to form formates, which probably can be also decomposed to CO_2 (see also discussion to Fig. 3 below). However, signals of gaseous CO_2 (2400 cm^{-1} – 2270 cm^{-1}) are present above 613 K. Upon re-cooling no gas phase signal for CO_2 is present below 573 K and the signal referred to the oxygen vacancy shifts to 2340 cm^{-1} below 373 K. Molecular adsorption of CO, when re-cooling, is not as pronounced as before the heating and cooling routine and results in a less-shifted peak at 2175 cm^{-1} . Applying vacuum leads to effective



removal of all gaseous and molecularly adsorbed species. The fact that the increase of oxygen vacancies due to surface reduction with CO does not lead to an increased signal for molecularly adsorbed CO species, is consistent with the statement that such an oxygen vacancy rather is a basic adsorption spot. Interestingly, also the amount of acidic Lewis sites such as Zr^{4+} centers is decreased upon the heating and cooling routine. Aside from the molecular adsorption and surface reaction of CO, also a high amount of chemisorbed species is detected. This is illustrated in Fig. 3, which shows the temperature dependency of tetragonal ZrO_2 in flowing CO.

The general absorbance change in Fig. 3A is originating from the pronounced reduction of the sample in pure flowing CO. For a better comprehension of this phenomenon, we refer to the discussion of Fig. 5B, where the absorbance at 2000 cm^{-1} was plotted *versus* the temperature. Derived from similar investigations on TiO_2 and CeO_2 , an increase of the absorbance is associated with the increase of electrons in the conduction band, which, upon reduction *e.g.* in H_2 , is the case for TiO_2 and CeO_2 due to the possibility of donor levels near the conduction band, such as Ti^{3+} and Ce^{3+} .^{39,41} In case of the characterized isolating tetragonal ZrO_2 sample, the absorbance is decreasing with increasing reduction. Thus, CO reduction surprisingly leads to fewer donor states near the conduction band (compare also the discussion about reduction in H_2 , displayed in Fig. 5C). Remarkable is the onset temperature for the absorbance-affecting reduction of 373 K, which is a hundred K before the observation of carbonates, but in good correlation with the detection of oxygen vacancies at 2326 cm^{-1} above 373 K. In Fig. 5B, signals for bidentate formates ($\nu_{\text{as}}(\text{OCO}) + \nu_{\text{s}}(\text{OCO}) = 2962\text{ cm}^{-1}$, $\nu(\text{CH}) = 2878\text{ cm}^{-1}$, $\nu_{\text{s}}(\text{OCO}) + \nu(\text{CH}) = 2743\text{ cm}^{-1}$, $\nu_{\text{as}}(\text{OCO}) = 1583\text{ cm}^{-1}$, $\delta(\text{CH}) = 1385\text{ cm}^{-1}$ and $\nu_{\text{s}}(\text{OCO}) = 1367\text{ cm}^{-1}$ (ref. 32)) are observed above 473 K, increasing up to 573 K. The signals then stay more or less constant upon further heating and re-cooling. Furthermore, negative peaks (species that were present when saving the background or on the initially oxidized surface) of terminal (3774 cm^{-1} , t-) and multi-coordinated (3686 cm^{-1} , m-) OH groups are present above 373 K. In addition, above 373 K bands for polydentate carbonates ($\nu_{\text{as}}(\text{CO}_3) = 1470\text{ cm}^{-1}$, $\nu_{\text{s}}(\text{CO}_3) = 1433\text{ cm}^{-1}$) are observed in flowing CO. The formation of polydentate carbonates can arise also from direct oxidation of CO at *cus* Zr^{4+} centers³⁴ or by chemisorption of initially formed CO_2 on a neighboring O^{2-} .¹⁶ Such reactions are believed to be catalyzed at steps of the crystals.³⁰

Summarizing the spectroscopic results of CO on tetragonal ZrO_2 , two questions arise: (i) which role do the chemisorbed formate species play in terms of surface reduction and (ii) which surface sites are changed, created or blocked during the heating and cooling routine? Upon heating, the signals indicating additional oxygen vacancies, together with signals for polydentate carbonates, already rise a hundred degrees (373 K) before signals for formates (473 K) are observed. Signals for gaseous CO_2 are present at significantly higher temperatures (above $\sim 600\text{ K}$). Upon re-cooling a considerable amount of formates and polydentate carbonates is present, accompanied by a

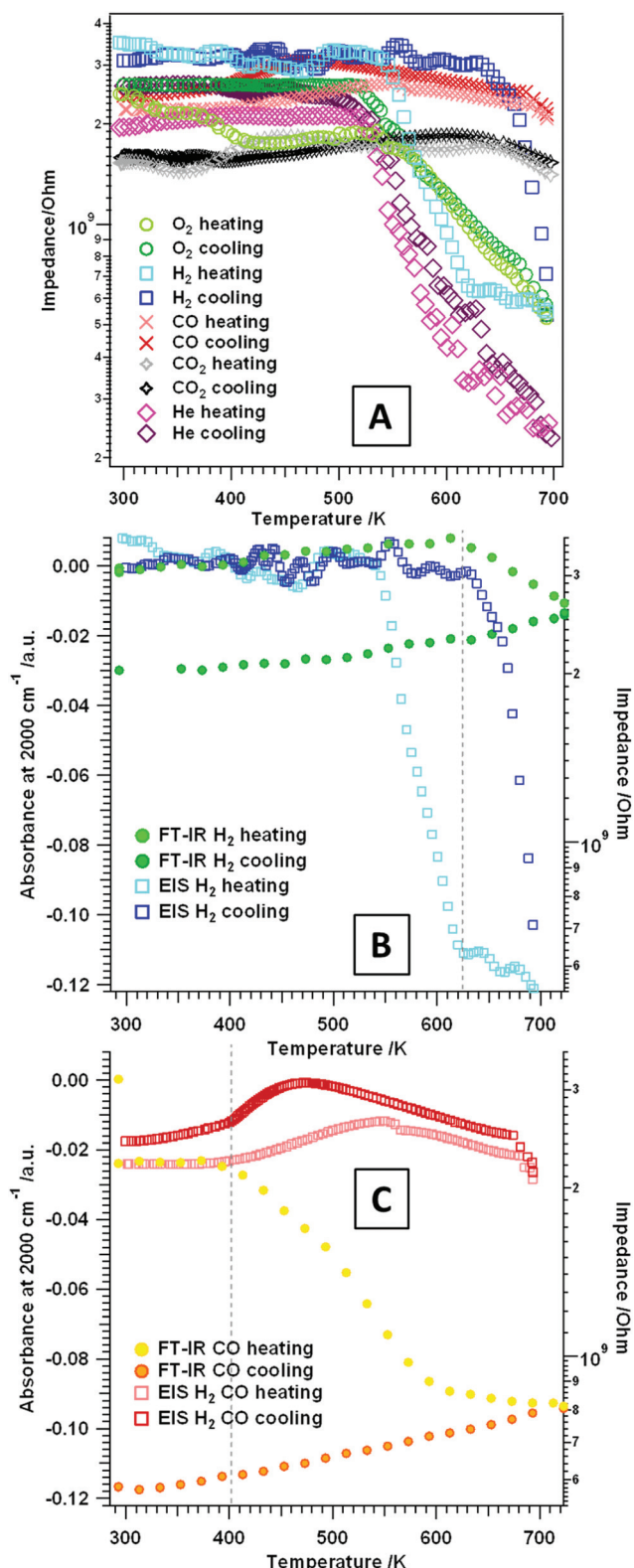


Fig. 5 (A) Temperature-programmed electrochemical impedance results on t- ZrO_2 in dry H_2 , CO , He , O_2 and CO_2 . Panel (B) and (C): plot of the FT-IR absorbance at 2000 cm^{-1} (left y-axis) of tetragonal ZrO_2 in flowing (B) H_2 and (C) CO *versus* the temperature, and the impedance (right axis) *versus* the temperature. Heating/cooling rates = 10 K min^{-1} , flow $\sim 1.0\text{ mL s}^{-1}$, experiments were performed between RT–723 K.

strong reduction of the amount of Lewis acidic centers capable of molecular CO adsorption. This leads to the conclusion that the chemisorbed species (especially the carbonates) block the Lewis surface activity. To verify this effect, an additional experiment with 5% CO₂ seeded in flowing CO was performed (see Fig. S2†). Due to chemisorption of CO₂ already at room temperature, a high amount of (bi-)carbonates is present (for detailed assignment of the species see section CO₂ adsorption). Similar to the experiment in pure CO above 473 K, also formates are observed. The signals for bicarbonates and bidentate carbonates are no more present above 473 K, but the signals for polydentate carbonates persist over the whole evaluated temperature range. Striking is the fact that the decrease of the overall absorbance is strongly suppressed with respect to treatment in pure CO, indicating a much less pronounced (surface) reduction. The reasons could be the quenching/re-oxidation of CO-induced defects by CO₂ or simply the “priming” of the vacancy mediated charge carriers already at low temperatures due to the presence of (bi-)carbonates. Due to the suppressed decrease of the absorbance a reversible decomposition of formates between 633 K and 723 K is visible, which can be correlated with the formation of gaseous CO₂. Upon re-cooling in the 5% CO₂ contaminated CO flow only very weak signals for carbonates are visible, indicating blockage of the essential active sites already during heating. For CO₂ adsorption these are mainly OH groups and O²⁻ anions. Adsorption of CO₂ without the presence of CO is perfectly reversible (see section CO₂ adsorption). So, clear answers to the questions raised above can be stated: (i) formates do decompose to CO₂ and, thus, play a certain role in surface reduction – nevertheless, reduction is pronounced if direct CO oxidation is possible. The vacancy mediated mechanism of CO oxidation is present already at lower temperatures. (ii) A high amount of formates, together with carbonates, effectively blocks surface sites that are related to a direct oxidation mechanism of CO to CO₂. These surface sites are most likely assigned to *cus* Zr⁴⁺ and O²⁻ pairs on steps, edges or corners.^{1,30} The results in CO clearly show that the used tetragonal ZrO₂ sample provides both, reactive Lewis acidic sites, alongside a high amount of hydroxylation.

CO₂ adsorption. Similar to the spectra in CO, also for the CO₂ adsorption studies pure tetragonal ZrO₂ was pre-treated in dry O₂ up to 723 K and the spectrum of the fully oxidized sample at room temperature was taken as background for all further spectra. The increasing pressure of CO₂ between 0.003 mbar and 20 mbar at room temperature is displayed in Fig. S3.† The used tetragonal ZrO₂ is very sensitive to CO₂ chemisorption leading to distinct signals for (bi-)carbonates already at very low pressures around 0.003 mbar. The observed species are monodentate and bidentate bicarbonates m/b-HCO₃⁻ ($\nu_{\text{as}}(\text{CO}_3) \sim 1700 \text{ cm}^{-1}/\nu_{\text{as}}(\text{CO}_3) = 1640 \text{ cm}^{-1}$, $\nu_{\text{s}}(\text{CO}_3)$ = superimposed, $\delta(\text{OH}) = 1223 \text{ cm}^{-1}$), bidentate carbonates b-CO₃²⁻ ($\nu_{\text{as}}(\text{CO}_3)$ = superimposed, $\nu_{\text{s}}(\text{CO}_3) = 1295 \text{ cm}^{-1}$) and polydentate carbonates p-CO₃²⁻ ($\nu_{\text{as}}(\text{CO}_3) = 1470 \text{ cm}^{-1}$, $\nu_{\text{s}}(\text{CO}_3) = 1432 \text{ cm}^{-1}$). The observed chemisorbed species are a product of the reaction of CO₂ with basic sites of

the oxides such as OH groups (leading to m- and b-bicarbonates) and other basic sites such as O²⁻ (leading to b- and p-carbonates).^{16,18,30,32,35} In contrast, molecular adsorption of CO₂ molecules is not observed up to a pressure of ~1 mbar, but at higher pressures a distinct signal at 2352 cm⁻¹ is visible, which gets superimposed by gas phase signals of CO₂ above ~8 mbar. The relatively small shift of the molecularly adsorbed CO₂ peak relative to the gas phase peak vibration indicates a low geometrical distortion and a weak bond.³⁰ Nevertheless, the principal observation of this peak corroborates the presence of a substantial amount of Lewis acid (*i.e.* oxygen-under-coordinated) sites even on the oxidized surface of the used tetragonal ZrO₂.

The distribution of the different kinds of (bi-)carbonates is appalling, especially if compared to literature of CO₂ adsorption on tetragonal ZrO₂. In ref. 32, 34 and 35 already at room temperature the dominant species are polydentate carbonates. In the present case the bicarbonates are predominant, but also the amount of b-CO₃²⁻ seems to be higher than reported in most literature data. This implies not only a very high hydroxylation degree, but also a generally high Brønsted basicity of the used ZrO₂ compared to the reference data. The reason for these differences could be the different preparation routines, which is either precipitation from an aqueous solution of zirconyl chloride^{32,34} or doping with silica³⁵ (leading to doped tetragonal ZrO₂ samples). In contrast, in the present work a sol-gel method by water-quenching a solution of zirconium(iv) isopropoxide in isopropanol (without any other additives) is used.¹⁹

The evolution of the surface species upon temperature treatment of tetragonal ZrO₂ in 20 mbar CO₂ is presented in Fig. 4A. The same experiment was done in flowing CO₂ and the general evolution of the (bi-)carbonate species is the same, but we chose to display the measurement in 20 mbar due to the better separation of the peaks.

With increasing temperature the bicarbonates are stable up to ~473 K, and the signals for these species strongly decrease up to 673 K. In contrast, the bidentate and polydentate carbonate species are not decomposed or removed up to the maximum applied temperature of 723 K. In principal, the removal and formation of all observed (bi-)carbonate species is entirely reversible upon re-cooling and the distribution at room temperature before and after heating is exactly the same (not only in the static measurement but also under flowing conditions). Fig. 4B shows the spectrum of tetragonal ZrO₂ in vacuum after the heating/cooling cycle in CO₂ and the observed bands are highlighted. Additionally to the already assigned wavenumbers of the (bi-)carbonate bands, in vacuum also negative peaks for consumed OH-groups (t-OH at 3784 cm⁻¹ and m-OH at 3700 cm⁻¹) are visible (insertion of a CO₂ molecule into an OH group leads to formation of bicarbonates). The peak at 3610 cm⁻¹ represents the stretching mode of the OH-groups of a bicarbonate adsorbate. The peak at 2340 cm⁻¹ is the same peak that was observed after the heating/cooling cycle in CO but is less pronounced. Due to the fact that this peak is not removed in vacuum (also in active



vacuum over several hours), it is again interpreted as a feature of oxygen vacancies or defective lattice sites.

3.1.2. Electrochemical conductivity

Temperature-dependent measurements between RT and 693 K. Alternating current (AC) impedance analysis was carried out on the t-ZrO₂ sample in different gas atmospheres: H₂ and CO, O₂ and He as an inert reference gas. The aim was to detect changes in the conductivity, as well as eventual stoichiometry changes/phase transformations (especially at higher temperatures, see section 3.2) during the respective gas treatment. In general, when analyzing pressed and sintered powder samples at lower temperatures, the contribution of electronic conduction in the bulk does play a negligible role on samples with such a high band gap. In this case, at lower temperatures conduction is strongly influenced by surface-related mechanisms and grain boundary effects, such as hydroxylation, surface defects or adsorbates.⁴²

To get a better understanding of simple temperature effects and to correlate the influence of the respective gas (oxidative or reductive properties) on the surface chemistry of the studied oxide, EIS measurements in an inert gas (He) were conducted (Fig. 5A pink traces). During treatment in dry He no heating/cooling hysteresis is observed. Furthermore, just as in O₂ (heating/cooling) and H₂ (heating), an insulating period between RT–520 K and semiconductive behavior (*i.e.* decreasing impedance values with increasing temperature) at higher temperatures is detectable. The impedance value at 693 K is interestingly lower ($\sim 2.5 \times 10^8 \Omega$) than the ones obtained at the same temperature in either dry O₂ or dry H₂ ($\sim 5 \times 10^8 \Omega$), but also in any other gas atmosphere. This indicates absence of any reactive surface influences in dry He and, thus, only the pure temperature effect of most likely defect-mediated conductivity is observed.

Fig. 5A also shows the impedance course during treatment of t-ZrO₂ in dry O₂ (green traces). Basically two distinct temperature regions are visible upon heating up to 693 K: the first one is between RT–540 K, where a plateau is visible in the temperature-dependent EIS spectra, followed by the second one starting at 540 K, where the impedance shows semiconductive behavior. The impedance value at 693 K is $5.2 \times 10^8 \Omega$. Upon re-cooling to room temperature a very similar trend is shown: semiconductive behavior between 693–540 K and insulating properties at $T < 540$ K.

Exposure of the sample to dry H₂ and heating up to ~ 540 K (Fig. 5A and B light blue trace) only shows impedance values in the GΩ area, exactly similar as under O₂. Upon further heating, again an impedance decrease (steeper than for dry O₂) of about one order of magnitude is visible. Starting at 623 K an almost plateau-like behavior is apparent with a corresponding impedance value of $\sim 6.5 \times 10^8 \Omega$ (673 K: $5.5 \times 10^8 \Omega$). A different behavior is apparent during re-cooling: between 693–630 K the impedance starts to increase (about one magnitude of order) and at temperatures < 630 K up to RT the sample shows insulating properties with impedance values in the GΩ area. The correlated FT-IR spectra show a decrease in the total transmittance above 623 K which is irreversible

upon re-cooling (see Fig. 5B: plot of the absorbance at 2000 cm^{-1} versus the temperature). A decrease of the absorbance can be referred to the depletion of excitable donor states in the conduction band, which perfectly corresponds to the stagnation of the impedance decrease above 623 K. The analysis in H₂ up to higher temperatures (3.2) prove a dissolution of hydrogen into the used t-ZrO₂ sample and a significant change in the (at least) surface near oxidation state of Zr atoms. Thus, this distinct hysteresis between heating and cooling, in both the impedance as well as the absorbance versus temperature plots, may be an effect of the onset of this process.

A very interesting impedance trend during heating and cooling in dry CO and CO₂ is visible in Fig. 5A. In summary, over the whole temperature region a plateau of the impedance in the GΩ area is apparent. This indicates that the sample is essentially an insulator in these gases and this particular temperature region. It is known from the FT-IR experiments that the sample provides a pronounced surface chemistry, but it is also proven that in both gases the acid/base properties change due to blockage of special surface sites with adsorbates. Due to the fact that especially the Lewis acid sites/*cus* acid–base pairs are affected, this indicates that these sites do contribute to the conductivity of the sample in an “unblocked” state. Also the evaluation of the total absorbance at 2000 cm^{-1} in dry CO (Fig. 5C) confirms, that especially in CO the excitable electronic states toward the conduction band are strongly reduced with increasing temperature ($T > \sim 400$ K) in the evaluated temperature range.

In essence, the conductivity of the characterized t-ZrO₂ sample is considerably influenced by different partially counteracting effects: hydroxylation, the presence of “free” Lewis acid sites, oxygen vacancies and chemical and electronic surface blockage due to chemisorbed adsorbates capable of pinning otherwise thermally excitable states. Up to a temperature of 673 K the sample in dry He shows the highest conductivity with an essentially reversible heating/cooling cycle. In all other gases the surface is reactively influenced and/or changed, which is directly visible in the course of the impedance.

3.2. Gas-phase dependency of the phase transition tetragonal-to-monoclinic ZrO₂

To further shed light on the kinetics of the transformation from tetragonal to monoclinic ZrO₂, the electrochemical impedance experiments in Fig. 6 were deliberately conducted up to temperatures where the phase transformation should occur. Previous high-temperature X-ray diffraction studies revealed that the crystallization of the initially amorphous gel-precursor to the tetragonal phase is finished at 673 K if the aerogel is heated very slowly (effective heating rate: 0.01 K s^{-1}).¹⁹ Upon further heating in air, starting at 873 K, the phase transformation to m-ZrO₂ sets in ref. 19. Therefore, the experiments in Fig. 5 were limited to only 693 K to guarantee phase purity and a high surface area. Note that in the referenced study¹⁹ the *in situ* XRD measurements were conducted in air. To put the



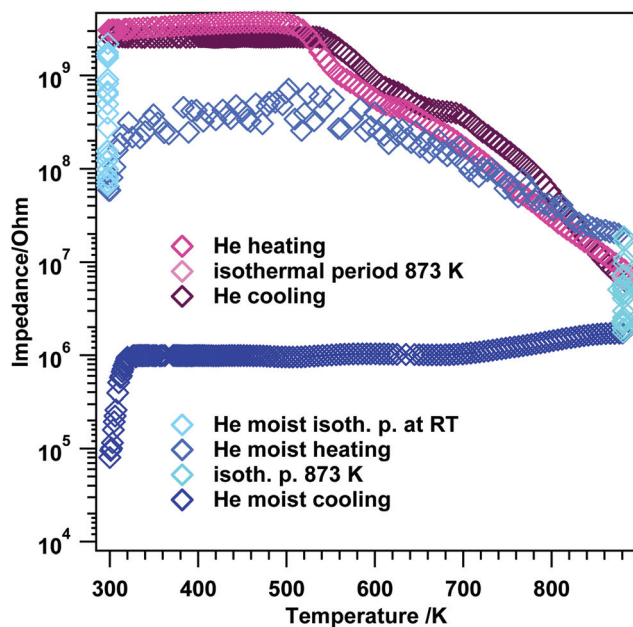


Fig. 6 Temperature-programmed electrochemical impedance spectra on t-ZrO₂ in dry and moist He. Heating/cooling rates = 10 K min⁻¹, flow = ~1.0 mL s⁻¹. The experiments were performed between room temperature and 873 K.

work into literature context, some works^{8,20,43–45} claim that water vapor does strongly promote the phase transformation to m-ZrO₂. Up to a temperature of 693 K the absolute phase stability of the herein used ZrO₂ in moist atmosphere was already proven in ref. 19. Thus, we conducted an EIS experiment where we heated the system up to the exact temperature, where phase transformation is strongly promoted in air (873 K), in dry and moist He (to also check the influence of moist/dry conditions) and routinely checked with *ex situ* XRD to see if the phase transformation to m-ZrO₂ sets in with this comparably fast heating and cooling rates.

3.2.1. Dry and moist He between room temperature and 873 K. Correlated experiments under moist conditions were also performed with the FT-IR setup but already at room temperature there are massive changes in the total transmittance, obviously due to soaking of the sample with water and thus, no reasonable interpretation of further heating experiments was possible. The EIS experiment in dry He (Fig. 6 pink traces) looks very similar to the one conducted up to 693 K (*cf.* Fig. 5). In the beginning, an induction period is visible up to ~520 K after which the impedance starts to decrease leading to a value of $7.0 \times 10^6 \Omega$ at 873 K. During re-cooling to room temperature a very similar trend is apparent. Rietveld analysis (Fig. 7) of subsequent XRD measurements (Fig. S4†) show that during this heating/cooling routine, only 3% of the sample was converted to monoclinic ZrO₂.

At the beginning of the EIS experiment in moist He (Fig. 6 blue traces), the sample was kept at RT for about 10 minutes to account for eventual hydroxylation and/or hydration. During this isothermal period, the impedance decreases almost two

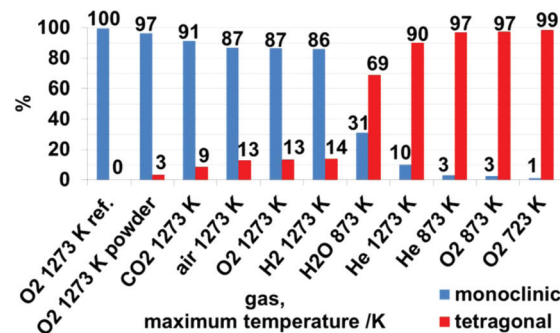


Fig. 7 Weight-percentage of monoclinic and tetragonal phases after treatment in different gases and maximum temperatures, as derived from Rietveld analysis of X-ray patterns in Fig. S4 and S5† and summarized in Table S1.†

orders of magnitude (start: $2.2 \times 10^9 \Omega$, after 10 minutes: $6.6 \times 10^7 \Omega$). Upon heating between room temperature and 323 K the impedance slightly increases again to a value of $3.1 \times 10^8 \Omega$ (de-hydration), after this an induction period up to 573 K is visible and upon further heating, semiconductive behavior leading to a value of $2.0 \times 10^7 \Omega$ at 873 K is apparent. The maximum temperature was again held for 10 minutes, with an impedance decrease to $1.8 \times 10^6 \Omega$. Upon re-cooling to room temperature, basically an almost constant impedance value up to ~323 K is obtained and at temperatures $T < 323$ K, the impedance starts to decrease again leading to a final value of $8.0 \times 10^4 \Omega$ at room temperature.

Rietveld analysis (Fig. 7 and S4†) show that upon heating and cooling in the impedance setup with a comparable fast rate of 10 K min⁻¹, pronounced crystallization of the monoclinic phase is observed for the moist treated sample (31 wt% monoclinic ZrO₂). In contrast, heating to lower maximum temperatures (compared to ref. 19) in dry He resulted in a comparably lower extent of transformation to monoclinic ZrO₂ (3 wt% monoclinic ZrO₂). Thus, these data confirm the water promoting role of the phase transformation to a certain extent above 693 K. According to ref. 8 a moist atmosphere promotes grain growth (and related phase transformation) and leads to enhanced surface diffusion. Nevertheless, note that for the tetragonal ZrO₂ sample characterized in this work the impact of water vapor is not as pronounced as one could expect from literature data, since in the present case the promoted phase transformation is only observed at temperatures, where kinetically controlled phase transformation would take place anyway. The very unusual course of the impedance in this experiment also indicates a strong interaction of water vapor with the sample. In fact, a 30% increased bulk conductivity (most likely proton conduction) due to incorporation of water into doped tetragonal ZrO₂ is also observed by Guo *et al.*⁴⁵ As a matter of fact, water can replenish the oxygen vacancies, which reduces the amount of defects, subsequently leading to phase transition. The main difference to ref. 45 is that the conductivity measurement in water lasts 8 h. After this treatment, 70 wt% monoclinic ZrO₂ is found in the structural mixture.



Our sample, however, is chemically and structurally pure (*i.e.* not Y-doped) and the admission time of water is much shorter.

3.2.2. Gas phase dependent characterization up to 1273 K.

In general, the allotropic phase transformation of tetragonal to monoclinic ZrO_2 is described as a martensitic twinning mechanism, with a complex dependence on temperature, stress and environment.⁴⁶ However, based on the study of Shukla and Seal,⁸ the phase transformation of pure undoped tetragonal ZrO_2 to monoclinic ZrO_2 is influenced by the following factors: basically derived from calculations for surface energy, stabilization of small particles exceeding a critical particle size of 10 nm leads to phase transformation. Possible aggregation of nano-crystallites enhances the critical size up to 33 nm (additional interfacial energy has to be taken into account). The metastable tetragonal phase at lower temperatures is stabilized by defects (critical defect concentration ~ 1.7 mol%),⁴⁵ which is also one of the discussed origins (besides lattice entropy increase) of initial phase transformation from m- ZrO_2 to t- ZrO_2 and c- ZrO_2 at high temperatures.⁴⁶ The existence of room temperature meta-stable tetragonal particles with even larger particle sizes (< 200 nm) may, thus, be explained by strain energy of aggregated ZrO_2 domains below the critical diameter of 33 nm. However, these energy-calculations do still not take possible additional stabilization effects such as OH^- anions (or hydrogen “impurities”) into account. A recent DFT study by Youssef and Yildiz also suggests that zirconium vacancies can act as hydrogen accumulators.⁴⁷

To focus on the phase transformation at even higher temperatures, the tetragonal ZrO_2 sample used in this work was heated up to 1273 K in pure He, O_2 , H_2 and CO_2 to examine a potential gas phase dependency of the phase transformation to monoclinic ZrO_2 . The standard temperature program was heating and cooling with 10 K min^{-1} and a gas flow of $\sim 1 \text{ mL s}^{-1}$ with a maximum temperature of 1273 K, *i.e.* a rather fast heating routine. This routine was performed with the *in situ/operando* setups (EIS and FT-IR), using pelletized samples rather than powders, but sintered and oxidatively pretreated as described in the Experimental section. Subsequently, the samples were checked by XRD and XPS.

Qualitative and quantitative phase analysis by XRD. As expected (compare kinetic phase transformation investigations in air in ref. 19) for O_2 , H_2 and CO_2 the main part of the samples was transformed to monoclinic ZrO_2 . As already discussed in the section above the presence of H_2O vapor drastically promotes the phase transformation. The corresponding XRD data of the measurements up to 1273 K in the different gas atmospheres are shown in Fig. S5.†

Derived from Fig. 7, the samples treated in dry O_2 and dry He up to 873 K show a low conversion to m- ZrO_2 of 3%, in contrast to moist He up to 873 K, as discussed above. Heating the sample in dry He up to 1273 K also leads to a low amount of 10% m- ZrO_2 – which is a significantly low amount. Based on the assumption that a sufficient defect concentration is crucial for the stabilization of the tetragonal phase,⁴⁵ this is originat-

ing from the missing defect quenching (such as in oxidative reactants or other possible reactive interaction with gases like H_2 and CO_2 , which is confirmed in the discussion to Fig. 8) and the absent active surface chemistry influencing the transformation kinetics upon heating in dry He. Treatment of the sample in dry O_2 , H_2 and air up to 1273 K leads to similar amounts of 86–87% m- ZrO_2 . Heating/cooling in CO_2 results in 91% m- ZrO_2 . The striking differences to He can be explained by a substantial reduction of the defect concentration of the sample because of reaction with the other gases. In the case of O_2 this is simply quenching of a sub-stoichiometric oxygen-deficient lattice. Derived from subsequent XP spectra of the sample pellets (see detailed discussion below, Fig. 8) treatment in H_2 and CO_2 leads to formation of dissolved hydrogen and Zr-(oxy-)carbide – at least at the surface. Such reactive modifications of the sample are not present at identical treatments of the monoclinic reference sample, and this indicates an initially highly defective oxide.

For characterization in the *in situ/operando* setups (but also for the preparation for the *ex situ* XRD, XPS and AFM measurements) the starting material was pressed into pellets and *in situ* sintered in the respective impedance and FT-IR setups. To check for effects derived from these procedures, an identical heating/cooling routine in O_2 up to 1273 K was performed for a powder of the tetragonal starting material, resulting in 97% m- ZrO_2 , which is 10% more than for the pressed sample. As will be discussed in detail below this can be explained by surface, crystallite sizes and agglomeration effects that enhance the stability of the tetragonal phase.⁸

***In situ* heating synchrotron XRD analysis.** To study the effect of crystallite size growth on the transformation temperature of t- ZrO_2 to m- ZrO_2 under different gas atmosphere, the pathway of phase transformation was investigated by *in situ* synchrotron XRD in H_2 and CO_2 atmospheres. Before the *in situ* XRD experiments, the t- ZrO_2 sample were obtained by calcination of the powdery amorphous aerogel at 723 K inside the capillary under air flow until crystallization to tetragonal ZrO_2 is completed. The contour plots in the Fig. 8a–d show the recorded diffraction data for one heating/cooling cycle in H_2 and CO_2 atmospheres. The *in situ* data reveals that the initially phase-pure t- ZrO_2 remains stable upon heating to 1273 K in both gases. The tetragonal to monoclinic transformation takes place during cooling down from 1273 K to room temperature at 625 and 715 K in H_2 and CO_2 atmosphere, respectively. The additional h- SiO_2 reflections result from a partial crystallization of the fused silica quartz capillary and are not related to the zirconia sample. The final patterns of the cooled samples show m- ZrO_2 as the major phase and a minor t- ZrO_2 side phase. The Rietveld refinement reveals that the cooled samples to room temperature still have 14.9 wt% and 9.8 wt% of tetragonal ZrO_2 in H_2 and CO_2 atmosphere, respectively, which in good agreement with *ex situ* XRD analysis (Fig. 7). Selected patterns are presented in Fig. 9.

The stabilization of the metastable t- ZrO_2 phase can be explained by a combined influence of (a) a high defect density and (b) a very small crystallite size, where a decreased defect



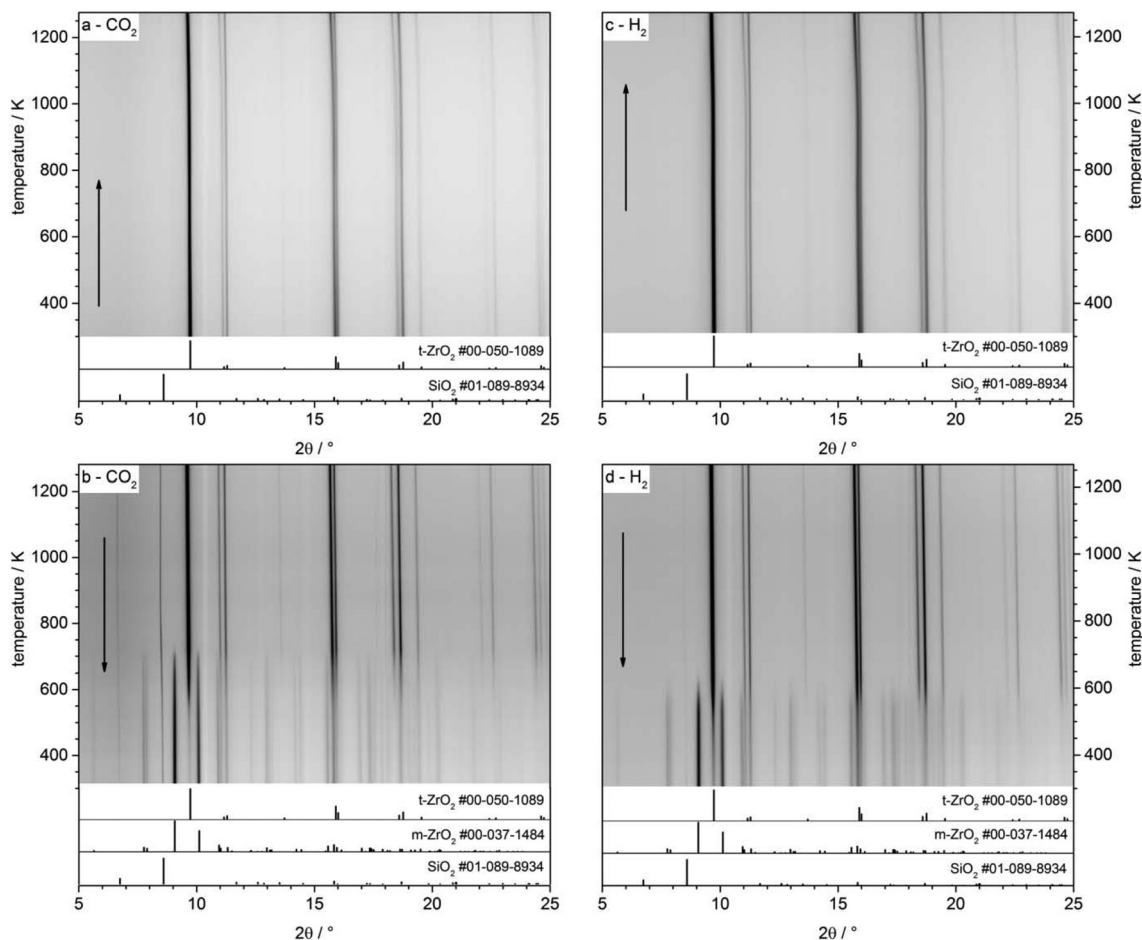


Fig. 8 Temperature vs. 2 theta contour plots of *in situ* high temperature synchrotron XRD characterization of tetragonal ZrO_2 heated to 1273 K and subsequently cooled down to room temperature with 20 K min^{-1} in carbon dioxide (a-heating and b-cooling) and hydrogen (c-heating and d-cooling). Dark colors denote high intensities, grey-white colors background intensities. ICDD reference diffraction data for t- ZrO_2 , m- ZrO_2 and h- SiO_2 shown at the bottom of each plot.

density and/or an increased crystallite size favors the stable monoclinic ZrO_2 . Rietveld refinement (strain was considered with Stokes-Wilson model, see strain plot in ESI†) results revealed that the initial t- ZrO_2 samples have a crystallite size of 26.5–32 nm (around the critical diameter of $\approx 30 \text{ nm}$) ref. 8. The results of Rietveld refinement were further confirmed by calculating the crystallite size D (20.4 nm) from S_{BET} surface area ($50 \text{ m}^2 \text{ g}^{-1}$) using the well-known relationship ($S = 6/\rho D$) ref. 8 where ρ the density of t- ZrO_2 (6.121 g cm^{-3}). The crystallite size of the heated samples was almost constant up to 1073 K followed by a sharp increase above this temperature in both gases, see Fig. 10 (left panel). The maximum crystallite size of t- ZrO_2 at 1273 K was found to be 41 and 43.5 nm in H_2 and CO_2 atmosphere, respectively. The size of crystallites remain constant during cooling in both atmospheres up to the transformation temperatures followed by a sharp decrease below transformation temperatures. Strain is released in the tetragonal phase above 1000 K when the grain size increases (see Fig. S6†). The newly formed monoclinic ZrO_2 shows higher microstrains – especially in H_2 – which can be

attributed to a defect rich material. The higher transformation temperature in CO_2 in comparison to H_2 atmosphere may be due to the higher crystallite size of t- ZrO_2 in CO_2 than in H_2 atmosphere. This finding is consistent with previous studies showed that as the crystallite size of ZrO_2 increases, the monoclinic-to-tetragonal phase transformation temperature increases ref. 8, 48 and 49. However, the interaction of different gas atmosphere with site defects on ZrO_2 could be also another reason to lower the transformation temperature in H_2 atmosphere as will be discussed below.

Electronic structure analysis by XPS. The *ex situ* XP spectra in Fig. 11 reveal changes in the chemical states of the surface-near region. Mixed Gaussian/Lorentzian peak models were fitted for the $3d_{5/2}$ and $3d_{3/2}$ peaks (as outlined in the Experimental section 2.4). The assignment of the oxidation states was done according to literature values found in ref. 24 and 50. For the sample heated in dry He to 873 K (Fig. 11A), a relatively large fraction (almost 50 at%) of the total accessible Zr species (inelastic mean free path = 2.3 nm; 1304 eV kinetic energy, calculated *via* the G-1 predictive formula⁵¹) is found to



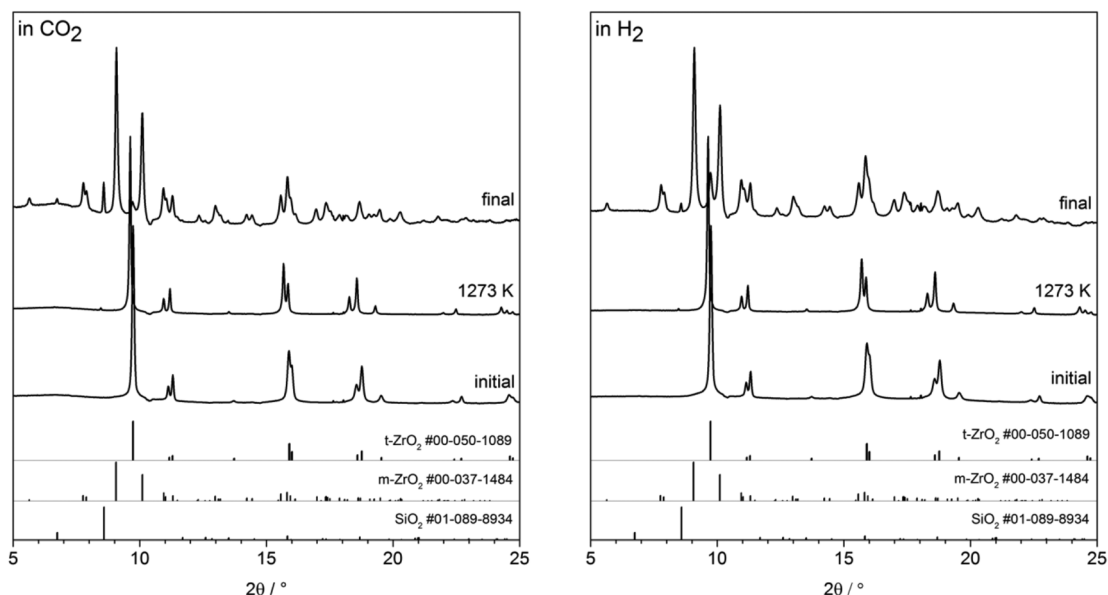


Fig. 9 Selected XRPD patterns from the *in situ* series. Patterns of initial sample, at 1273 K and final sample are shown for CO₂ (left) and H₂ (right) atmosphere. ICDD reference diffraction data for t-ZrO₂, m-ZrO₂ and h-SiO₂ shown at the bottom of each plot.

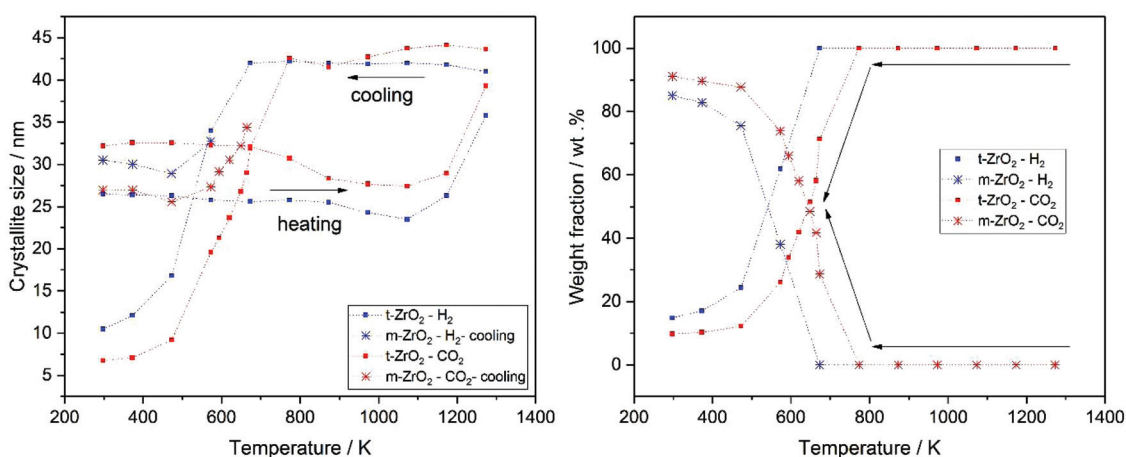


Fig. 10 Crystallite size evolution of t-ZrO₂ crystals during heating and cooling in H₂ (blue) and CO₂ (red) (left) and tetragonal and monoclinic phase fractions during cooling (right) as derived from Rietveld refinement of selected *in situ* XRPD data.

be in a lower oxidation state than Zr(IV). This is also derived from the general peak shape, as a distinct shoulder at a binding energy of about 181 eV, as well as the apparent weaker peak splitting between the 3d_{5/2} and 3d_{3/2} components is observed. 34 at% of the Zr ions can be assigned to an oxidation state of +III, which is shifted relative to the +IV state (182.8 eV) by 1.1 eV, and the remaining 15.5 at% appear reduced to Zr(II) species (with a shift of 2.1 eV to the Zr(IV) species). After treatment at 1273 K (Fig. 11B), the amount of Zr(IV) at 63 at% is significantly larger. The fraction of Zr(III) is reduced to 25 at%, whereas the concentration of Zr(II) is only decreased slightly (to 12 at%). As no 'external' oxidation of the sample is possible during treatment in helium, the now larger fraction of Zr(IV) has to originate from the sample itself. As the

portion of Zr(II) did not increase, a disproportionation reaction of Zr(III) to form Zr(IV) and Zr(II) must take place, possibly overlaid with a higher oxygen mobility at elevated temperatures, *i.e.* bulk oxygen segregates to surface-near regions. That the majority of thus oxidized species seem to be Zr(III), indicates that the reaction from Zr(III) to Zr(IV) is favored compared to the oxidation of Zr(II) to form Zr(III).

Fig. 11C shows the spectrum obtained after heating the sample to 873 K in moist He. The peak is not as broad as compared to the above-discussed spectra, without the presence of a strong shoulder. Both can be directly related to the complete absence of Zr(II). The experimental spectrum is composed of Zr(IV) (72%) and Zr(III) (28%). This suggests that water is able to oxidize the Zr(II) that was found for the dry He treatment at



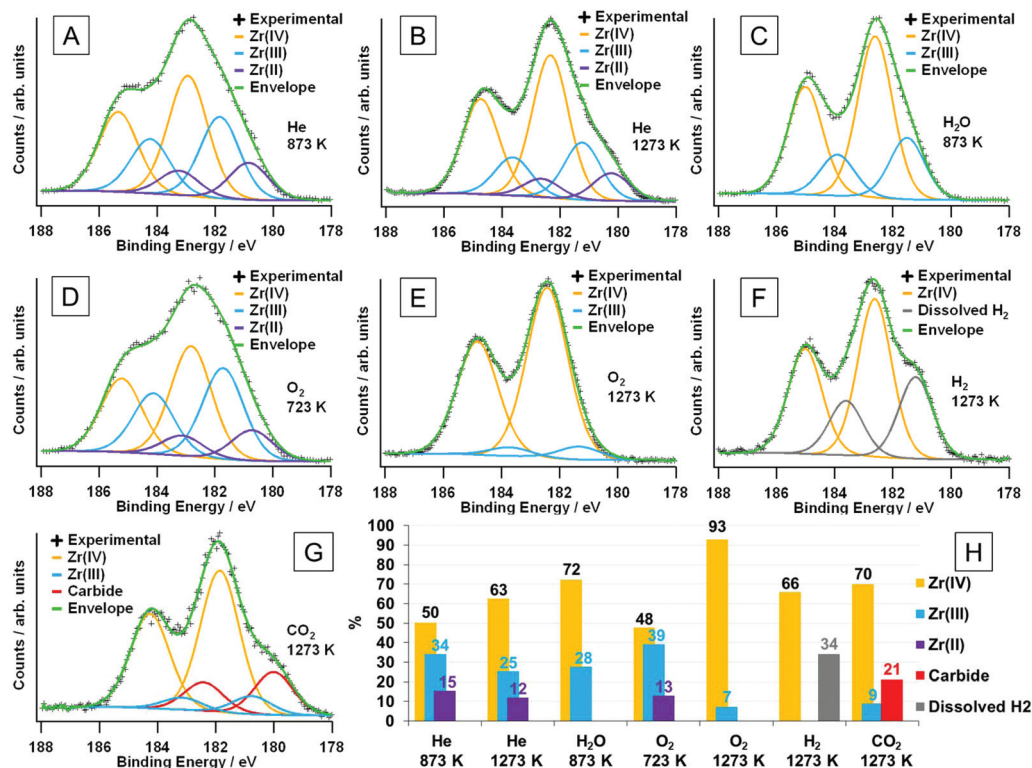


Fig. 11 XPS Zr 3d regions for the samples after various treatments, including fits of the oxidation states. Samples were treated (heating/cooling rate of 10 K min⁻¹) in flowing (flow ~1 mL s⁻¹) dry He up to (A) 873 K and (B) 1273 K, moist He up to 873 K (C), dry O₂ up to 723 K (D) and 1273 K (E), and dry H₂ up to 1273 K (F) and dry CO₂ up to 1273 K (G). The relative at% of the observed surface species (as determined by the chemical state fits) are shown in panel (H). The maximum depth of emission for the Zr 3d peak, obtained via the inelastic mean free path for photoelectrons with this energy (1304 eV kinetic energy) calculated via the G-1 predictive formula⁵¹ is 2.3 nm. Corresponding C 1s peaks are shown in Fig. S7.†

the same temperature, as well as to oxidize some of the Zr(III), since the amount of that state is also lower. In pure dry O₂ at 723 K (Fig. 11D), the spectrum looks very similar to that for He (873 K), which indicates that the temperature is too low to effectively oxidize the sample. Only 48% are present in the form of Zr(IV), whereas nearly the same amount (39%) is present as Zr(III), and 13% are Zr(II). The Zr(IV)/Zr(III) ratio is even lower than for the helium-treated sample. Upon increasing the oxidation temperature to 1273 K the suboxides vanish almost completely, and there is only a minor fraction of 7% Zr(III) left (Fig. 11E).

After reduction in dry H₂ at 1273 K (Fig. 11F), the general peak separation is comparable to treatments either in O₂ at 1273 K or in H₂O at 873 K with a well-defined shoulder at about 181 eV. The fits reveal that – in addition to Zr(IV) – there is a second component, which is neither Zr(III) nor Zr(II), as its shift relative to Zr(IV) is 1.4 eV, and is, thus, found in-between those two oxidation states (which would be at 1.1 eV and 2.1 eV, respectively). The shift, nevertheless, is also not large enough for it to be zirconium hydride, ZrH₂, which should yield peaks at a binding energy of 178.8 eV.⁵² This could indicate hydrogen dissolution inside the crystal structure, which either shifts the neighboring Zr(IV) or Zr(III) energetically to lower binding energies.⁴⁷ What would be in favor of the Zr(III) ions being affected is that even the spectra taken after O₂ treat-

ment contain a small fraction of suboxides. A sample reduced in pure H₂ should feature an even larger portion, and the quantification of the respective spectra would yield a sum formula of ZrO_{1.88}, which agrees with the presence of Zr(III). This hydrogen-influenced species gives rise to 34% of the total Zr 3d signal.

In Fig. 11G, the sample was annealed to 1273 K in CO₂, which gives rise to another distinct shoulder, albeit at lower binding energies as compared to treatment in hydrogen. It is located at about 180 eV and is caused by a component relatively shifted to the Zr(IV) peak by 1.9 eV. The C 1s region (Fig. S7†) reveals a severely asymmetric peak being due to a Zr carbide component at 282.9 eV,⁵³ which appears as a shoulder in the Zr 3d region. The latter peak (as the C 1s component) is shifted to higher binding energies when compared to the reference values for ZrC.⁵³ The reason for this could be that it is not a pure ZrC layer, but rather a ZrO_xC_y oxy-carbide, where the remaining oxygen could influence the surrounding Zr and C atoms and cause the respective chemical shifts. This could be either because the ZrC layer itself contains oxygen or that it is indeed ZrC, but is so thin that the oxide/carbide interface is sampled as well. Taking into account the different escape depths between the Zr 3d and C 1s regions, the carbide components in both spectra yield the same absolute amounts of carbide: 21% of the surveyed zirconium and 18% of the



present carbon species (the rest being adventitious carbon), with a total of Zr/C ratio of 1.2.

Surface modifications including dissolved hydrogen species and Zr-carbide in H_2 and CO_2 are not observed in comparable experiments of monoclinic ZrO_2 (preceding study of H_2 on m- ZrO_2 in ref. 54 no carbide in the XPS spectra in Fig. S8†). Starting with a tetragonal sample, which is highly defective and hydroxylated and, thus, more active, seems to strongly promote such surface modifications.

The essentially different phase transformation behavior in He compared to the other gases can be explained as follows: starting with a defect concentration sufficient enough to stabilize the tetragonal phase at room temperature, simple heating only leads to an unaffected increase of the defect concentration and further stabilization of the tetragonal phase. Also upon cooling, no quenching or reaction of these defects is possible in dry He and, thus, a less pronounced phase transformation to monoclinic ZrO_2 is observed. In oxidative gases such as O_2 and H_2O direct defect quenching is possible and phase transformation upon cooling is unavoidable. Also the formation of (oxy)-carbides or dissolved hydrogen species can be seen as a unique form of defect alteration and blockage.

A final note on the optical change of the sample color should be added at this point. The tetragonal sample annealed at 673 K exhibits a grayish/brownish color, likely due to a certain amount of reduced centers. The sample treated in He up to 1273 K is dark gray, indicating an increased amount of color centers. Treatment in dry H_2 leads to an almost black pellet. The samples after treatment in water up to 873 K and O_2 and CO_2 up to 1273 K are white, which matches the assumption of different, but nevertheless oxidative quenching mechanisms.

Surface morphology as determined by AFM: influence of powder pressing. To sum up the XRD and XPS data, derived from the distinct surface modifications in H_2 and CO_2 , compared to pure monoclinic ZrO_2 the surface chemistry is strikingly different. Another remarkable effect is the influence of pressing and *in situ* sintering the sample compared to treatment of a loose powder as starting material. The XRD pattern in Fig. S5† and the corresponding Rietveld analysis in Fig. 7 and Table S1† clearly show that it makes a difference whether the original aerogel is heated in a pressed or a powderous form. Whereas the powder sample (starting with a fine milled aerogel) is 97% monoclinic after heating in dry O_2 the pressed pellet still provides 13% of tetragonal phase. To further shed light on this phenomenon, AFM measurements of the pellet after treatment in O_2 up to 1273 K were performed (Fig. 12). The AFM images obtained at a larger scale ($5 \times 5 \mu\text{m}$, Fig. 12C) clearly show that the pellet is composed of large grains (with a grain size of about 200 nm, as obtained from a histogram). Fig. 12D makes it easier to visualize the surface, as in this 3D image, the x-, y- and z-axes all have the same scaling, revealing that the surface is rather flat, with the grains having a low profile (the z-extent of the image is 283.9 nm). Upon zooming into one such grain (Fig. 12A, z: 65.2 nm), it can be seen that a lot of small crystallites sit at its surface, which exhibit an

improved contrast in the respective phase image (Fig. 12B). By measuring the size of more than 200 such grains in the phase image and fitting a Gaussian function to the resulting histogram, the most frequent crystallite size could be determined to be 16 nm (Fig. 12E). According to ref. 8 this particle size implicates stabilization of the tetragonal phase. We conclude that the main bulk of the pressed and sintered sample is transferred into monoclinic phase upon heating to 1273 K, but the surface still exhibits small tetragonal particles. In this context, Shukla and Seal⁸ also claim that agglomeration of many small crystals does stabilize possible μm -sized sintered tetragonal structures. Note that for the present experiments the aerogel was pressed into pellets and subsequently sintered in this “shape” inside the *in situ* setups. Treating such a pellet in O_2 up to 1273 K leads to phase transformation to monoclinic ZrO_2 but still a distinct amount of tetragonal ZrO_2 is present (13 at% t- ZrO_2), as derived from the AFM measurements: very likely on the surface of the pellet derived from the 16 nm size of the particles on the surface of the pellet.

Conductivity measurements at elevated temperatures between RT–1273 K. Based on the XRD, XPS and AFM data, the corresponding EIS measurements in different gas atmospheres up to 1273 K can now be critically discussed. For detailed discussion of data obtained with a maximum temperature of 693 K, we refer to section 3.1.2. Additional comparative reference experiments were also done for monoclinic ZrO_2 , which are displayed in Fig. S9.†

Exposure of t- ZrO_2 to dry O_2 and heating up to 1273 K (Fig. 13, green trace) shows semiconductive behavior starting at $T > 560$ K with a step between 560–700 K, after which the impedance starts to decrease faster up to ~ 1013 K. At higher temperatures, the impedance decrease flattens, leading to a value of $5.8 \times 10^5 \Omega$ at 1273 K. The same trend is observed during cooling with minor differences in the insulating period between 660–RT and a missing step between 560 and 700 K. Compared to m- ZrO_2 during treatment in O_2 (Fig. S9A†), a somewhat similar trend for both oxides is observed up to 973 K. At temperatures above 973 K the impedance decreases less strongly for t- ZrO_2 than for m- ZrO_2 leading to slightly different final values at 1273 K (m- ZrO_2 : $1.2 \times 10^4 \Omega$; t- ZrO_2 : $5.8 \times 10^5 \Omega$). This could be tentatively assigned to kinetic phase transformation/oxygen quenching processes in the tetragonal sample.

Treatment in H_2 up to 1273 K (Fig. 13 blue traces) reveals slight differences especially in the semiconductive region, leading to a final impedance value of $1.4 \times 10^4 \Omega$ at 1273 K. Again, re-cooling yields similar results between 1273–973 K, but at lower temperatures, differences to the heating process are evident. Two plateaus are apparent between 760–652 K ($3.8 \times 10^8 \Omega$) and 652–RT ($4.8 \times 10^8 \Omega$). In contrast, exposure of m- ZrO_2 to dry H_2 (Fig. S9B†) leads to semiconductive behavior at temperatures $T > 415$ K and a value of $1.2 \times 10^5 \Omega$ at 1273 K. This indicates that in dry H_2 on t- ZrO_2 the impedance at the highest temperature is about one order of magnitude lower than on m- ZrO_2 . Together with the strongly deviating behavior of re-cooling the t- ZrO_2 sample in H_2 , this effect may be due to



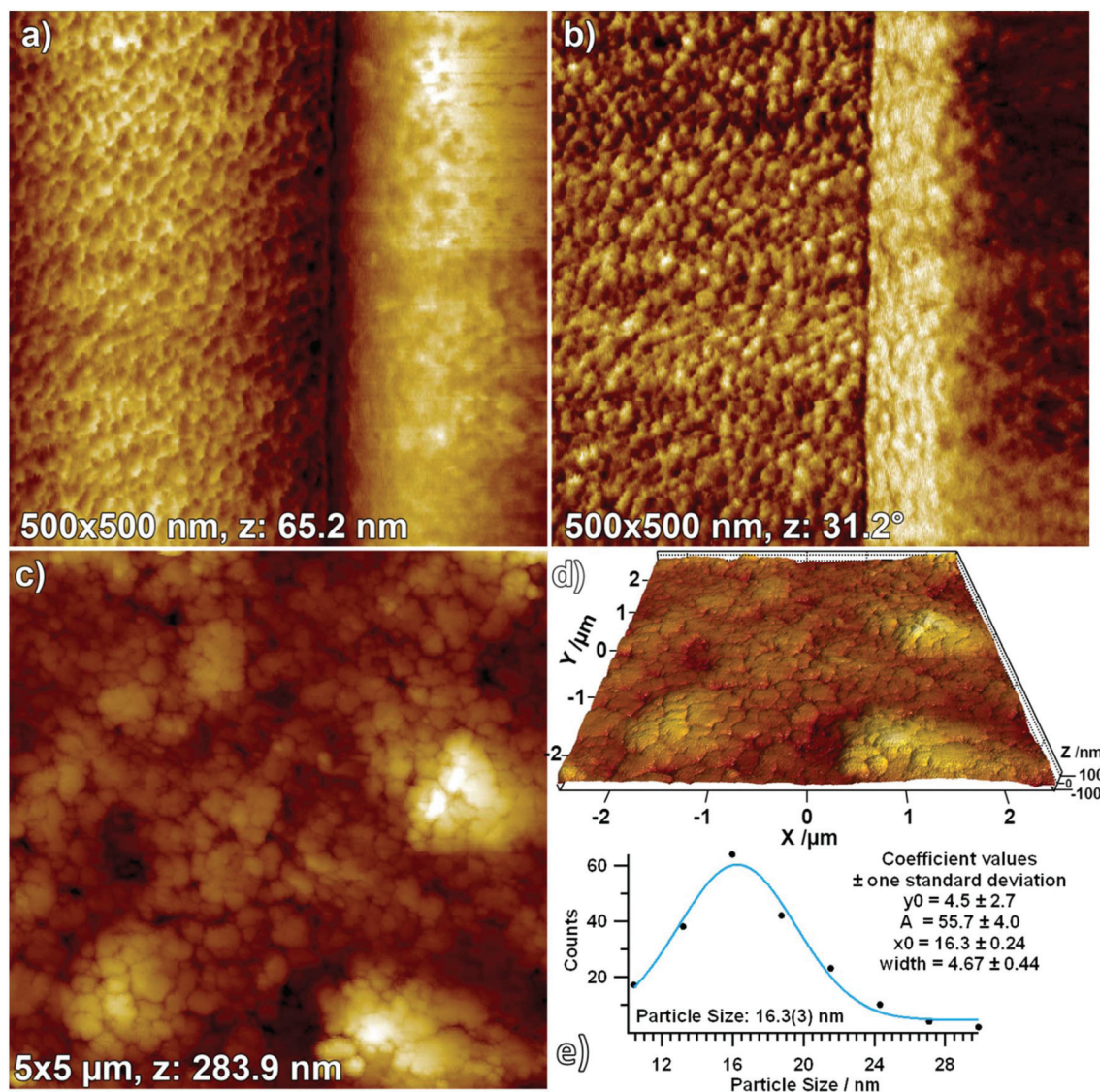


Fig. 12 500 × 500 nm AFM topography (a) and phase (b) images, as well as a 5 × 5 μm topography image (c) with 3D rendering (d). (e) Histogram of crystallite sizes as obtained from (b) with a fitted Gaussian function.

the above-mentioned dissolution of hydrogen in the sample, which is not the case for pure m-ZrO₂.⁵⁴

Striking differences during the temperature-dependent EIS analysis in CO₂ are illustrated in Fig. 13 (grey and black traces). Between RT–673 K a pronounced insulating period in combination with semiconductive behavior, exhibiting several steps in the decrease of the impedance, is apparent ($1.0 \times 10^6 \Omega$ at 1273 K). As in hydrogen, during re-cooling a similar trend as during heating in the high-temperature region 1273–1126 K is apparent, but at lower temperatures differences arise. Higher impedance values as during heating are obtained down to ~820 K, followed by the insulating period. The monoclinic sample (Fig. S9C†) exhibits an insulating period during heating between RT–673 K and 860 K–RT during cooling as well as an impedance value of $2.5 \times 10^5 \Omega$ at 1273 K. From the FT-IR experiments one can deduce that the surface chemistry

of the used t-ZrO₂ is strongly enhanced at lower temperatures compared to the used monoclinic ZrO₂ sample.^{42,55} The XPS investigations show that on the t-ZrO₂ sample (oxy-)carbides are formed (which is not the case for the monoclinic sample – compare Fig. S9†). Since a fully percolated (oxy-)carbide layer would lead to a better conductivity, a full (oxy-)carbide surface percolation can be excluded due to the increased impedance values of t-ZrO₂ at 1273 K compared to m-ZrO₂.

The reference measurement in dry He again provide interesting results (pink traces Fig. 13). Upon heating to 1273 K a very short insulating period (RT–520 K) together with semiconductive behavior at temperatures >520 K is apparent. However, during He treatment the impedance starts to decrease faster as compared to O₂, but comparable to H₂. The value at 1273 K is lower ($4.3 \times 10^4 \Omega$) than in O₂ and CO₂ and a little bit higher than in H₂ ($1.4 \times 10^4 \Omega$). The comparison between m- and



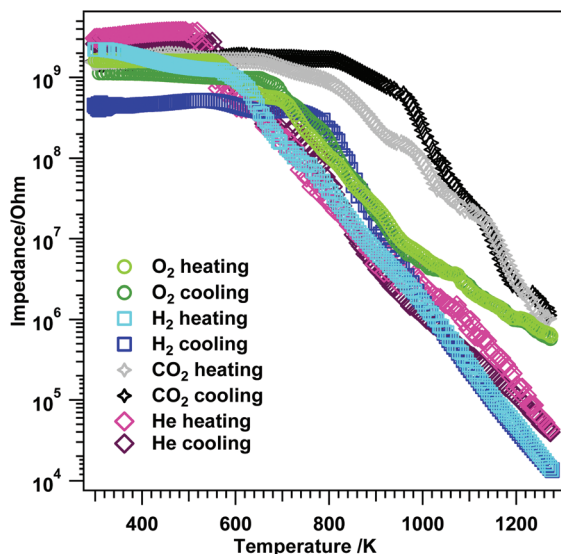


Fig. 13 Electrochemical impedance versus temperature results on t-ZrO₂ in dry O₂, H₂, CO₂ and He. Heating/cooling rates = 10 K min⁻¹, flow = ~1.0 mL s⁻¹, experiments were performed between RT–1273 K.

t-ZrO₂ in dry He (Fig. S9D†) again helps to provide information on pure temperature-induced conductivity effects on the samples. Basically, almost an identical impedance course is observed for both oxides with some slight differences in the value at the highest temperature (m-ZrO₂: $1.1 \times 10^5 \Omega$ and t-ZrO₂: $4.3 \times 10^4 \Omega$). This again emphasizes the huge influence of the gas atmosphere to the overall electrochemical response of the sample.

To sum up the EIS investigations up to 1273 K, we note a very strong dependence of the conductivity on the nature of the gas atmosphere, which corroborates the XPS data. This might especially affect the surface and grain boundary contribution conductivity mechanisms. The phase transformation itself cannot be observed “directly”, but confirms that the latter is most likely a rather continuous process.

4. Conclusions

The conclusions derived from the presented work can be summarized as follows:

- the characterization of the pure tetragonal ZrO₂ compound below 723 K reveal an active surface with a high degree of hydroxylation. Lewis-acidic and Brønsted-basic centers are both found on the surface. Both CO and CO₂ are effectively blocking coordinately unsaturated sites and, thus, no conductivity has been observed in these C1 gases. However, already in this temperature region, significant differences in the surface conductivity as a function of the gas atmosphere arise.

- Water promotes the transformation from tetragonal to monoclinic ZrO₂ especially around 873 K. This has been worked out in a combined spectroscopic and structural study by impedance, FT-IR and X-ray diffraction measurements. Moreover, the tetragonal ZrO₂ sample exhibits pronounced

interactions with water, leading to additional protonic conduction. Most importantly, this transformation is observed at much higher temperatures than reported in literature.

- The high-temperature investigations up to 1273 K as a function of the gas atmosphere reveal distinct differences, ranging from being basically unaffected (He), over dissolved hydrogen (in H₂) to reaction ((oxy-)carbide formation in CO₂). The phase transformation is also reported to be strongly affected by the sintering and pressing history of the sample.

- Both defects and hydroxylation are the decisive parameters for steering the structural stability and reactivity of the phase-pure tetragonal ZrO₂ modification. Due to critical control of these parameters during synthesis, the pure ZrO₂ material shares some common behavior with the doped samples reported in literature, but also major differences. The latter is basically reflected in the much higher structural stability, especially at low temperatures and a totally different surface reactivity.

- This then enables a proper physico-chemical characterization of this prominent modification in the ZrO₂ polymorphic system. These investigations demonstrate unique aspects of the used t-ZrO₂ sample. Despite the fact that ZrO₂, and especially in this case metastable tetragonal ZrO₂, seems to be a well-investigated polymorph in literature, the unique properties point out the significant impact of sample preparation routine and sample history.

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